Integrating the Healthcare Enterprise

IHE IT Infrastructure (ITI) Technical Framework

Volume 1 (ITI TF-1) Integration Profiles

Revision 5.0 – Final Text

December 12, 2008

Content

1 Introduction

Integrating the Healthcare Enterprise (IHE) is an initiative designed to stimulate the integration of the information systems that support modern healthcare institutions. Its fundamental objective is to ensure that in the care of patients all required information for medical decisions is both correct and available to healthcare professionals. The IHE initiative is both a process and a forum for encouraging integration efforts. It defines a technical framework for the implementation of established messaging standards to achieve specific clinical goals. It includes a rigorous testing process for the implementation of this framework. And it organizes educational sessions and exhibits at major meetings of medical professionals to demonstrate the benefits of this framework and encourage its adoption by industry and users.

The approach employed in the IHE initiative is to support the use of existing standards, e.g HL7, ASTM, DICOM, ISO, IETF, OASIS and others as appropriate, rather than to define new standards. IHE profiles further constrain configuration choices where necessary in these standards to ensure that they can be used in their respective domains in an integrated manner between different actors. When clarifications or extensions to existing standards are necessary, IHE refers recommendations to the relevant standards bodies.

This initiative has numerous sponsors and supporting organizations in different medical specialty domains and geographical regions. In North America the primary sponsors are the Healthcare Information and Management Systems Society (HIMSS) and the Radiological Society of North America (RSNA). IHE Canada has also been formed. IHE Europe (IHE-EUR) is supported by a large coalition of organizations including the European Association of Radiology (EAR) and European Congress of Radiologists (ECR), the Coordination Committee of the Radiological and Electromedical Industries (COCIR), Deutsche Röntgengesellschaft (DRG), the EuroPACS Association, Groupement pour la Modernisation du Système d'Information Hospitalier (GMSIH), Société Francaise de Radiologie (SFR), Società Italiana di Radiologia Medica (SIRM), and the European Institute for health Records (EuroRec). In Japan IHE-J is sponsored by the Ministry of Economy, Trade, and Industry (METI); the Ministry of Health, Labor, and Welfare; and MEDIS-DC; cooperating organizations include the Japan Industries Association of Radiological Systems (JIRA), the Japan Association of Healthcare Information Systems Industry (JAHIS), Japan Radiological Society (JRS), Japan Society of Radiological Technology (JSRT), and the Japan Association of Medical Informatics (JAMI). Other organizations representing healthcare professionals are invited to join in the expansion of the IHE process across disciplinary and geographic boundaries.

1.1 Overview of the Technical Framework

This document, the IHE IT Infrastructure Technical Framework (ITI TF), defines specific implementations of established standards to achieve integration goals that promote appropriate sharing of medical information to support optimal patient care. It is expanded annually, after a period of public review, and maintained regularly through the identification and correction of errata. The current version, Revision 5.0 for Final Text, specifies the IHE transactions defined and implemented as of October 2008. The latest version of the document is always available via the Internet at http://www.ihe.net/Technical_Framework .

The IHE IT Infrastructure Technical Framework identifies a subset of the functional components of the healthcare enterprise, called IHE actors, and specifies their interactions in terms of a set of coordinated, standards-based transactions. It describes this body of transactions in progressively greater depth. The present volume (ITI TF-1) provides a high-level view of IHE functionality, showing the transactions organized into functional units called integration profiles that highlight their capacity to address specific IT Infrastructure requirements.

Volume 2 of the IT Infrastructure Technical Framework (ITI TF-2) provides detailed technical descriptions of each IHE transaction used in the IT Infrastructure Integration Profiles. These two volumes are consistent and can be used in conjunction with the Integration Profiles of other IHE domains.

The other domains within the IHE initiative also produce Technical Frameworks within their respective areas that together form the IHE Technical Framework. For example, the following IHE Technical Framework(s) are some of those which are available:

- IHE IT Infrastructure Technical Framework
- IHE Cardiology Technical Framework
- IHE Laboratory Technical Framework
- IHE Patient Care Coordination Technical Framework
- IHE Radiology Technical Framework

Where applicable, references are made to other technical frameworks. For the conventions on referencing other frameworks, see Section 1.6.3 within this volume.

1.2 Overview of the IT Infrastructure Volume I

The remainder of Section 1 further describes the general nature, purpose and function of the Technical Framework. Section 2 introduces the concept of IHE Integration Profiles that make up the Technical Framework.

Section 3 and the subsequent sections of this volume provide detailed documentation on each integration profile, including the IT Infrastructure problem it is intended to address and the IHE actors and transactions it comprises.

The appendices following the main body of the document provide a summary list of the actors and transactions, detailed discussion of specific issues related to the integration profiles and a glossary of terms and acronyms used.

1.3 Audience

The intended audience of this document is:

- IT departments of healthcare institutions
- Technical staff of vendors participating in the IHE initiative
- Experts involved in standards development
- Those interested in integrating healthcare information systems and workflows

1.4 Relationship to Standards

The IHE Technical Framework identifies functional components of a distributed healthcare environment (referred to as IHE actors), solely from the point of view of their interactions in the healthcare enterprise. At its current level of development, it defines a coordinated set of transactions based on ASTM, DICOM, HL7, IETF, ISO, OASIS and W3C standards. As the scope of the IHE initiative expands, transactions based on other standards may be included as required.

In some cases, IHE recommends selection of specific options supported by these standards; however, IHE does not introduce technical choices that contradict conformance to these standards. If errors in or extensions to existing standards are identified, IHE's policy is to report them to the appropriate standards bodies for resolution within their conformance and standards evolution strategy.

IHE is therefore an implementation framework, not a standard. Conformance claims for products must still be made in direct reference to specific standards. In addition, vendors who have implemented IHE integration capabilities in their products may publish IHE Integration Statements to communicate their products' capabilities. Vendors publishing IHE Integration Statements accept full responsibility for their content. By comparing the IHE Integration Statements from different products, a user familiar with the IHE concepts of actors and integration profiles can determine the level of integration between them. See Appendix C for the format of IHE Integration Statements.

1.5 Relationship to Real-world Architectures

The IHE actors and transactions described in the IHE Technical Framework are abstractions of the real-world healthcare information system environment. While some of the transactions are traditionally performed by specific product categories (e.g. HIS, Clinical Data Repository, Radiology Information Systems, Clinical Information Systems or Cardiology Information Systems), the IHE Technical Framework intentionally avoids associating functions or actors with such product categories. For each actor, the IHE Technical Framework defines only those functions associated with integrating information systems. The IHE definition of an actor should therefore not be taken as the complete definition of any product that might implement it, nor should the framework itself be taken to comprehensively describe the architecture of a healthcare information system.

The reason for defining actors and transactions is to provide a basis for defining the interactions among functional components of the healthcare information system environment. In situations where a single physical product implements multiple functions, only the interfaces between the product and external functions in the environment are considered to be significant by the IHE initiative. Therefore, the IHE initiative takes no position as to the relative merits of an integrated environment based on a single, all-encompassing information system versus one based on multiple systems that together achieve the same end. IHE demonstrations emphasize the integration of multiple vendors' systems based on the IHE Technical Framework.

1.6 Conventions

This document has adopted the following conventions for representing the framework concepts and specifying how the standards upon which the IHE Technical Framework is based should be applied.

1.6.1 IHE Actor and Transaction Diagrams and Tables

Each integration profile is a representation of a real-world capability that is supported by a set of actors that interact through transactions. Actors are information systems or components of information systems that produce, manage, or act on categories of information required by operational activities in the enterprise. Transactions are interactions between actors that communicate the required information through standards-based messages.

The diagrams and tables of actors and transactions in subsequent sections indicate which transactions each actor in a given profile must support.

The transactions shown on the diagrams are identified both by their name and the transaction number as defined in ITI TF-2. The transaction numbers are shown on the diagrams as bracketed numbers prefixed with the specific Technical Framework domain.

In some cases, a profile is dependent on a prerequisite profile in order to function properly and be useful. For example, Enterprise User Authentication depends on Consistent Time. These dependencies can be found by locating the desired profile in Table 2-1 to determine which profile(s) are listed as prerequisites. An actor must implement all required transactions in the prerequisite profiles in addition to those in the desired profile.

1.6.2 Process Flow Diagrams

The descriptions of integration profiles that follow include process flow diagrams that illustrate how the profile functions as a sequence of transactions between relevant actors.

These diagrams are intended to provide an overview so the transactions can be seen in the context of an institution's workflow. Certain transactions and activities not defined in detail by IHE are shown in these diagrams in *italics* to provide additional context on where the relevant IHE transactions fit into the broader scheme of healthcare information systems.

These diagrams are not intended to present the only possible scenario. Often other actor groupings are possible, and transactions from other profiles may be interspersed.

In some cases the sequence of transactions may be flexible. Where this is the case there will generally be a note pointing out the possibility of variations. Transactions are shown as arrows oriented according to the flow of the primary information handled by the transaction and not necessarily the initiator.

1.6.3 Technical Framework Cross-references

When references are made to another section within a Technical Framework volume, a section number is used by itself. When references are made to other volumes or to a Technical Framework in another domain, the following format is used:

<domain designator> TF-<volume number>: <section number>, where

<domain designator> is a short designator for the IHE domain (ITI = IT Infrastructure, RAD = Radiology)

<volume number> is the applicable volume within the given Technical Framework (e.g., 1, 2, 3), and

<section number> is the applicable section number.

For example: ITI TF-1: 3.1 refers to Section 3.1 in volume 1 of the IHE IT Infrastructure Technical Framework. RAD TF-3: 4.33 refers to Section 4.33 in volume 3 of the IHE Radiology Technical Framework. ITI TF-2: Appendix B refers to Appendix B in volume 2 of the IHE IT Infrastructure Technical Framework.

When references are made to Transaction numbers in the Technical Framework, the following format is used:

[<domain designator>-<transaction number>], where

<transaction number> is the transaction number within the specified domain.

For example: [ITI-1] refers to Transaction 1 from the IHE IT Infrastructure Technical Framework.

1.7 Scope of Changes Introduced in the Current Year

The IHE Technical Framework is updated annually to reflect new profiles, corrections and new transactions (refer to ITI TF-2) used in those profiles.

This document expands the V4.0 IT Infrastructure Technical Framework and includes integration profiles developed in the previous years as well as the new profiles finalized in the 2007-2008 cycle of the IHE IT Infrastructure initiative. It will be the basis for the 2009 connectathon testing and exhibition process associated in particular with the HIMSS 2009 annual meeting.

Retrieve Information for Display (RID) – a simple and rapid read-only access to patient information necessary for provision of better care. It supports access to existing persistent documents in well-known presentation formats such as CDA, PDF, JPEG, etc. It also supports access to specific key patient-centric information such as allergies, current medications, summary of reports, etc. for presentation to a clinician.

Enterprise User Authentication (EUA) – a means to establish one name per user that can then be used on all of the devices and software that participate in this integration profile, greatly facilitating centralized user authentication management and providing users with the convenience and speed of a single sign-on. This profile leverages Kerberos (RFC 1510) and the HL7 CCOW standard (user subject).

Patient Identifier Cross-referencing (PIX) – provides cross-referencing of patient identifiers from multiple Patient Identifier Domains. These patient identifiers can then be used by identity consumer systems to correlate information about a single patient from sources that know the patient by different identifiers.

Patient Synchronized Applications (PSA) – a means for viewing data for a single patient using independent and unlinked applications on a user's workstation, reducing the repetitive tasks of selecting the same patient in multiple applications. Data can be viewed from different Identifier Domains when used with the Patient Identifier Cross-referencing Integration Profile to resolve multiple identifications for the same patient. This profile leverages the HL7 CCOW standard specifically for patient subject context management. .

Consistent Time (CT) – mechanisms to synchronize the time base between multiple actors and computers. Various infrastructure, security, and acquisition profiles require use of a consistent time

base on multiple computers. The Consistent Time Profile provides a median synchronization error of less than 1 second.

This Version 3.0 IT Infrastructure Technical Framework finalizes four new Integration Profiles developed and tested in the 2004-2005 cycle:

Patient Demographics Query (PDQ) – provides ways for multiple distributed applications to query a central patient information server for a list of patients, based on user-defined search criteria, and retrieve a patient's demographic (and, optionally, visit or visit-related) information directly into the application.

Audit Trail and Node Authentication (ATNA) – establishes the characteristics of a Basic Secure Node:

1. It describes the security environment (user identification, authentication, authorization, access control, etc.) assumed for the node so that security reviewers may decide whether this matches their environments.
2. It defines basic auditing requirements for the node
3. It defines basic security requirements for the communications of the node using TLS or equivalent functionality.
4. It establishes the characteristics of the communication of audit messages between the Basic Secure Nodes and Audit Repository nodes that collect audit information.

This profile has been designed so that specific domain frameworks may extend it through an option defined in the domain specific technical framework. Extensions are used to define additional audit event reporting requirements, especially actor specific requirements. The Radiology Audit Trail option in the IHE Radiology Technical Framework is an example of such an extension.

Personnel White Pages (PWP) – provides access to basic human workforce user directory information. This information has broad use among many clinical and non-clinical applications across the healthcare enterprise.

Cross-Enterprise Document Sharing (XDS) – enables a number of healthcare delivery organizations belonging to an XDS Affinity Domain (e.g. a community of care) to cooperate in the care of a patient by sharing clinical records in the form of documents as they proceed with their patients' care delivery activities. This profile is based upon ebXML Registry standards, SOAP, HTTP and SMTP. It describes the configuration of an ebXML Registry in sufficient detail to support Cross Enterprise Document Sharing.

Cross-Enterprise User Assertion Profile (XUA) - provides a means to communicate claims about the identity of an authenticated principal (user, application, system...) in transactions that cross enterprise boundaries. To provide accountability in these cross-enterprise transactions there is a need to identify the requesting principal in a way that enables the receiver to make access decisions and generate the proper audit entries. The XUA Profile supports enterprises that have chosen to have their own user directory with their own unique method of authenticating the users, as well as others that may have chosen to use a third party to perform the authentication.

Patient Administration Management (PAM) - provides patient identity, registration, and encounter management transactions in a healthcare enterprise as well as across enterprises.

Cross-Enterprise Document Media Interchange (XDM) - provides document interchange using a common file and directory structure over several standard media. This permits the patient to use physical media to carry medical documents. This also permits the use of person-to-person email to convey medical documents.

Basic Patient Privacy Consents (BPPC) - provides a mechanism to record the patient privacy consent(s), a method to mark documents published to XDS with the patient privacy consent that was used to authorize the publication, and a method for XDS Consumers to use to enforce the privacy consent appropriate to the use. The XDS profile provides little guidance on supporting privacy policies within an XDS Affinity Domain. Documents can be marked with a confidentialityCode, but no information is provided on how to use this information to support patient privacy concerns. This profile complements XDS by describing a mechanism whereby an XDS Affinity Domain can develop and implement multiple privacy policies, and describes how that mechanism can be integrated with the access control mechanisms supported by the XDS Actors (e.g. EHR systems).

Cross Enterprise Sharing of Scanned Documents (XDS-SD) – A profile which associates structured, healthcare metadata with non-healthcare specific document format to maintain the integrity of the patient health record as managed by the source system.

1.8 Security Implications

IHE transactions often contain information that must be protected in conformance with privacy laws and regulations, such as HIPAA or similar requirements in other regions. IHE includes a few security and privacy-focused profiles listed below. Other IHE Profiles generally do not have specific privacy protections, but rather expect a proper grouping with one or more of the security profiles:

- The Audit Trail and Node Authentication (ATNA) profile specifies a means to ensure that nodes in a network are authenticated.
- The ATNA profile specifies an audit message for reporting security- and privacy-relevant events.
- The Enterprise User Authentication (EUA) profile specifies a means to authenticate system users and to share knowledge of the authenticated users among applications.
- The Personnel White Pages (PWP) profile provides a repository that may be used to hold system users' identification data.

Implementers may follow these IHE profiles to fulfill some of their security needs. It is understood that institutions must implement policy and workflow steps to satisfy enterprise needs and to comply with regulatory requirements.

1.9 Comments

HIMSS and RSNA welcome comments on this document and the IHE initiative. They should be directed to the discussion server at http://ihe.rsna.org/ihetf/ or to:

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1.10 Copyright Permission

1.11 IHE Technical Framework Development and Maintenance Process

The IHE IT Infrastructure Technical Framework is continuously maintained and expanded on an annual basis by the IHE IT Infrastructure Technical Committee. The development and maintenance process of the Framework follows a number of principles to ensure stability of the specification so that both vendors and users may use it reliably in specifying, developing and acquiring systems with IHE integration capabilities.

The first of these principles is that any extensions, clarifications and corrections to the Technical Framework must maintain backward compatibility with previous versions of the framework in order to maintain interoperability with systems that have implemented IHE Actors and Integration Profiles defined there.

The IHE IT Infrastructure Technical Framework is developed and re-published annually following a three-step process:

1. The IT Infrastructure Technical Committee develops supplements to the current stable version of the Technical Framework to support new functionality identified by the IHE Strategic and Planning Committees and issues them for public comment.
2. The Committee addresses all comments received during the public comment period and publishes an updated version of the Technical Framework for “Trial Implementation.” This version contains both the stable body of the Technical Framework from the preceding cycle and the newly developed supplements. It is the version of the Technical Framework used by vendors in developing trial implementation software for the annual IT Infrastructure Connectathon.
3. The Committee regularly considers change proposals to the Trial Implementation version of the Technical Framework, including those from implementers who participate in the Connectathon. After resolution of all change proposals received within 60 days of the Connectathon, the Technical Framework version is published as “Final Text”.

2 IT Infrastructure Integration Profiles

IHE IT Infrastructure Integration Profiles (Figure 2-1), offer a common language that healthcare professionals and vendors can use to discuss integration needs of healthcare enterprises and the integration capabilities of information systems in precise terms. Integration Profiles specify implementations of standards that are designed to meet identified clinical needs. They enable users and vendors to state which IHE capabilities they require or provide, by reference to the detailed specifications of the IHE IT Infrastructure Technical Framework.

Integration profiles are defined in terms of IHE Actors and transactions. Actors (see ITI TF-1, Appendix A) are information systems or components of information systems that produce, manage, or act on information associated with clinical and operational activities in the enterprise. Transactions (see ITI TF-1, Appendix B) are interactions between actors that communicate the required information through standards-based messages.

Vendor products support an Integration Profile by implementing the appropriate actor(s) and transactions. A given product may implement more than one actor and more than one integration profile.

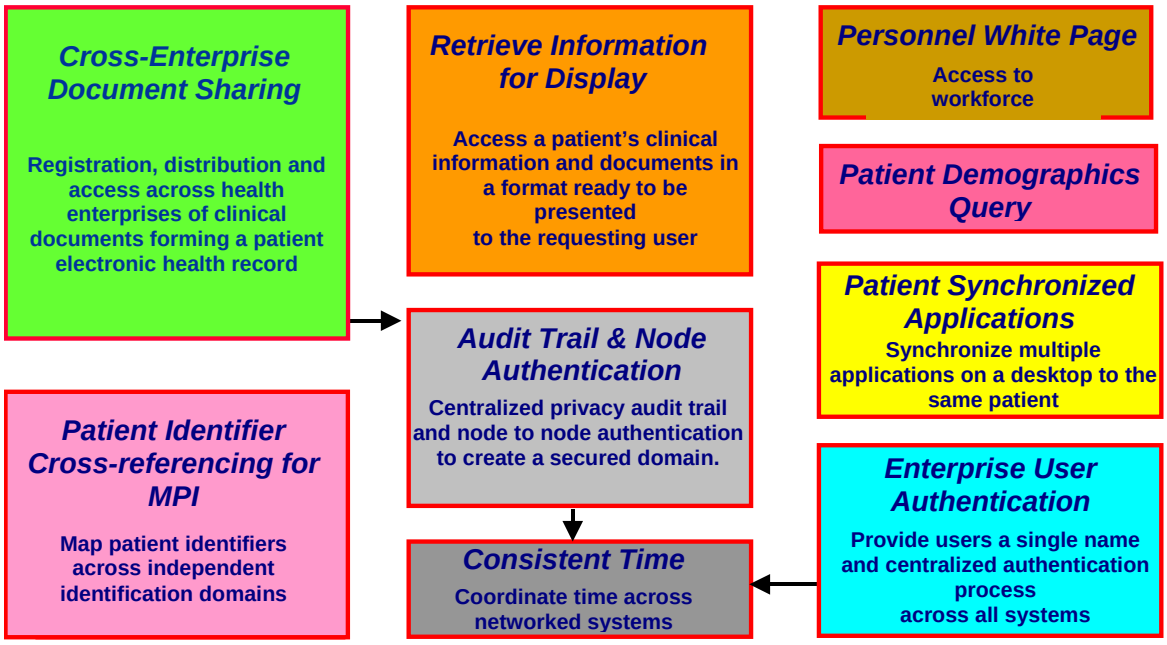


Figure 2-1 IHE IT Infrastructure Integration Profiles

2.1 Dependencies among Integration Profiles

Dependencies among IHE Integration Profiles exist when implementation of one integration profile is a prerequisite for achieving the functionality defined in another integration profile. Figure 2-1 provides a graphical view of the dependencies among IHE IT Infrastructure Integration Profiles. The arrows in the figure point from a given integration profile to the integration profile(s) upon which it depends. Table 2-1 defines these dependencies in tabular form.

Some dependencies require that an actor supporting one profile be grouped with one or more actors supporting other integration profiles. For example, Enterprise User Authentication (EUA) requires that different participating actors be grouped with the Time Client Actor that participates in the Consistent Time (CT) Integration Profile. The dependency exists because EUA actors must refer to consistent time in order to function properly.

Table 2-1 Integration Profiles Dependencies

Integration Profile	Depends on	Dependency Type	Purpose
Retrieve Information for Display Integration	*None*	None	-
Enterprise User Authentication	Consistent Time	Each actor implementing EUA shall be grouped with the Time Client Actor	- Required to manage expirations of authentication tickets
Patient Identifier Cross-referencing	Consistent Time	Each actor implementing PIX shall be grouped with the Time Client Actor	Required to manage and resolve conflicts in multiple updates.
Patient Synchronized Applications	*None*	*None*	-
Consistent Time	*None*	*None*	-
Patient Demographics Query	*None*	*None*	-
Personnel White Pages	*None*	*None*	-
Audit trail and Node Authentication	Consistent Time	Each actor implementing ATNA shall be grouped with the Time Client Actor	- Required for consistent time in audit logs.
Cross-Enterprise Document Sharing	Audit Trail and Node Authentication	Each XDS Actor must be grouped with the Secure Node Actor.	- Required to manage audit trail of exported PHI, node authentication and transport encryption.
Cross-Enterprise Document Sharing	Consistent Time	Each XDS actor must be grouped with the Time Client Actor	To ensure consistency among document and submission set dates.
Cross-Enterprise User Assertion	None	None	
Patient Administration Management	None	None	-
Cross-Enterprise Document Media Interchange	ATNA	Requires secure communication, and audit trails.	
Basic Patient Privacy Consent	XDS, XDM, XDR	XDS Metadata	Indicates Patient Privacy Consent Policy applied to document
Cross Enterprise Sharing of Scanned Documents (XDS-SD)	XDS, XDM or XDR	This content is created and consumed by actors grouped with actors in either XDS, XDR or XDM.	The content of this profile is intended for use in XDS, XDR and XDM.

To support a dependent profile, an actor must implement all required transactions in the prerequisite profiles in addition to those in the dependent profile. In some cases, the prerequisite is that the actor selects any one of a given set of profiles.

2.2 Integration Profiles Overview

In this document, each IHE Integration Profile is defined by:

- The IHE actors involved
- The specific set of IHE transactions exchanged by each IHE actor.

These requirements are presented in the form of a table of transactions required for each actor supporting the Integration Profile. Actors supporting multiple Integration Profiles are required to support all the required transactions of each Integration Profile supported. When an Integration Profile depends upon another Integration Profile, the transactions required for the dependent Integration Profile have not been included in the table.

Note that IHE Integration Profiles are not statements of conformance to standards, and IHE is not a certifying body. Users should continue to request that vendors provide statements of their conformance to standards issued by relevant standards bodies, such as HL7 and DICOM. Standards conformance is a prerequisite for vendors adopting IHE Integration Profiles.

Also note that there are critical requirements for any successful integration project that IHE cannot address. Successfully integrating systems still requires a project plan that minimizes disruptions and describes fail-safe strategies, specific and mutually understood performance expectations, well-defined user interface requirements, clearly identified systems limitations, detailed cost objectives, plans for maintenance and support, etc.

2.2.1 This section is reserved.

2.2.2 This section is reserved.

2.2.3 Retrieve Information for Display (RID)

Retrieve Information for Display enables simple and rapid access to patient information for better care. It supports access to existing persistent documents in well-known presentation formats such as CDA, PDF, JPEG, etc. It also supports access to specific key patient-centric information such as allergies, current medications, summary of reports, etc. for presentation to a clinician. It complements workflows from within the users' on-screen workspace or application. By linking it with two other IHE profiles - Enterprise User Authentication and Patient Identifier Cross-referencing, this profile's reach can extend across organization boundaries within an enterprise. This IHE Integration Profile leverages HTTP, Web Services, IT presentation formats and HL7 CDA Level 1.

2.2.4 Enterprise User Authentication (EUA)

Enterprise User Authentication defines a means to establish one name per user that can then be used on all of the devices and software that participate in this integration profile. It greatly facilitates centralized user authentication management and provides users with the convenience and speed of a single sign-on. This profile leverages Kerberos (RFC 1510) and the HL7 CCOW standard (user subject). User authentication is a necessary step for most application and data access operations and streamlines workflow for users. Future profiles will deal with other security issues, such as authorization management.

2.2.5 Patient Identifier Cross-referencing (PIX)

The PIX profile supports the cross-referencing of patient identifiers from multiple Patient Identifier Domains. These cross-referenced patient identifiers can then be used by "identity consumer" systems to correlate information about a single patient from sources that "know" the patient by different identifiers. This allows a clinician to have more complete view of the patient information.

2.2.6 Patient Synchronized Applications (PSA)

Patient Synchronized Applications supports viewing data for a single patient among otherwise independent and unlinked applications on a user's workstation. Its implementation reduces the repetitive tasks of selecting the same patient in multiple applications. It also improves patient safety by reducing the chance of medical errors caused by viewing the wrong patient's data. Its ability to work with the Patient Identifier Cross-referencing provides a seamless environment for clinicians and IT staff. This profile leverages the HL7 CCOW standard specifically for patient subject context management.

2.2.7 Consistent Time (CT)

Consistent Time Profile defines mechanisms to synchronize the time base between multiple actors and computers. Various infrastructure, security, and acquisition profiles require use of a consistent time base on multiple computers. The Consistent Time Profile provides median synchronization error of less than 1 second. Configuration options can provide better synchronization. The Consistent Time profile specifies the use of the Network Time Protocol (NTP) defined in RFC 1305.

2.2.8 Patient Demographics Query (PDQ)

Patient Demographics Query provides ways for multiple distributed applications to query a central patient information server for a list of patients, based on user-defined search criteria, and retrieve a patient's demographic (and, optionally, visit or visit-related) information directly into the application.

2.2.9 Audit Trail and Node Authentication (ATNA)

Audit Trail and Node Authentication establishes the characteristics of a Basic Secure Node:

1. It describes the security environment (user identification, authentication, authorization, access control, etc.) assumed for the node so that security reviewers may decide whether this matches their environments.
2. It defines basic auditing requirements for the node
3. It defines basic security requirements for the communications of the node using TLS or equivalent functionality.
4. It establishes the characteristics of the communication of audit messages between the Basic Secure Nodes and Audit Repository nodes that collect audit information.
5. It defines a Secure Application actor for describing product configurations that are not able to meet all of the requirements of a Secure Node.

Note: ATNA security considerations require the use of Secure Nodes. The Secure Application is defined to permit product configurations to indicate that the product is ready for easy integration into a Secure Node environment because it performs all of the security related functions that are directly related to the application function. See section 9.7 for more details.

This profile has been designed so that specific domain frameworks may extend it through an option defined in the domain specific technical framework. Extensions are used to define additional audit event reporting requirements, especially actor specific requirements. The Radiology Audit Trail option in the IHE Radiology Technical Framework is an example of such an extension.

2.2.10 Cross-Enterprise Document Sharing (XDS)

Cross-Enterprise Document Sharing enables a number of healthcare delivery organizations belonging to an XDS Affinity Domain (e.g. a community of care) to cooperate in the care of a patient by sharing clinical records in the form of documents as they proceed with their patients' care delivery activities. Federated document repositories and a document registry create a longitudinal record of information about a patient within a given XDS Affinity Domain. This profile is based upon ebXML Registry standards, SOAP, HTTP and SMTP. It describes the configuration of an ebXML Registry in sufficient detail to support Cross Enterprise Document Sharing.

2.2.11 Personnel White Pages (PWP)

Personnel White Pages Profile (PWP) provides access to basic human workforce user directory information. This information has broad use among many clinical and non-clinical applications across the healthcare enterprise. The information can be used to enhance the clinical workflow (contact information), enhance the user interface (user friendly names and titles), and ensure identity (digital certificates). This Personnel White Pages directory will be related to the User Identity provided by the Enterprise User Authentication (EUA) Integration Profile previously defined by IHE.

2.2.12 This section is reserved for Notification of Document Availability (NAV)

2.2.13 Cross Enterprise User Assertion (XUA)

Cross-Enterprise User Assertion provides a means to communicate claims about the identity of an authenticated principal (user, application, system...) in transactions that cross-enterprise boundaries. To provide accountability in these cross enterprise transactions there is a need to identify the requesting principal in a way that enables the receiver to make access decisions and generate the proper audit entries. The XUA Profile supports enterprises that have chosen to have their own user directory with their own unique method of authenticating the users, as well as others that may have chosen to use a third party to perform the authentication.

2.2.14 Patient Administration Management (PAM)

The Patient Administration Management Integration Profile establishes the continuity and integrity of patient data, and additional information such as related persons (primary caregiver, guarantor, next of kin, etc.). It coordinates the exchange of patient registration and update information among systems that need to be able to provide current information regarding a patient's encounter status and location. This profile supports ambulatory and acute care use cases including patient identity

feed, admission and discharge, and transfer and encounter management, as well as explicit and precise error reporting and application acknowledgment.

The PAM profile supports two patient encounter management scenarios: either one single central patient registration system serving the entire institution, or multiple patient registration systems collaborating as peers serving different clinical settings in an institution.

2.2.15 Cross-Enterprise Document Media Interchange (XDM)

Cross-Enterprise Document Media Interchange provides document interchange using a common file and directory structure over several standard media. This permits the patient to use physical media to carry medical documents. This also permits the use of person-to-person email to convey medical documents.

2.2.16 Basic Patient Privacy Consents (BPPC)

The Basic Patient Privacy Consents profile provides a mechanism to record the patient privacy consent(s), a method to mark documents published to XDS with the patient privacy consent that was used to authorize the publication, and a method for XDS Consumers to use to enforce the privacy consent appropriate to the use. The XDS profile provides little guidance on supporting privacy policies within an XDS Affinity Domain. Documents can be marked with a confidentialityCode, but no information is provided on how to use this information to support patient privacy concerns. This profile complements XDS by describing a mechanism whereby an XDS Affinity Domain can develop and implement multiple privacy policies, and describes how that mechanism can be integrated with the access control mechanisms supported by the XDS Actors (e.g. EHR systems).

2.2.17 Scanned Documents Integration Profile (XDS-SD)

A variety of legacy paper, film, electronic and scanner outputted formats are used to store and exchange clinical documents. These formats are not designed for healthcare documentation, and furthermore, do not have a uniform mechanism to store healthcare metadata associated with the documents, including patient identifiers, demographics, encounter, order or service information. The association of structured, healthcare metadata with this kind of document is important to maintain the integrity of the patient health record as managed by the source system. It is necessary to provide a mechanism that allows such source metadata to be stored with the document.

2.3 Product Implementations

Developers have a number of options in implementing IHE actors and transactions in product implementations. The decisions cover three classes of optionality:

- For a system, select which actors it will incorporate (multiple actors per system are acceptable).
- For each actor, select the integration profiles in which it will participate.
- For each actor and profile, select which options will be implemented.

All required transactions must be implemented for the profile to be supported (refer to the transaction descriptions in ITI TF-2).

Implementers should provide a statement describing which IHE actors, IHE integration profiles and options are incorporated in a given product. The recommended form for such a statement is defined in ITI TF-1, Appendix C.

In general, a product implementation may incorporate any single actor or combination of actors. When two or more actors are grouped together, internal communication between actors is assumed to be sufficient to allow the necessary information flow to support their functionality; for example, the Context Manager uses the Patient Identifier Cross-reference Consumer Actor to obtain the necessary patient identifier mapping information from the Patient Identifier Cross-reference Manager. The exact mechanisms of such internal communication are outside the scope of the IHE Technical Framework.

When multiple actors are grouped in a single product implementation, all transactions originating or terminating with each of the supported actors shall be supported (i.e., the IHE transactions shall be offered on an external product interface).

The following examples describe which actors typical systems might be expected to support. This is not intended to be a requirement, but rather to provide illustrative examples.

A departmental system, such as a laboratory information system or a radiology picture archiving and communication system might include an Information Source Actor as well as a Kerberized Server Actor.

A clinical repository might include an Information Source Actor as well as a Kerberized Server Actor and a Patient Identifier Cross-reference Consumer Actor.

A context management server might include a Context Management Actor as well as a Patient Identifier Cross-reference Consumer Actor.

3 Retrieve Information for Display (RID)

The ***Retrieve Information for Display Integration Profile (RID)*** provides simple and rapid read-only access to patient-centric clinical information that is located outside the user's current application but is important for better patient care (for example, access to lab reports from radiology department). It supports access to existing persistent documents in well-known presentation formats such as CDA (Level 1), PDF, JPEG, etc. It also supports access to specific key patient-centric information such as allergies, current medications, summary of reports, etc. for presentation to a clinician. It complements workflows with access from within the users' on-screen workspace or application to a broad range of information.

In this profile, the Information Source is solely responsible to turn the healthcare specific semantics into what this IHE Integration Profile calls a "presentation" format. As a consequence the Display actor may process and render this "presentation" format with only generic healthcare semantics knowledge. Different formats have specific characteristics in terms of (1) server imposed limitations and (2) flexibility of display on the client side to render within its display constraints (e.g. a generic CDA level 1 style sheet).

The Information Source is entirely responsible for the information returned for display and its clinical accuracy.

This profile offers the capability to leverage industry standards that address both the structure and content of documents that may be returned by information sources. Where this profile references HL7 Clinical Documentation Architecture (CDA), it limits itself to the approved CDA Level 1. Furthermore, it only uses a subset of CDA Level 1 that facilitates making information available for display.

Future extensions to the IHE IT Infrastructure TF will more fully leverage CDA Release 2 and other industry standards, and will incorporate vocabularies such as SNOMED and Clinical LOINC as well as clinical templates.

This profile does not provide specific requirements on the means of assuring access control or security of information in transit. Such measures shall be implemented through appropriate security-related integration profiles, such as Enterprise User Authentication (see Section 4). Appendix E describes the process flows for usage of the Retrieve Information for Display Integration Profile in conjunction with the Enterprise User Authentication and Patient Identifier Cross-referencing Integration Profiles.

3.1 Actors/ Transactions

Figure 3.1-1 shows the actors directly involved in the Retrieve Information for Display Integration Profile and the relevant transactions between them. Other actors that may be indirectly involved due to their participation in User Authentication and Patient Identifier Cross-referencing are not shown.

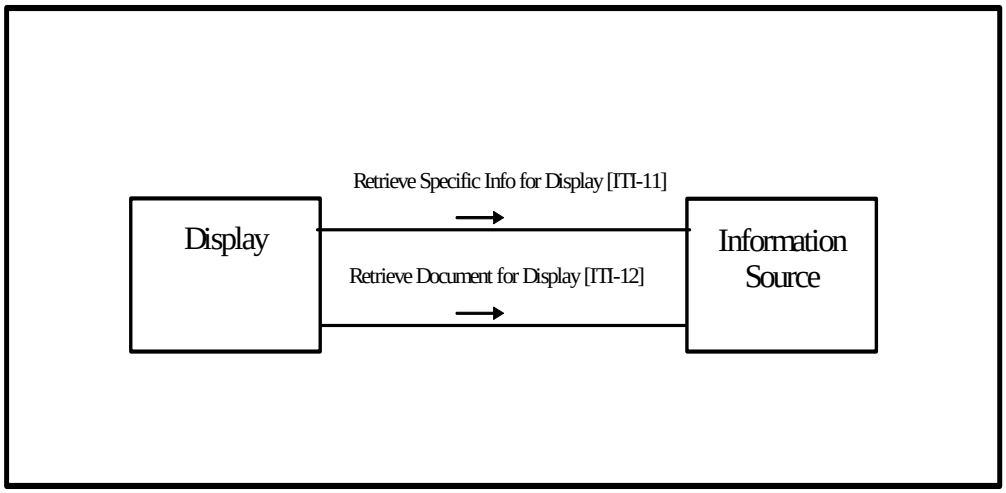


Figure 3.1-1. Retrieve Information for Display Actor Diagram

Table 3.1-1 lists the transactions for each actor directly involved in the Retrieve Information for Display Integration Profile. In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled "R"). A complete list of options defined by this Integration Profile and that implementations may choose to support is listed in ITI TF-1: 3.2.

Table 3.1-1 Retrieve Information for Display Integration Profile - Actors and Transactions

Actors	Transactions	Optionality	Section in Vol. 2
Display	Retrieve Specific Info for Display[ITI-11]	R	ITI TF-2: 3.11
	Retrieve Document for Display[ITI-12]	R	ITI TF-2: 3.12
Information Source	Retrieve Specific Info for Display[ITI-11]	R (see below)	ITI TF-2: 3.11
	Retrieve Document for Display[ITI-12]	R (see below)	ITI TF-2: 3.12

Transaction [ITI-11] is required if one of the following Options is selected by the Information Source Actor (See Section 3.2):

Summary of All Reports
Summary of Laboratory Reports
Summary of Radiology Reports
Summary of Cardiology Reports
Summary of Surgery Reports
Summary of Intensive Care Reports
Summary of Emergency Reports
Summary of Discharge Reports
Summary of Prescriptions
List of Allergies and Adverse Reactions
List of Medications

Transaction [ITI-12] is required if the Persistent Document Option is selected by the Information Source Actor (See Section 3.2).

The means for a Display Actor to obtain documents' unique identifiers in order to retrieve them via Transaction [ITI-11] may be either via Transaction [ITI-12] or by other means that are outside the scope of the RID Integration Profile.

3.2 Retrieve Information for Display Integration Profile Options

Options that may be selected for this Integration Profile are listed in the Table 3.2-1 along with the IHE actors to which they apply.

Table 3.2-1 Retrieve Information for Display - Actors and Options

Actor	Options	Vol & Section
Display	*None*	- -
Information Source	*Persistent Document*	ITI TF-2: 3.12
	Summary of All Reports (note2)	ITI TF-2: 3.11
	Summary of Laboratory Reports (note2)	ITI TF-2: 3.11
	Summary of Radiology Reports (note2)	ITI TF-2: 3.11
	Summary of Cardiology Reports (note2)	ITI TF-2: 3.11
	Summary of Surgery Reports (note2)	ITI TF-2: 3.11
	Summary of Intensive Care Reports (note2)	ITI TF-2: 3.11
	Summary of Emergency Reports (note2)	ITI TF-2: 3.11
	Summary of Discharge Reports (note2)	ITI TF-2: 3.11
	Summary of Prescriptions (note2)	ITI TF-2: 3.11
	List of Allergies and Adverse Reactions	ITI TF-2: 3.11
	List of Medications (note1)	ITI TF-2: 3.11

Note1: List of Medications includes the list of medications currently known to be administered to the patient. It differs from the Summary of Prescriptions, in that the latter reflects what has been prescribed to the patient, but are not necessarily any longer administered.

Note2: In all the above options, "summary of reports" means that a general patient context (patient name, etc.) is provided along with a list of entries, where an entry includes key attributes such as date, specialty, and additional information sufficient to allow the viewer to select an entry. An entry may reference a persistent document for RID or other application defined RID summaries. Beyond these general guidelines, the specific content may likely be influenced by the context of use and customer desires. Such summaries are non-persistent in that they are likely to be updated in the course of patient care.

3.3 Retrieve Information for Display Process Flow

This section describes the process and information flow when displayable patient information is retrieved from an information source. Three cases are distinguished.

Case 1-Retrieve *Specific* Information for Display: The first case describes use cases when the display actor and the person associated are requesting some information related to a patient. A somewhat specific request for information is issued (e.g. Retrieve a summary of laboratory reports) for a specific Patient ID to an Information Source Actor. The patient ID is assumed to be unambiguous as fully qualified with the assigning authority. A number of additional filtering keys may be used (last N reports, date range, etc.) depending on the specific type of request issued. The Information Source Actor responds with presentation-ready information that it considers relevant to the request. This Integration Profile leaves entire flexibility to the Information Source Actor to organize the content and presentation of the information returned. The Display Actor simply displays the information to the person that triggered the request. The Information Source Actor shall respond with an error message when it does not support the specific type of request or does not hold any records for the requested patient ID.

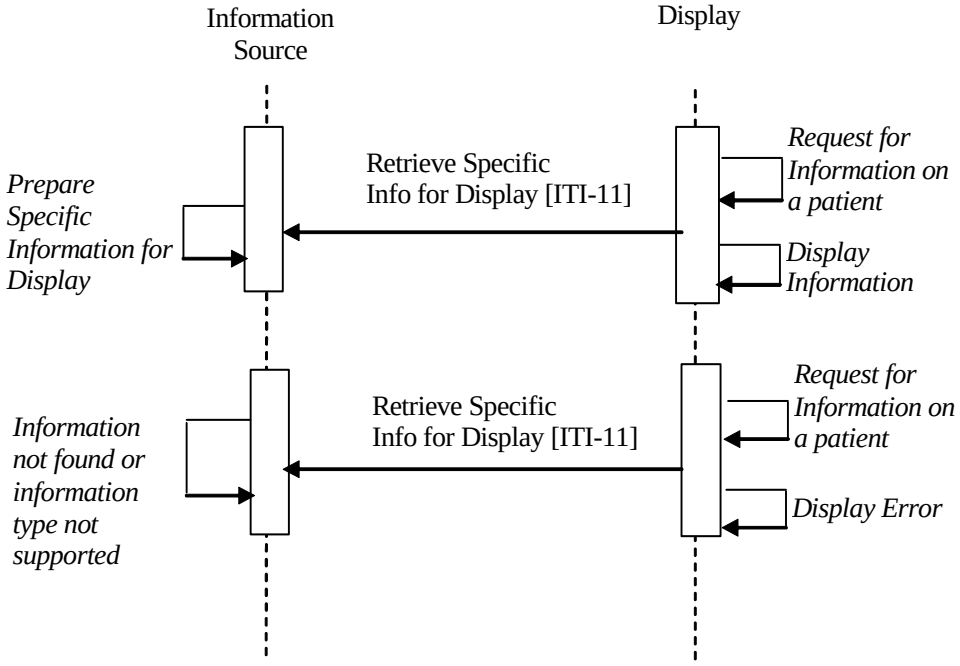


Figure 3.3-1 Case 1: Retrieve Specific Information for Display Process Flow

Case 2 - Retrieve a Document: The second case describes use cases when the Display Actor and the person associated are requesting a uniquely identified document such as a report, an image, an ECG strip, etc. The Information Source Actor responds to the request by using one of the proposed formats to provide the presentation-ready content of the object it manages. The detailed presentation and the clinical integrity of the content of the document are under the control of the Information Source Actor. The Display Actor simply displays the presentation-ready document content to the person that triggered the request. The Information Source Actor shall respond with an error message when the requested document is unknown or when none of the formats acceptable to the Display Actor is suitable to present the requested document.

The main difference between the Retrieve *Specific* Information and the Retrieve *Document* transactions is that the latter applies to a uniquely identifiable persistent object (i.e. retrieving the same document instance at a different point in time will provide the same semantics for its presented content). For the Retrieve Specific Information transaction, this information is always related to a well-identified patient (Patient ID), but its content, although of a specific type (lab summary, or radiology summary, list of allergies) is generally dynamic (i.e. retrieving the same type of specific information at a different point in time is likely to result in different content; for example, a list of allergies may have been updated between two requests).

Note: This Integration profile is not intended for highly dynamic information such as that used for patient monitoring.

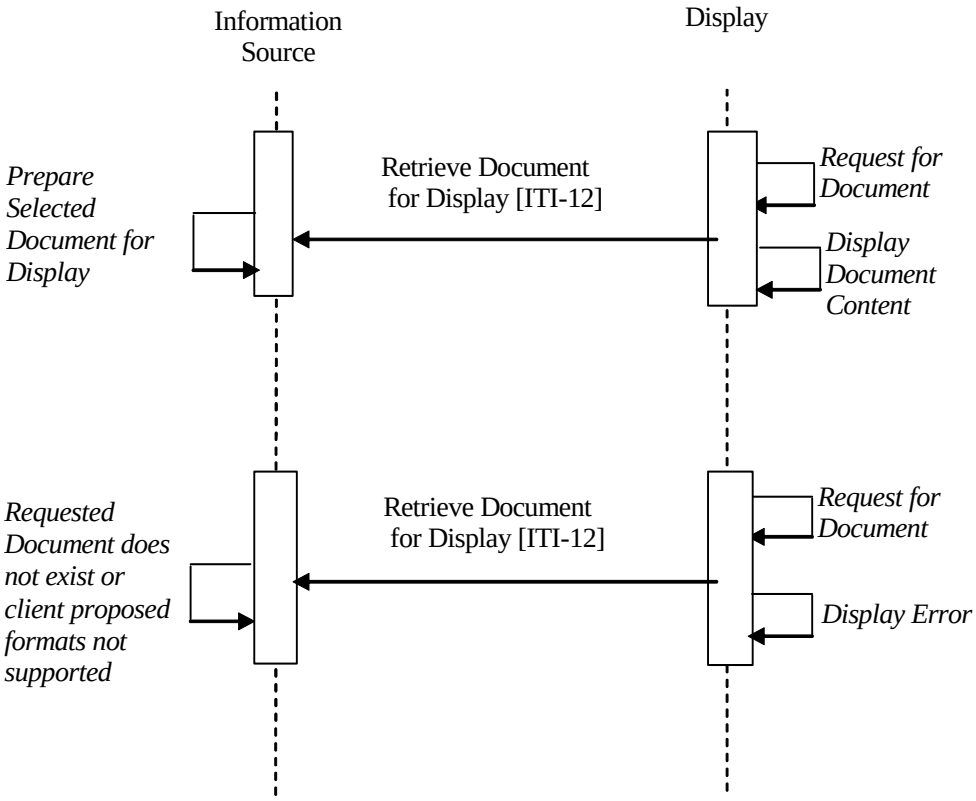
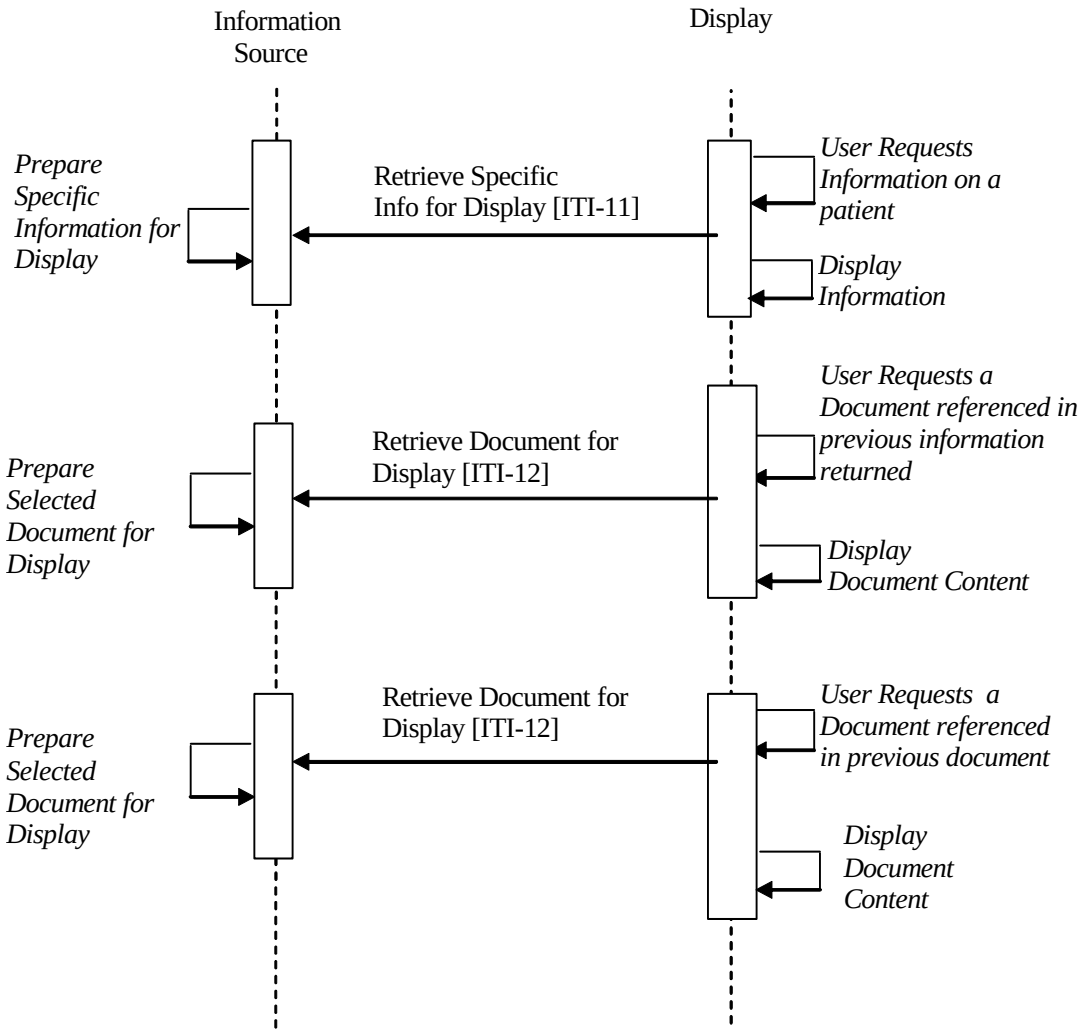


Figure 3.3-2 Case 2: Retrieve a Document Process Flow

Case 3 - Retrieve Specific Information for Display *and* Retrieve several Documents Process Flow: The third case combines the two cases above with the capability to associate in sequence the Retrieve Specific Information and the Retrieve Document for Display transactions. This allows for links to persistent documents within the returned specific information or for having persistent documents reference other persistent documents. For example, the user requests a summary of recent discharge reports, and then selects a specific document referenced in that summary list. From the discharge report displayed to the user, the user selects a specific surgery report. This surgery report is retrieved and displayed.



780

Figure 3.3-3 Case 3: Retrieve Summary Information for Display and Retrieve several Documents Process Flow

The same Display Actor may involve more than one Information Source Actor by sequentially issuing different transactions. This Integration Profile assumes that the Display Actors may be
785 configured a priori with one or more remote Information Source Actors along with the type of retrieve transactions/type of requests/specific keys suitable for the application context from which this Retrieve Information for Display requests are issued. Future Integration Profiles may facilitate such site-specific configuration tasks.

4 Enterprise User Authentication (EUA)

Enterprise User Authentication Profile (EUA) **–** This defines a means to establish one name per user that can then be used on all of the devices and software that participate in this integration profile. It greatly facilitates centralized user authentication management and provides users with the convenience and speed of a single sign-on. This profile leverages Kerberos (RFC 1510) and the HL7 CCOW standard (user subject). User authentication is a necessary step for most application and data access operations and it is a workflow improvement for the users. The IHE EUA Profile adds value to the CCOW specification for the user subject by specifying the user subject and CCOW user subject suffix. This profile does not address security features such as audit trails, access control, authorization management and PKI. Future profiles will be developed to address these security features in a manner complementary to this EUA profile.

The environment is assumed to be a single enterprise, governed by a single security policy and having a common network domain. Unsecured domains -- in particular, Internet access -- are of interest, but not in the scope of this profile. Considerations for applications such as telemedicine and patient remote access to healthcare data are therefore also not in its scope. See Appendix G.

Node and machine authentication is specified in the IHE Basic Security Profile as specified in the IHE Radiology Technical Framework and is not part of this profile.

4.1 Actors/ Transactions

A number of transactions used in this profile conform to the Kerberos v5 standard, defined in RFC 1510. This standard has been stable since 1993, is widely implemented on current operating system platforms, has successfully withstood attacks in its 10-year history, and is fully interoperable among platforms. For example, Sun Solaris, Linux, AIX, HPUX, IBM-z/OS, IBM-OS400, Novell, MAC OS X, and Microsoft Windows 2000/XP all implement Kerberos in an interoperable manner. This is not a complete list; many other vendors also support Kerberos.

For additional detailed information on Kerberos, beyond what is specified in this profile, we suggest these references:

- RFC 1510 - http://www.ietf.org/rfc/rfc1510.txt
- MIT's Kerberos home page - http://web.mit.edu/kerberos/www/
- The Moron's Guide to Kerberos - http://www.isi.edu/~brian/security/kerberos.html
- Microsoft Kerberos information http://www.microsoft.com/TechNet/prodtechnol/windows2000serv/deploy/kerberos.asp

Kerberos implementations are widely available worldwide. Kerberos does include cryptography that may have restricted use laws in some countries. The US export regulations can be found at http://www.bxa.doc.gov/Encryption.

Figure 4.1-1 shows the actors directly involved in the Enterprise User Authentication Profile and the relevant transactions between them. The box labeled "Other IHE Actor" represents actors from other integration profiles that are meant to be grouped with the nearby actor from within this profile. Other actors that may be indirectly involved due to their use of authentication, etc. are not shown.

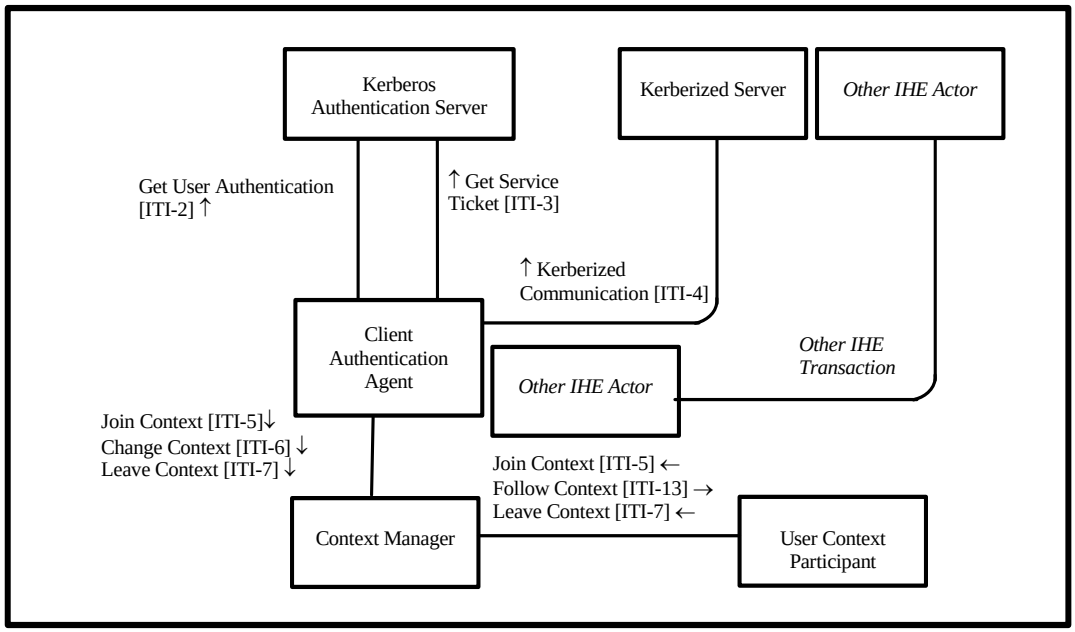


Figure 4.1-1 Enterprise Authentication Actor Diagram

Table 4.1-1 lists the transactions for each actor directly involved in the Enterprise User Authentication Profile. In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled "R"). Transactions labeled "O" are optional. A complete list of options defined in this Integration Profile and that implementations may choose to support is listed in ITI TF-1: 4.2.

Table 4.1-1 Enterprise User Authentication Profile - Actors and Transactions

Actors	Transactions	Optionality	Section in Vol. 2
Kerberos Authentication Server	Get User Authentication [ITI-2]	R	ITI TF-2: 3.2
	Get Service Ticket [ITI-3]	R	ITI TF-2: 3.3
Client Authentication Agent	Get User Authentication [ITI-2]	R	ITI TF-2: 3.2
	Get Service Ticket [ITI-3]	R	ITI TF-2: 3.3
	Kerberized Communication [ITI-4]	R	ITI TF-2: 3.4
	Join Context [ITI-5]	O [Note1]	ITI TF-2: 3.5
	Change Context [ITI-6]	O [Note1]	ITI TF-2: 3.6
	Leave Context [ITI-7]	O [Note1]	ITI TF-2: 3.7
Kerberized Server	Kerberized Communication [ITI-4]	R	ITI TF-2: 3.4
User Context Participant	Join Context [ITI-5]	R	ITI TF-2: 3.5
	Follow Context [ITI-13]	R	ITI TF-2: 3.13
	Leave Context [ITI-7]	R	ITI TF-2: 3.7
Context Manager	Join Context [ITI-5]	R	ITI TF-2: 3.5
	Follow Context [ITI-13]	R	ITI TF-2: 3.13
	Leave Context [ITI-7]	R	ITI TF-2: 3.7
	Change Context [ITI-6]	R	ITI TF-2: 3.6

Note 1: When the Authentication for User Context Option is supported, then the transaction is required.

CCOW facilitates the sharing of the identity of a EUA authentication user but does not provide for the authentication of users. In order for the Context Manager and User Context Participant to participate in the EUA profile it is required that the Client Authentication Agent supports the Authentication for User option. This design provides the User Context Participant with a consistent and enterprise recognized user identity, but does not define access to the Kerberos credentials. Future IHE profiles may address this limitation. Note that the Client Authentication Agent is the key actor when PSA and EUA are combined. See the use case outlined in Section 4.3.2. Applications that implement both the Client Authentication Agent Actor and the User Context Participant Actor shall support configurations where either Actor is disabled.

In any single user environment there shall be only one Client Authentication Agent for one user. In a multi-user environment there shall not be more than one Client Authentication Agent per user.

4.2 Enterprise User Authentication Integration Profile Options

Options that may be selected for this Integration Profile are listed in Table 4.2-1 along with the Actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 4.2-1 Enterprise User Authentication - Actors and Options

Actor	Options	Vol & Section
Kerberos Authentication Server	*No options defined*	- -
Client Authentication Agent	*Authentication for User Context*	ITI TF-2: 3.6
Kerberized Server	*No options defined*	- -
Context Manager	*No options defined*	- -
User Context Participant	*No options defined*	- -

4.3 Enterprise User Authentication Profile Process Flow

4.3.1 Basic User Authentication Process Flow

The following diagram describes the sequence of events in the use of Enterprise User Authentication:

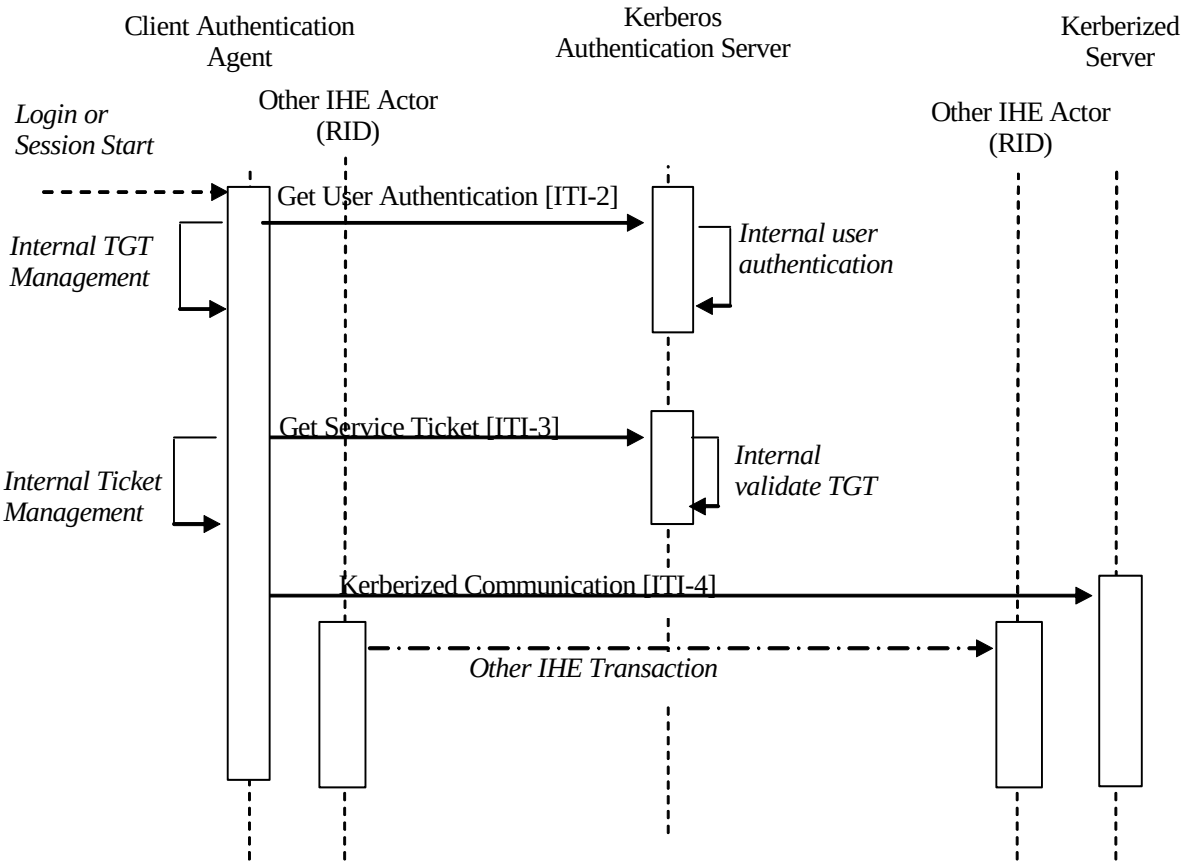


Figure 4.3.1-1. Basic Process Flow in Enterprise User Authentication Profile

The sequence of events in the use of Enterprise User Authentication is:

- The user begins the session. This initiates a local username/password authentication that is converted into the challenge/response system used by Kerberos to avoid transmitting the password over the network. This information is used as part of the Get User Authentication Transaction to get a "Ticket Granting Ticket" (TGT).
- The TGT is saved and managed internally by the Client Authentication Agent Actor. The TGT acts as confirmation that the user has been authenticated.
- For each service that has been Kerberized, the Client Authentication Agent Actor uses the Get Service Ticket Transaction to obtain a service ticket. The service ticket is then used as part of the Kerberized Communication Transaction.

A Kerberized Communication is a Kerberos data exchange that is integrated into another protocol, such as HL7 or DICOM, which is used in another IHE transaction. The details of Kerberization vary and are described separately for the protocols that have been Kerberized. The Kerberization enables the other IHE Actors involved in the other transaction to use the identity of the authenticated user for purposes such as user authorization or audit messages.

The Client Authentication Agent Actor also maintains an internal cache of credentials such as the TGT and service tickets. It renews the tickets as necessary to deal with ticket expirations, re-uses tickets while they are still valid, and removes credentials from the cache when the user session ends. The Client Authentication Agent shall make the Kerberos credentials available using the local operating system mechanisms. Other IHE Actors that need the Kerberos credentials are strongly

encouraged to obtain them using the local operating system mechanisms. Operating system support for ticket management has been implemented and has been defined for various operating systems.

4.3.2 User Authentication with User Synchronized Applications Process Flow

In this use case an application supporting user authentication on the same desktop as another application is synchronized to the same user identity, thus giving the user a single-sign-on experience.

The following diagram describes the sequence of events in the use of User Authentication with User Synchronized Applications:

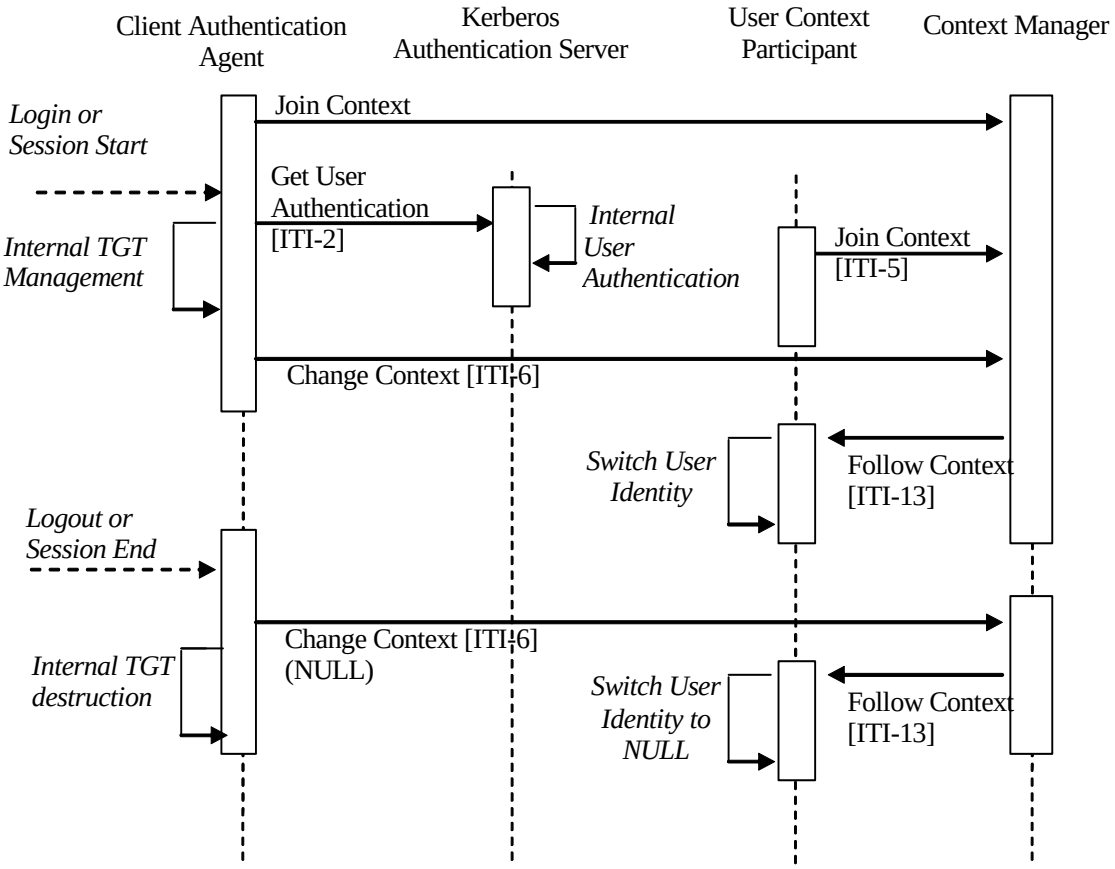


Figure 4.3.2-1 Process Flow with User Synchronized Applications

The sequence of events of the User Authentication with User Synchronized Applications is:

- The user initiates a login by starting the Client Authentication Agent.
- The Client Authentication Agent joins the CCOW user context by sending a Join Context Transaction to the Context Manager Actor. At this point there is no user identity in the context.
- The user provides their username and password to the Client Authentication Agent. This authentication information is converted into the challenge/response system used by Kerberos to avoid transmitting the password over the network. This information is used as part of the Get User Authentication Transaction to get a "Ticket Granting Ticket" (TGT).

- The TGT is saved and managed internally by the Client Authentication Agent Actor. The TGT acts as confirmation that the user has been authenticated.
- A Change Context Transaction is sent to the Context Manager Actor with the users fully qualified user name.
- The user is now logged in to the User Context Participant.
- When the user ends the session, a Change Context Transaction is sent to the Context Manager Actor with a NULL user name.
- The user is logged out of the User Context Participant.

4.3.3 Fast User Switching with Multiple Applications Process Flow

The use model in the clinical environment can be characterized as multiple clinicians using the same workstation for short intervals of time many times a day. In this shared workstation environment the user requires quick access to the patient data contained in the applications. Traditional methods of logging in and out of the workstation at the operating system or network level can take too long and typically force the applications to terminate. This means that the application clients will potentially need to initialize and establish new database connections, introducing further delay to the Clinician access to patient data. The CCOW standard and more specifically the “user” subject provides a means in combination with the Enterprise Authenticator to allow the user to authenticate at the application level and have all of the other applications tune to the new user.

The following diagram describes the sequence of events in the case of Fast User Switching with Multiple Applications:

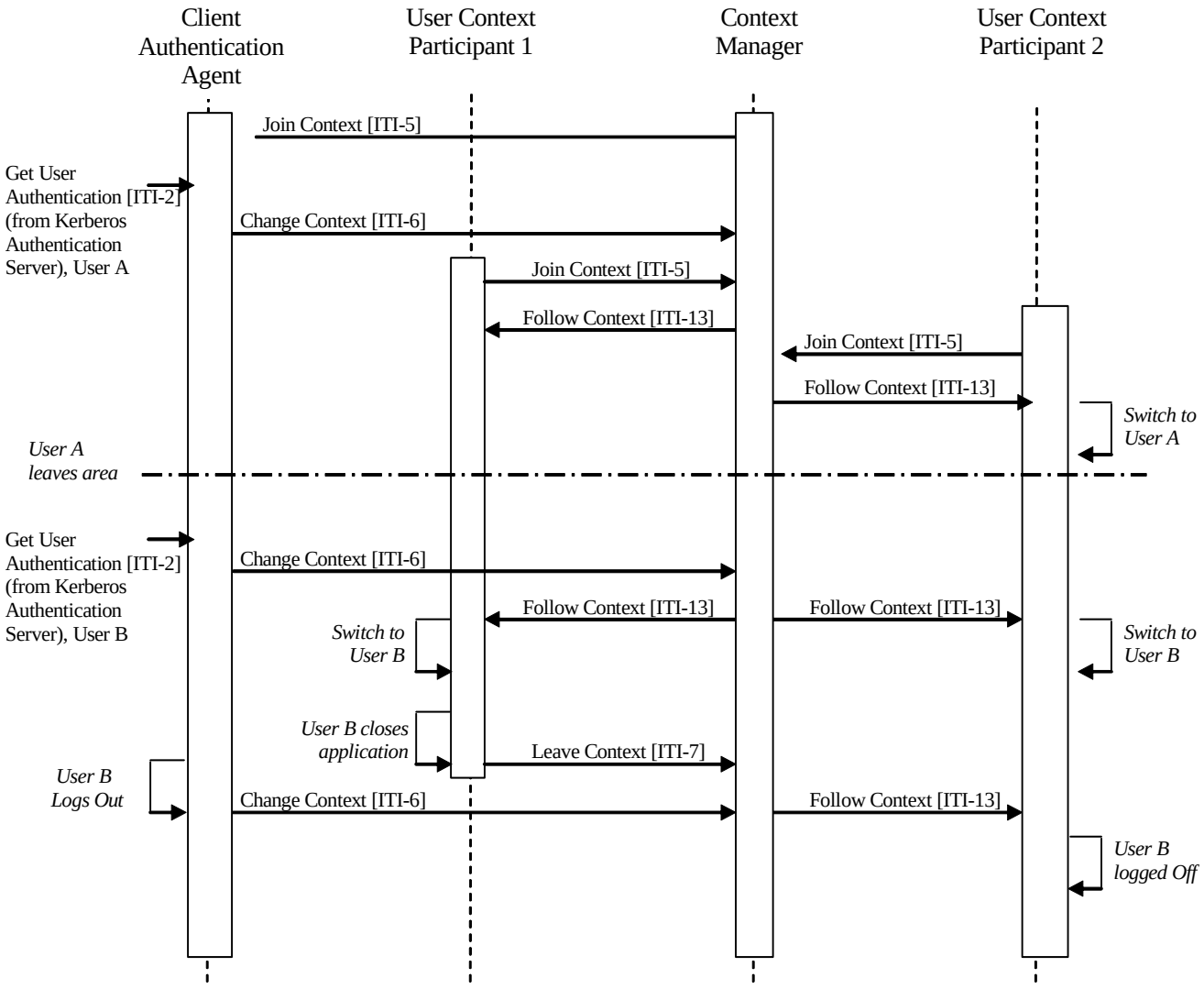


Figure 4.3.3-1. Fast User Switching when using Multiple Applications

The process flow would be similar to the following:

Clinician A launches and authenticates via an application containing the Client Authentication Agent (refer to Figure 4.3.3-1 for details). This actor joins the context session and performs a context change to set Clinician A as the user in context.

Clinician A launches the clinical data repository application, containing a User Context Participant Actor, depicted as User Context Participant 1. The actor joins the context session, gets the current user from the Context Manager, and logs clinician A into the application.

Clinician A launches a cardiology application, containing a User Context Participant Actor, depicted as User Context Participant 2. The actor joins the context session, gets the current user from the Context Manager, and logs clinician A into the application.

Clinician A does his job and then gets called away and leaves the workstation.

Clinician B approaches the workstation and authenticates using the Client Authentication Agent. This results in a context change from Clinician A to Clinician B being set in context without the delay typically associated with a logout and login at the operating system level. The clinical data repository and the cardiology application are notified of the context change by the Context Manager resulting in Clinician A being logged out of both applications and Clinician B being logged into both applications.

Clinician B does his job and then closes the clinical data repository application, which leaves the context prior to terminating the application.

Clinician B is finished reviewing patient data within the cardiology application and logs out using the Client Authentication Agent. This forces a context change to remove the current user from the context, which results in the user being logged out of the cardiology application.

5 Patient Identifier Cross-referencing (PIX)

The ***Patient Identifier Cross-referencing Integration Profile (PIX)*** is targeted at healthcare enterprises of a broad range of sizes (hospital, a clinic, a physician office, etc.). It supports the cross-referencing of patient identifiers from multiple Patient Identifier Domains via the following interactions:

- The transmission of patient identity information from an identity source to the Patient Identifier Cross-reference Manager.
- The ability to access the list(s) of cross-referenced patient identifiers either via a query/ response or via update notification.

By specifying the above transactions among specific actors, this integration profile does not define any specific enterprise policies or cross-referencing algorithms. By encapsulating these behaviors in a single actor, this integration profile provides the necessary interoperability while maintaining the flexibility to be used with any cross-referencing policy and algorithm as deemed adequate by the enterprise.

The following diagram shows the intended scope of this profile (as described above).

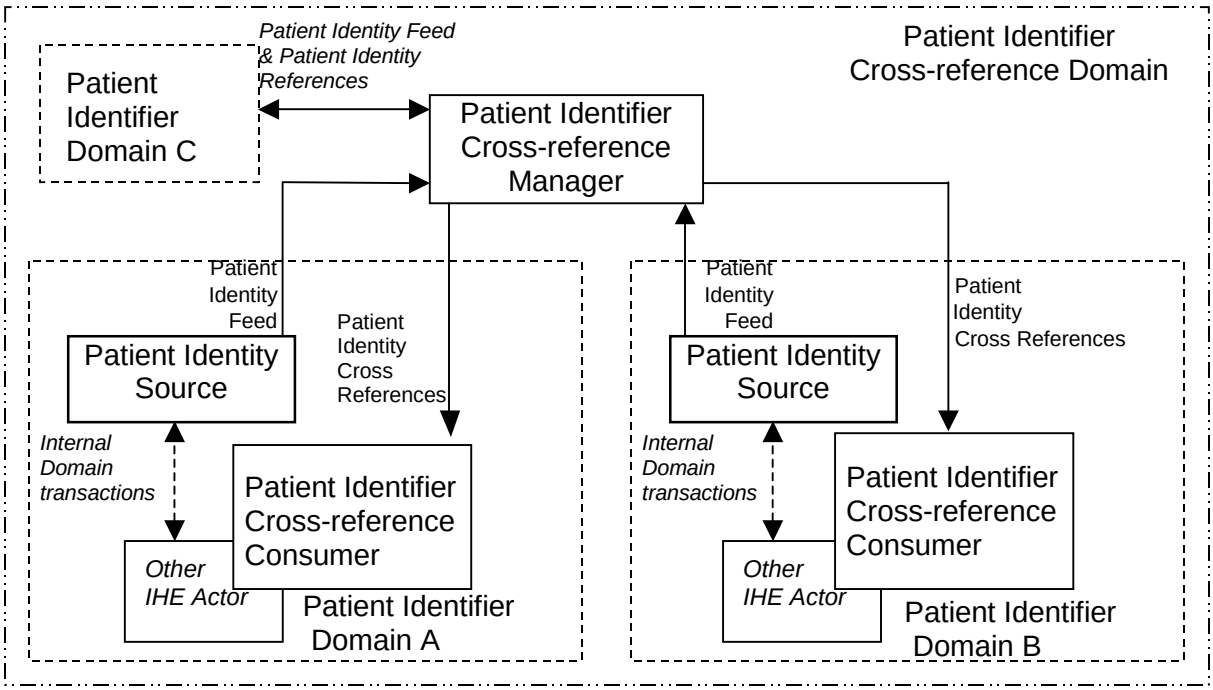


Figure 5-1 Process Flow with Patient Identifier Cross-referencing

The diagram illustrates two types of Identifier Domains: a Patient Identifier Domain and a Patient Identifier Cross-reference Domain.

A Patient Identifier Domain is defined as a single system or a set of interconnected systems that all share a common identification scheme (an identifier and an assignment process to a patient) and issuing authority for patient identifiers. Additionally, a Patient Identifier Domain has the following properties:

- A set of policies that describe how identities will be defined and managed according to the specific requirements of the domain.
- An administration authority for administering identity related policies within the domain.
- A **single** system, known as a patient identity source system, that assigns a unique identifier to each instance of a patient-related object as well as maintaining a collection of identity traits.
- Ideally, only one identifier is uniquely associated with a single patient within a given Patient Identifier Domain, though a single Patient Identity Source Actor may assign multiple identifiers to the same patient and communicate this fact to the Patient Identifier Cross-reference Manager. For a description of how the Patient Identifier Cross-reference Manager Actor responds to requests for a list of cross-referenced identifiers that include these "duplicates" see ITI TF-2: 3.9.4.2.2.6).
- An "Identifier Domain Identifier" (known as assigning authority) that is unique within a Patient Identifier Cross-reference Domain.
- Other systems in the Patient Identifier Domain rely upon the identifiers assigned by the patient identity source system of the domain to which they belong.

A Patient Identifier Cross-reference Domain consists of a set of Patient Identifier Domains known and managed by a Patient Identifier Cross-reference Manager Actor. The Patient Identifier Cross-reference Manager Actor is responsible for creating, maintaining and providing lists of identifiers that are aliases of one another across different Patient Identifier Domains.

The Patient Identifier Cross-reference Domain embodies the following assumptions about agreement within the group of individual Identifier Domains:

- They have agreed to a set of policies that describe how patient identities will be cross-referenced across participating domains;
- They have agreed to a set of processes for administering these policies;
- They have agreed to an administration authority for managing these processes and policies.

All these assumptions are critical to the successful implementation of this profile. This integration profile imposes minimal constraints on the participating Patient Identifier Domains and centralizes most of the operational constraints for the overall Patient Identification Cross-reference Domain in the Patient Identifier Cross-reference Manager Actor. If the individual Identifier Domains cannot agree to the items outlined above, implementation of this profile may not provide the expected results.

The Patient Identifier Cross-reference Manager Actor is not responsible for improving the quality of identification information provided to it by the Identity Source Actors. It is assumed that the Identity Source actors are responsible for providing high quality data to the Patient Identifier Cross-reference Manager. For example, the Patient Identifier Cross-reference Manager Actor is NOT responsible to provide a single reference for patient demographics. The intent is to leave the responsibility for the quality and management of its patient demographics information and the integrity of the identifiers it uses within each Patient Identity Domain (Source actors). When receiving reports and displays from multiple PIX domains, it is inevitable that some of those reports and displays will have inconsistent names.

The Patient Identifier Cross-reference Consumer may use either a query for sets of cross-reference patient identifiers or use both a notification about cross-reference changes and a query transaction. In the case of using a notification, the Patient Identifier Cross-reference Consumer may also use the PIX Query Transaction to address situations where the Patient Identifier Cross-reference Consumer may be out of synch with the Patient Identifier Cross-reference Manager. This Integration Profile does not specify the consumer policies in using the PIX Query Transaction (ITI TF-2: 3.9).

For a discussion of the relationship between this Integration Profile and an enterprise master patient index (eMPI) see Section 5.4.

5.1 Actors/ Transactions

Figure 5.1-1 shows the actors directly involved in the Patient Identifier Cross-referencing Integration Profile and the relevant transactions between them. Other actors that may be indirectly involved due to their participation in other related profiles are not shown.

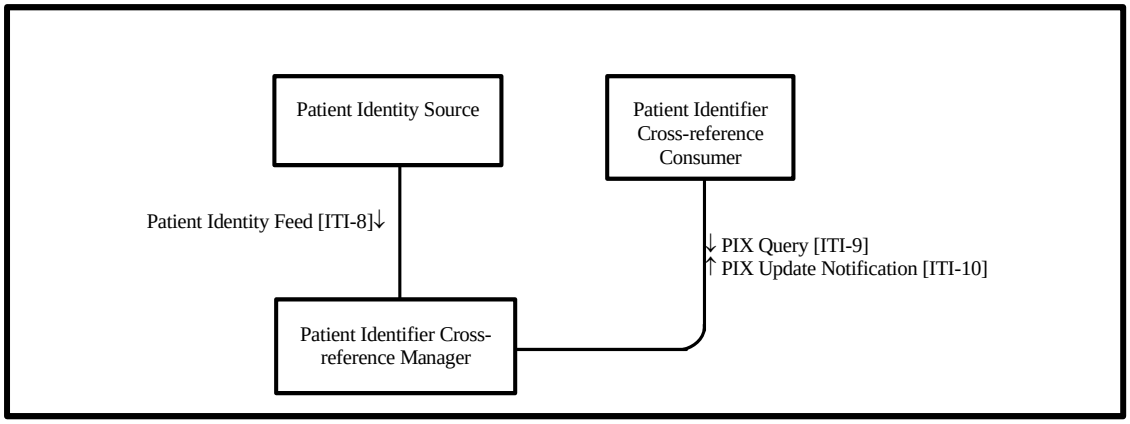


Figure 5.1-1 Patient Identifier Cross-referencing Actor Diagram

Table 5.1-1 lists the transactions for each actor directly involved in the Patient Identifier Cross-referencing Profile. In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled "R"). Transactions labeled "O" are optional. A complete list of options defined by this Integration Profile and that implementations may choose to support is listed in the ITI TF-1: 5.2.

Table 5.1-1 Patient Identifier Cross-referencing Integration for MPI Profile - Actors and Transactions

Actors	Transactions	Optionality	Section in Volume 2
Patient Identity Source	Patient Identity Feed[ITI-8]	R	ITI TF-2: 3.8
Patient Identifier Cross-reference Consumer	PIX Query[ITI-9]	R	ITI TF-2: 3.9
	PIX Update Notification[ITI-10]	O	ITI TF-2: 3.10
Patient Identifier Cross-reference Manager	Patient Identity Feed[ITI-8]	R	ITI TF-2: 3.8
	PIX Query[ITI-9]	R	ITI TF-2: 3.9
	PIX Update Notification[ITI-10]	R	ITI TF-2: 3.10

5.2 Patient Identifier Cross-referencing Integration Profile Options

Options that may be selected for this Integration Profile are listed in the Table 5.2-1 along with the Actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 5.2-1 Patient Identifier Cross-referencing - Actors and Options

Actor	Options	Vol & Section
Patient Identity Source	*No options defined*	- -
Patient Identifier Cross-reference Manager	*No options defined*	- -
Patient Identifier Cross-reference Consumer	*PIX Update Notification*	ITI TF-2: 3.10

5.3 Patient Identifier Cross-referencing Profile Process Flows

The following sections describe use cases that this profile addresses.

5.3.1 Use Case: Multiple Identifier Domains within a Single Facility/ Enterprise

A clinician in the Intensive Care Unit at General Hospital is reviewing a patient chart on the Intensive Care information system and wishes to review or monitor the patient's glucose level, which is included in a laboratory report stored in the hospital's main laboratory system. The Intensive Care system needs to map its own patient ID, which it generates internally, to the patient's medical record number (MRN), which is generated from the hospital's main ADT system and is used as the patient identity by the lab system. In this case the Intensive Care system is essentially in a different identifier domain than the rest of the hospital since it has its own notion of patient identity.

In this scenario, the hospital's main ADT system (acting as a Patient Identity Source) would provide a Patient Identity Feed (using the patient's MRN as the identifier) to the Patient Identifier Cross-reference Manager. Similarly, the Intensive Care system would also provide a Patient Identity Feed to the Patient Identifier Cross-reference Manager using the internally generated patient ID as the patient identifier and providing its own unique identifier domain identifier.

Once the Patient Identifier Cross-reference Manager receives the Patient Identity Feed transactions, it performs its internal logic to determine which, if any, patient identifiers can be "linked together" as being the same patient based on the corroborating information included in the Feed transactions it has received. The cross-referencing process (algorithm, human decisions, etc.) is performed within the Patient Identifier Cross-reference Manager and is outside the scope of IHE. (See ITI TF-2: 3.9.4.2.2.6 for a more complete description of the scope of the cross-referencing logic boundary).

The Intensive Care system wants to get lab information associated with a patient that the Intensive Care system knows as patient ID = 'MC-123'. It requests the lab report from the lab system using its own patient ID (MC-123) including the domain identifier/ assigning authority. Upon receipt of the request, the lab system determines that the request is for a patient outside of its own identifier domain (ADT Domain). It requests a list of patient ID aliases corresponding to patient ID = 'MC-123' (within the "Intensive Care domain") from the Patient Identifier Cross-reference Manager. Having linked this patient with a patient known by medical record number = '007' in the 'ADT Domain', the Patient Identifier Cross-reference Manger returns this list to the lab system so that it may retrieve the lab report for the desired patient and return it to the Intensive Care system. Figure 5.3-1 illustrates this process flow.

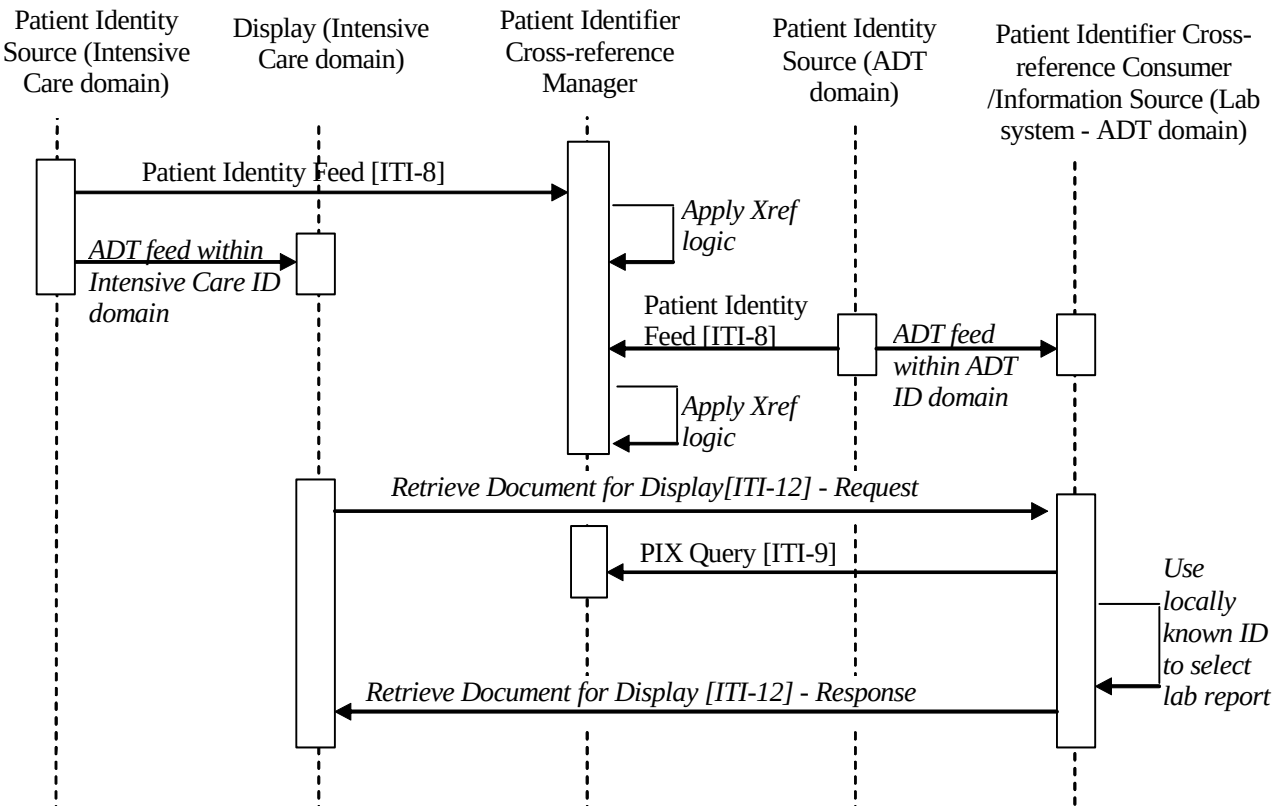


Figure 5.3-1. Multiple ID Domains in a Single Facility Process Flow in PIX Profile

Note: Request and Response portions of the Retrieve Document for Display transaction are not part of this profile and included for illustration purposes only.

5.3.2 Use Case: Multiple ID Domains Across Cooperating Enterprises

A healthcare enterprise is established by the consolidation of two hospitals, each having its own separate patient registration process run by different hospital information systems. When a patient is treated in one hospital, the access to its electronic records managed by the other hospital is necessary. The following use case illustrates this scenario.

Hospitals A and B have been consolidated and have a single Patient Identifier Cross-reference Manager that maintains the ID links between the two hospitals. Each hospital has a different HIS that is responsible for registering patients, but they have consolidated their cardiology information systems. The cardiology system has been configured with a Patient Identifier Cross-reference Consumer to receive patient identity notifications when cross-referencing activity occurs.

A patient is registered and then has some diagnostic stress tests done at hospital A. The cardiology information system queries the Patient Identifier Cross-reference Manager to get a list of possible ID aliases for the patient to see if any past cardiology reports may be available. No patient ID aliases are found. Some time later the same patient goes to hospital B to have a second diagnostic stress test done. The patient is registered via the HIS in hospital B which then sends that identity information to the Patient Identifier Cross-reference Manager. The Patient Identifier Cross-reference Manager determines this is in fact the same patient as was registered previously at

hospital A. The cardiology information system was previously configured with the Patient Identifier Cross-reference Manager to receive notifications, thus a notification is sent to the cardiology system to inform it of the patient identifier aliases. This notification is done to allow systems that are aware of multiple identifier domains to maintain synchronization with patient identifier changes that occur in any of the identifier domains that they are aware of.

Figure 5.3-2 illustrates the process flow for this use case.

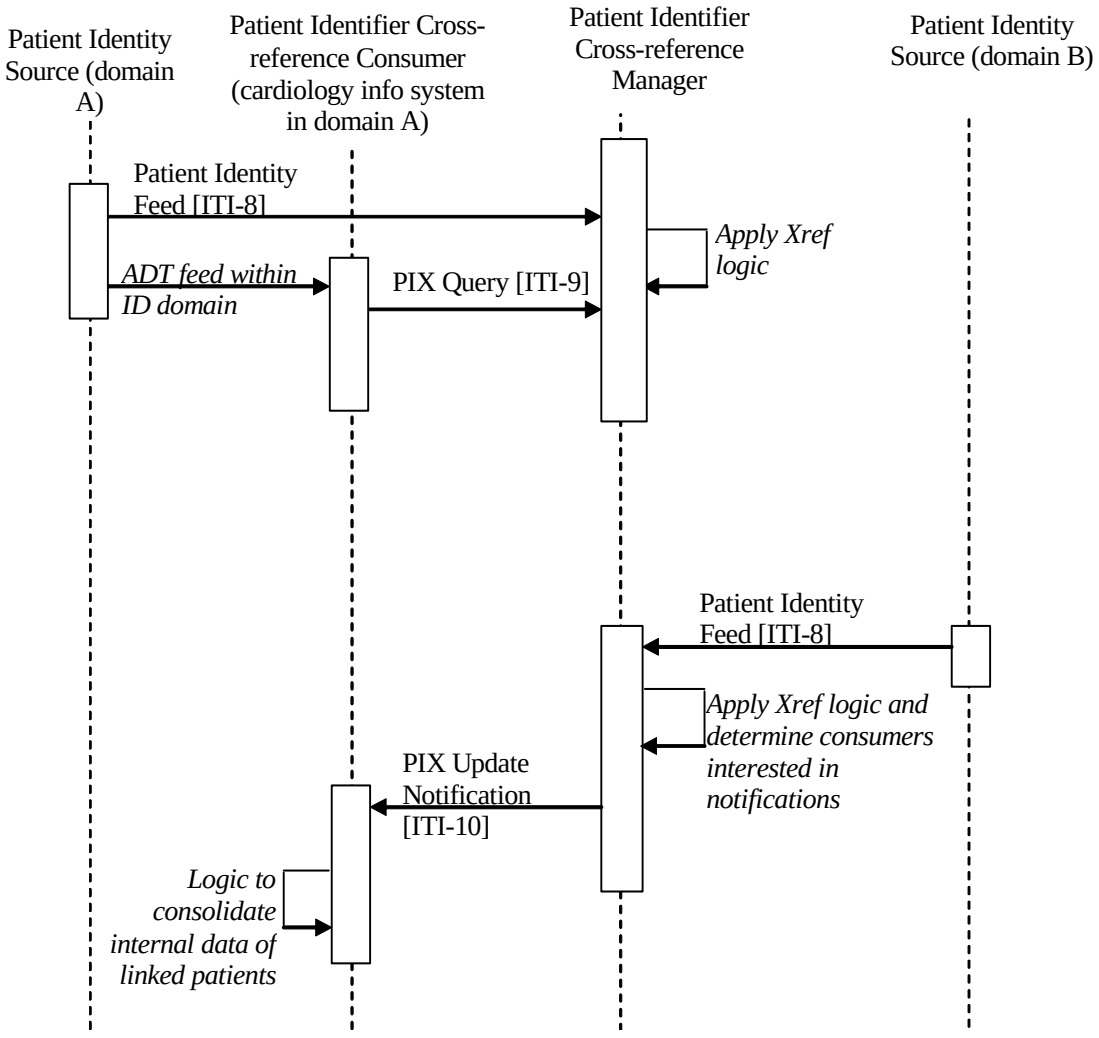


Figure 5.3-2 Multiple ID Domains Across Cooperating Enterprises Process Flow in PIX Profile

Note: PIX Update Notifications are not sent for the first Patient Identity Feed for a patient, since no cross-referencing activity occurred after this first Patient Identity Feed Transaction.

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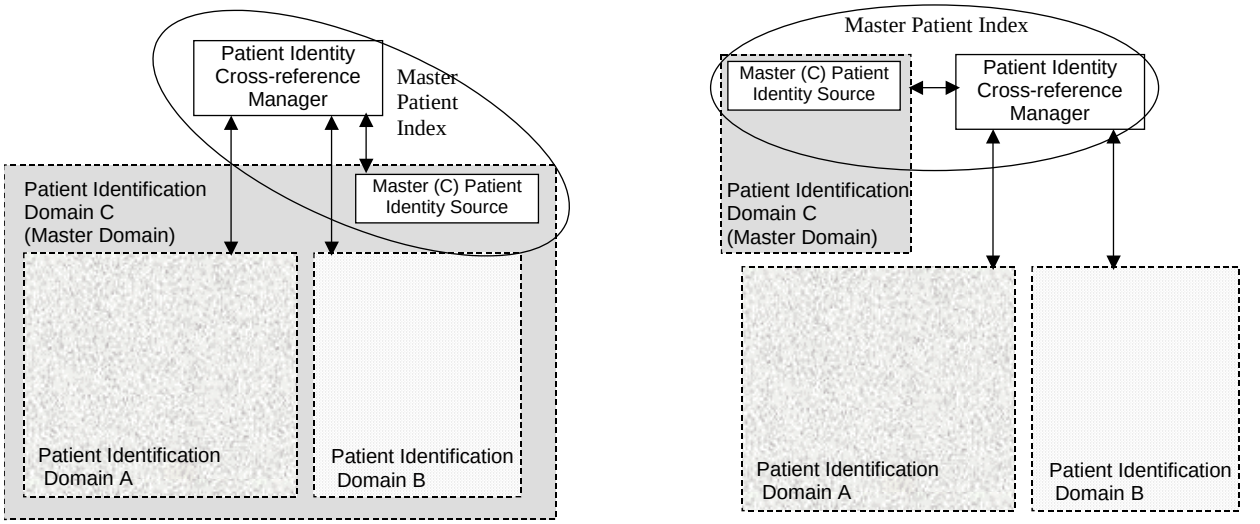
5.4 Relationship between the PIX Integration Profile and eMPI

The PIX Integration Profile achieves the integration of disparate Patient Identifier Domains by using a cross-referencing approach between Patient Identifiers associated with the same patient. This section discusses how this approach is compatible with environments that wish to establish

master patient identifiers (MPI), or enterprise MPI (eMPI) systems. An eMPI may be considered a particular variation in implementation of the PIX Integration Profile.

The concept of an MPI is a rather broad concept, yet it is most often associated with the creation of a master patient identifier domain. Such a master domain is considered more broadly applicable or more "enterprise-level" than the other patient identifier domains it includes. Such a hierarchical inclusion of patient identification domains into a "master patient identification domain" can be considered a particular case of patient cross-reference, where the patient identifiers in the various domains are cross-referenced to the patient identifiers of the master domain. Two possible configurations are depicted by Figure 5.4-1.



Two domains included in a Master Patient Index

The same configuration represented as 3 cross-referenced domains

Figure 5.4-1 PIX Profile Relationship to eMPI

Figure 5.4-1 above shows how the Master Patient Identifier Domain (Domain C), in a typical MPI approach, is simply another patient Identification Domain when considered in a Cross-referencing approach. The decision to place enterprise-wide systems such as Clinical Data Repositories into the so-called master domain is simply a configuration choice. In addition, such a configuration sometimes assumes that any system in Patient Domain A not only manages the patient Identifiers of Domain A but is also aware of those of Domain C. In the Patient Identifier Cross-reference Integration Profile, this is a configuration choice where certain systems have been designed and configured to operate across multiple domains. Thus the entity often called an MPI (shown by the oval) is actually the combination of a Patient Identity Source Actor (ADT) along with a Patient Identifier Cross-reference Manager.

The PIX Integration Profile can coexist with environments that have chosen to deploy a distinct MPI, and provides a more scalable approach. Many other configurations can also be deployed, in particular those where the creation of a master domain "including" the other domains is not necessary (i.e., a simple federation of domains where none is actually the master).

6 Patient Synchronized Applications (PSA)

The ***Patient Synchronized Applications Profile (PSA)*** enables single patient selection for the user working in multiple applications on a workstation desktop. With this Integration Profile patient selection in any of the applications causes all other applications to tune to that same patient. This allows a clinician to use the application they are most familiar with to select the patient and have that selection reflected in the other applications they are using follow along.

This profile leverages the HL7 CCOW standard, specifically for patient subject context management. The scope of this profile is for sharing of the CCOW Patient subject only. The IHE PSA profile adds value to the CCOW specification for the patient subject by further constraining the patient identifier to ensure consistency across applications supporting PSA, providing guidance for consistent behavior across applications supporting PSA and ensuring consistent interaction with the Patient Identifier Cross-reference Consumer Actor across the enterprise.

For applications that require user authentication, IHE recommends implementation of the Enterprise User Authentication Profile, as opposed to other means, such as a CCOW Authentication Repository. ITI TF-1: 4 describes the Enterprise User Authentication Profile and the use of the CCOW user subject.

6.1 Actors/ Transactions

Figure 6.1-1 shows the actors directly involved in the Patient Synchronized Applications Integration Profile and the relevant transactions between them. Other actors that may be indirectly involved due to their participation in other profiles are not shown.

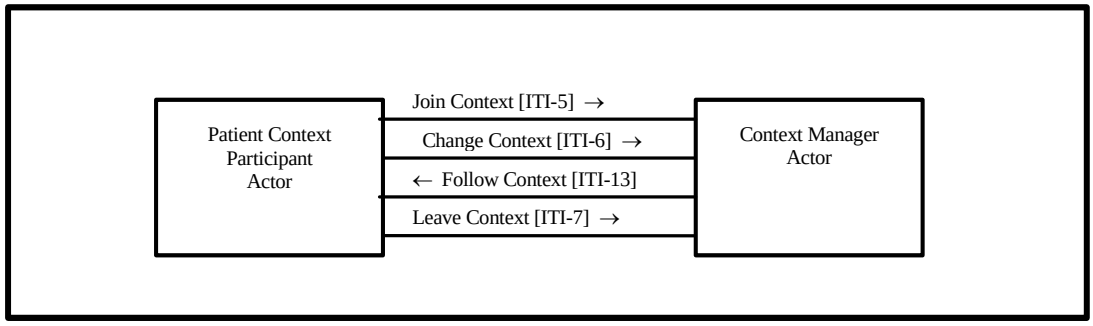


Figure 6.1-1 Patient Synchronized Applications Profile Actor Diagram

Table 6.1-1 lists the transactions for each actor directly involved in the PSA Profile. In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled “R”).

The Patient Context Participant Actor shall support all four transactions identified in Figure 6.1-1 as defined in ITI TF-2. The Patient Context Participant Actor shall respond to all patient context changes. This actor shall set the patient context provided the application has patient selection capability.

The IHE Context Manager Actor may encompass more than a CCOW context manager function. It may include a number of other components such as the context management registry and patient mapping agent.

The Context Manager Actor may be grouped with a Patient Identifier Cross-referencing (PIX) Consumer Actor of the Patient Identity Cross-referencing Profile; see ITI TF-2: Appendix D for a description of the additional responsibilities placed on the Context Manager Actor in this case.

Table 6.1-1 Patient Synchronized Applications Integration Profile - Actors and Transactions

Actors	Transactions	Optionality	Section
Patient Context Participant	Join Context [ITI-5]	R	ITI TF-2: 3.5
	Change Context [ITI-6]	R	ITI TF-2: 3.6
	Leave Context [ITI-7]	R	ITI TF-2: 3.7
	Follow Context [ITI-13]	R	ITI TF-2: 3.13
Context Manager	Join Context [ITI-5]	R	ITI TF-2: 3.5
	Change Context [ITI-6]	R	ITI TF-2: 3.6
	Leave Context [ITI-7]	R	ITI TF-2: 3.7
	Follow Context [ITI-13]	R	ITI TF-2: 3.13

6.2 Patient Synchronized Applications Integration Profile Options

Options that may be selected for this Integration Profile are listed in Table 6.2-1 along with the actors to which they apply. Dependencies between options, when applicable, are specified in notes.

Table 6.2-1 Patient Synchronized Applications - Actors and Options

Actor	Options	Vol & Section
Patient Context Participant	*No options defined*	- -
Context Manager	*No options defined*	- -

6.3 Patient Synchronized Applications Integration Profile Process Flows

The Patient Synchronized Applications Integration Profile provides maximum value when a user needs to use more than one application simultaneously. The process flow outlined in Section 6.3.1 depicts a use case where the applications only participate in the PSA profile. The process flow outlined in ITI TF-1: Appendix E illustrates when the PSA and Enterprise User Authentication (EUA) profiles are deployed together.

6.3.1 Use Case: Simple Patient Switching

When the PSA profile is not grouped with EUA profile only the patient identity is passed in context. This use case does not explicitly identify the method of user authentication, as it may not be required by the application or may be accomplished by other means. In this use case both applications share the same patient identifier domain. The process flow for this use case is:

The clinician launches the clinical data repository application, depicted as Patient Context Participant Actor 1. The clinical data repository application joins the context session for the clinician desktop.

The clinician selects patient A in the clinical data repository application. The clinical data repository application sets the identifier for patient A in context.

The clinician launches a cardiology application, depicted as Patient Context Participant Actor 2. The Cardiology application joins the context session, gets the identifier for patient A from context, and tunes its display to patient A.

The clinician selects patient B in the cardiology application. This action results in the initiation of a Change Context transaction by the cardiology application (Patient Context Participant Actor 2). All non-instigating applications participate via the Follow Context transaction, which results in the selected patient being displayed in the clinical data repository application (Patient Context Participant Actor 1).

The clinician closes the clinical data repository application. The clinical data repository application leaves the context prior to terminating the application.

The clinician closes the cardiology application. The cardiology application leaves the context prior to terminating the application.

Figure 6.3-1 illustrates the process flow for this use case.

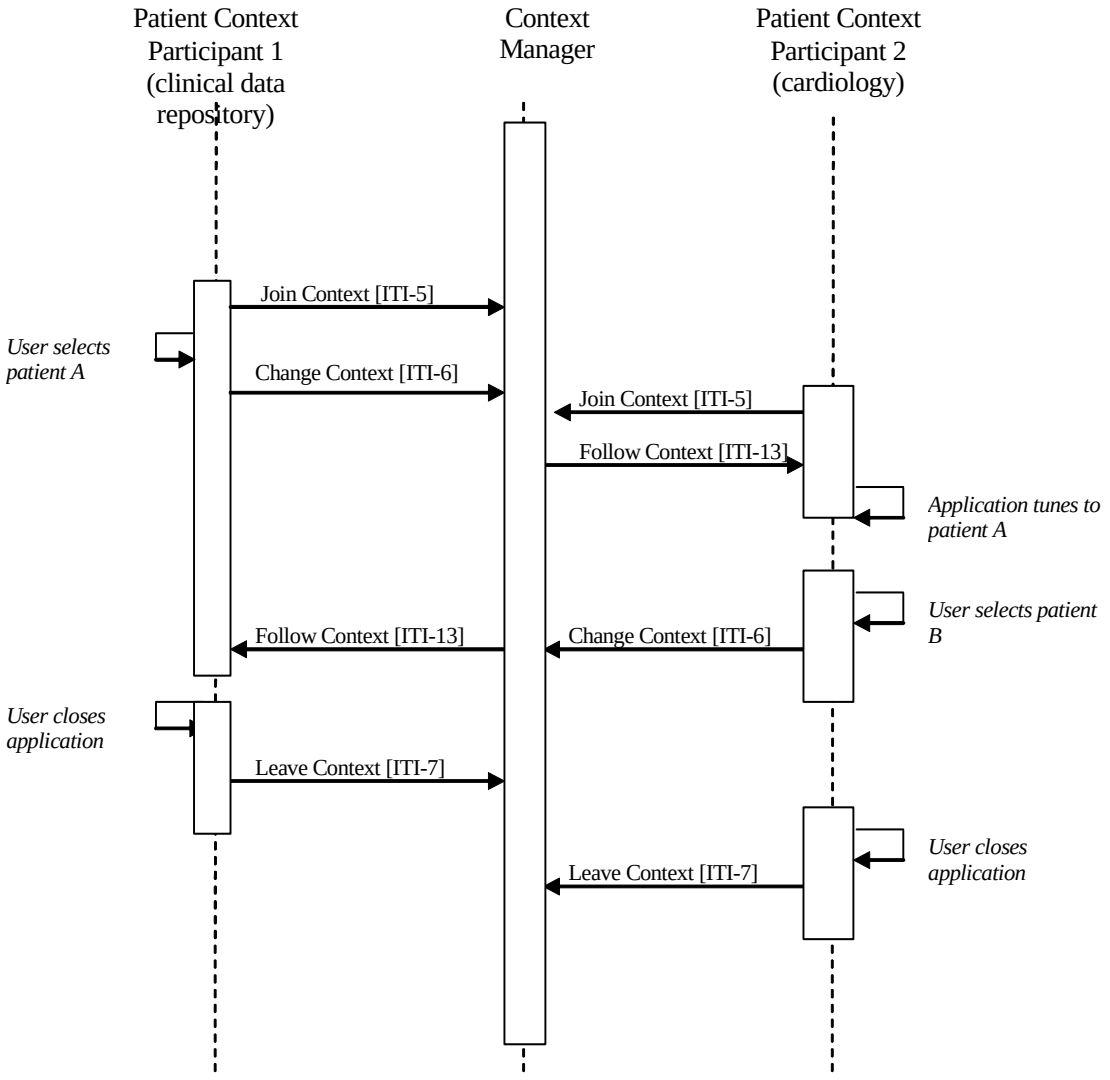


Figure 6.3-1 Simple Patient Switching Process Flow

7 Consistent Time (CT)

The ***Consistent Time Integration Profile (CT)*** provides a means to ensure that the system clocks and time stamps of the many computers in a network are well synchronized. This profile specifies synchronization with a median error less than 1 second. This is sufficient for most purposes.

The Consistent Time Integration Profile defines mechanisms to synchronize the time base between multiple actors and computers. Various infrastructure, security, and acquisition profiles require use of a consistent time base on multiple computers. The Consistent Time profile requires the use of the Network Time Protocol (NTP) defined in RFC 1305. When the Time Server is grouped with a Time Client to obtain time from a higher tier Time Server, the Time Client shall utilize NTP. For some Time Clients that are not grouped with a Time Server, SNTP may be usable.

This profile was previously a portion of the Radiology Basic Security Profile, but it has a variety of other infrastructure uses.

Note: This profile corresponds to a portion of the IHE Radiology Technical Framework, Basic Security Profile. It is required by more than just radiology systems. It is needed by several of the profiles in the IHE IT Infrastructure and will also be needed by Cardiology. It is therefore being re-located from IHE Radiology into IHE IT Infrastructure. There are no changes to the requirements, so actors that supported the Radiology Basic Secure Node or Time Server do not need modification. The Maintain Time [RAD TF-3: 4.33] transaction from Radiology and the Maintain Time [ITI TF-2: 3.1] transaction for IT Infrastructure are the same.

7.1 Actors/ Transactions

Figure 7.1-1 shows the actors directly involved in the Consistent Time Profile and the relevant transactions between them. Other actors that may be indirectly involved because of their participation in profiles that require consistent time are not shown.

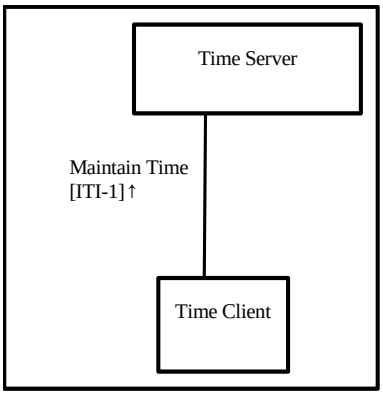


Figure 7.1-1: Consistent Time Profile Actor Diagram

Table 7.1-1 lists the transactions for each actor directly involved in the Consistent Time Integration Profile. In order to claim support of this integration profile, an implementation must perform the required transactions (labeled “R”).

Table 7.1-1: Consistent Time - Actors and Transactions

Actors	Transactions	Optionality	Section in

			Vol. 2
Time Server	Maintain Time [ITI-1]	R	ITI TF-2: 7.1
Time Client	Maintain Time [ITI-1]	R	ITI TF-2: 7.1

7.2 Consistent Time Integration Options

Options that may be selected for this integration profile are listed in the Table 7.2-1 along with the actors to which they apply.

Table 7.2-1: Consistent Time - Actors and Options

Actor	Options	Vol & Section
Time Server	*Secured NTP*	ITI TF-2: 3.1.4-1
Time Client	*SNTP, Secured NTP*	ITI TF-2: 3.1.4-1

7.3 Consistent Time Process Flow

This section describes the typical flow related to the Consistent Time Profile. In the process flow diagram 7.3-1, the Time Client B and Time Server B have been grouped. When a Client and Server are grouped they utilize internal communications mechanisms to synchronize their time.

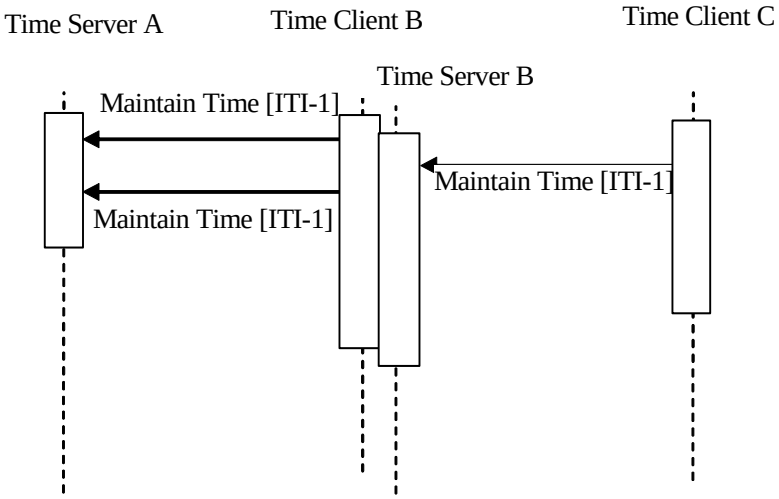


Figure 7.3-1 Basic Process Flow in Consistent Time Profile

The Time Client B maintains time synchronization with the Time Server A. The Time Server B is internally synchronized with Time Client B. The Time Client C maintains time synchronization with Time Server B.

The NTP protocol has been designed to provide network time services for synchronization with this kind of cascaded synchronization. The achievable accuracy is dependent on specific details of network hardware and topology, and on details of computer hardware and software implementation. The Time Server and Time Client are grouped to provide synchronization cascading and reduce network traffic.

8 Patient Demographics Query (PDQ)

8.1 Actors/ Transactions

Figure 8.1-1 shows the actors directly involved in the Patient Demographics Query Integration Profile and the relevant transactions between them. Other actors that may be indirectly involved due to their participation in Patient ID Cross-referencing, etc. are not necessarily shown.

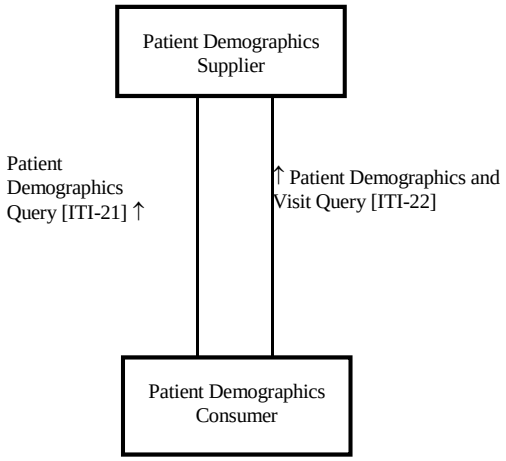


Figure 8.1-1. Patient Demographics Query Profile Actor Diagram

Table 8.1-1 lists the transactions for each actor directly involved in the Patient Demographics Query Profile. In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled "R"). Transactions labeled "O" are optional. A complete list of options defined by this Integration Profile and that implementations may choose to support is listed in Section 8.2.

Table 8.1-1. Patient Demographics Query Integration Profile - Actors and Transactions

Actors	Transactions	Optionality	Section in Vol. 2
Patient Demographics Consumer	Patient Demographics Query	R	ITI TF-2: 3.21
	Patient Demographics and Visit Query	O	ITI TF-2: 3.22
Patient Demographics Supplier	Patient Demographics Query	R	ITI TF-2: 3.21
	Patient Demographics and Visit Query	O	ITI TF-2: 3.22

8.2 Patient Demographics Query Integration Profile Options

Options that may be selected for this Integration Profile are listed in the table 8.2-1 along with the actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 8.2-1 Patient Demographics Query - Actors and Options

Actor	Options	Vol & Section
Patient Demographics Consumer	*Patient Demographics and Visit Query*	ITI TF-2: 3.22
Patient Demographics Supplier	*Patient Demographics and Visit Query*	ITI TF-2: 3.22

8.3 Patient Demographics Query Process Flow

The Patient Demographics Supplier performs the following functions.

- It receives patient registration and update messages from other systems in the enterprise (*e.g.*, ADT Patient Registration systems), which may or may not represent different Patient ID Domains. The method in which the Patient Demographics Supplier obtains the updated patient demographic information is not addressed by this profile.
- It responds to queries for information.

Specific methods for acquiring demographic information are beyond the scope of this Profile. It is a prerequisite that the Patient Demographics Supplier possess current demographic information. One method by which current demographic information may be obtained is for the Patient Demographic Supplier to be grouped with another IHE actor, such as Order Filler, that either maintains or receives such information.

In all cases, the Patient Demographics Supplier receives a Patient Demographics Query or Patient Demographics and Visit Query request from the Patient Demographics Consumer, and returns demographics (and, where appropriate, visit) information from the single domain that is associated with the application to which the query message is sent. Identifier information may be returned from multiple or single domains; see the "Using Patient Data Query (PDQ) in a Multi-Domain Environment" section (ITI TF-2: Appendix M) for a discussion of the architectural issues involved.

Use Case 1: Patient Information Entering at Bedside

An admitted patient is assigned to a bed. The patient may or may not be able to provide positive ID information. The nurse needs to enter patient identity information into some bedside equipment to establish the relationship of the assigned bed to the patient. The equipment issues a query for a patient pick list to a patient demographics supplier that provides data for a patient pick list. Search criteria entered by the nurse might include one or more of the following:

- Partial or complete patient name (printed on the patient record or told by the patient)
- Patient ID (this may be obtained from printed barcode, a bed-side chart, etc.)
- Partial ID entry or scan.
- Date of birth / age range
- Bed ID

The system returns a list of patients showing the MRN, full name, age, sex, room/bed, and admit date, and displays the list to the nurse. The nurse then selects the appropriate record to enter the patient identity information into the bedside equipment application.

Use Case 2: Patient Identity Information Entering in Physician Offices

A patient visits a physician office for the first time. The nurse needs to register the patient; in doing so, it is desired to record the patient's demographic data in the practice management information system (PMIS). The physician office is connected to a hospital enterprise's central patient registry. The nurse issues a patient query request to the central patient registry, with some basic patient demographics data as search criteria. In the returned patient list, she picks up an appropriate record for the patient, including the hospital's patient ID, to enter into the PMIS. (Note that the PMIS uses a different Patient ID domain than that of the central patient registry.)

The PMIS uses its own patient identifier, coordinating this identifier with the patient identifier returned in the pick list (sharing the hospital's Patient ID Domain) to retrieve information from the hospital's clinical repository.

Use Case 3: Patient Demographics Query in an Enterprise with Multiple Patient ID Domains

A lab technician enters some basic demographics data (*e.g.*, patient name) into a lab application to query a patient demographics supplier to identify a patient for his lab exams. As the application also needs the patient identifier in another Patient ID Domain in the enterprise for results delivery, the application is configured to receive patient IDs from other domains in the query response.

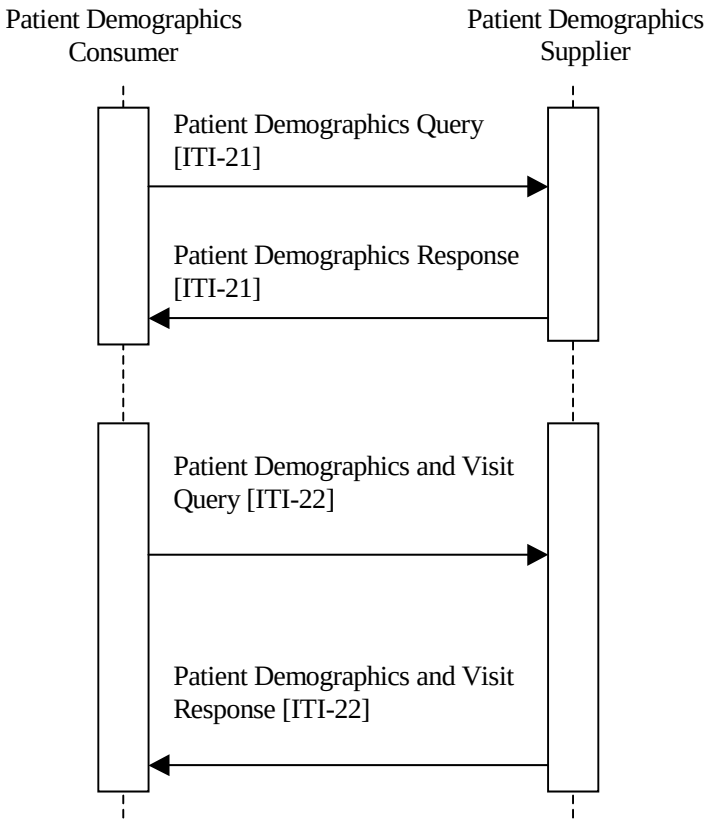


Figure 8.2-1. Basic Process Flow in Patient Demographics Query Profile

8.3.1 Combined Use of PDQ with Other IHE Workflow Profiles

When the Patient Demographics Supplier Actor is grouped with actors in other IHE profiles that perform patient information reconciliation activities (*e.g.*, Radiology PIR), the PDQ Supplier Actor may use the updated information to respond to PDQ Queries. In addition, the Patient Demographics Query Profile may play an integral workflow role in conjunction with other IHE Profiles.

8.3.2 Supplier Data Configuration

A Patient Demographics Supplier Actor that holds demographic information for a single Patient ID domain shall provide matches in that domain.

In the case where the Patient Demographics Supplier Actor holds demographic information for multiple Patient ID domains, the Patient Demographics Supplier Actor shall return information for the domain associated with *MSH-5-Receiving Application* and *MSH-6-Receiving Facility*. See the "Using Patient Data Query (PDQ) in a Multi-Domain Environment" section (ITI TF-2: Appendix M) for a further discussion of this case and an illustration of the supporting architecture.

9 Audit Trail and Node Authentication (ATNA)

The Audit Trail and Node Authentication (ATNA) Integration Profile establishes security measures which, together with the Security Policy and Procedures, provide patient information confidentiality, data integrity and user accountability. This environment is considered the Security Domain and can scale from a department, to enterprise or XDS Affinity Domain. The ATNA model considers that within the secure domain the following is true:

1. All machines are host authenticated. (There are various means of accomplishing this.) This authentication identifies the machine as being one that is known to the security system of the hospital, with known security characteristics. Unknown machines might be granted access, but with the caveat that they are only granted access to information that is authorized for disclosure to the public or to unknown machines. (A patient might choose to allow information such as appointment schedules to be at risk of machine disclosure by unknown machines while not allowing more sensitive PHI to be disclosed.)
2. The host identification is used to determine what (if any) access should be granted to automated processes on that host, and/or persons under the direction of that host's access controls. In practice the automated processes play a critical role, managing issues like pre-fetching, thus person authentication/identification is not sufficient.
3. The secure node is responsible for providing reasonable access controls. This typically includes user authentication and authorization. The value of this user authentication needs to be balanced against the possible safety and patient health impacts of delaying delivery of care by the additional authentication steps.
4. The secure node is also responsible for providing security audit logging to track security events. In healthcare this audit log is often more useful than strict access controls and should be relied upon even in emergencies.

This model is partially driven by the underlying assumption that there will be situations where documents are being exchanged between machines and stored on the recipient. This is partly driven by the need for healthcare systems to operate in disasters and overload situations, where the network operation is limited or destroyed. It is not safe to assume that clients are display only. So there will be semi-permanent copies of most information kept. Even in normal operation, healthcare providers may have only 15 minutes per patient. Good healthcare system design recognizes the need to not waste any of those seconds searching and transferring documents over a network. The documents are transferred in advance, and are kept locally until it is determined that they are no longer needed. There are thin client display only applications in healthcare, but they are limited to uses that can fail without introducing risks to safety or patient health, but a complete security/privacy design requires handling situations where data is stored after retrieval.

ATNA Governance Assumptions

The underlying assumptions are:

- All systems that are members of the secure domain implement a Secure Node Actor for the ATNA profile. The ATNA profile defines transactions between the secure nodes to create a secure domain that is under the management of a domain security officer.

- All applications on a secure node will comply with ATNA requirements, regardless of whether they are IHE Actors or not. They apply to all IT assisted activities that directly create, access, update, and delete PHI, not only those specified by IHE and performed by IHE actors.
- IHE addresses only those security requirements related to systems within the scope of IHE healthcare applications. It does not address other security requirements such as defending against network attacks, virus infection, etc. The principal objective of the Audit Trail mechanism is to track data access to PHI, not IHE transactions.
- Mobile equipment can participate in the Audit Trail and Node Authentication Integration Profile, but special issues related to mobile equipment are not explicitly addressed in this profile.
- ATNA assumes that physical access control, personnel policies and other organizational security considerations necessary to make an enterprise compliant with security and privacy regulations are in place.

9.1 Authentication

ATNA contributes to access control by limiting network access between nodes and limiting access to each node to authorized users. Network communications between secure nodes in a secure domain are restricted to only other secure nodes in that domain. Secure nodes limit access to authorized users as specified by the local authentication and access control policy.

9.1.1 User Authentication

The Audit Trail and Node Authentication Integration Profile requires only local user authentication. The profile allows each secure node to use the access control technology of its choice to authenticate users. The use of Enterprise User Authentication is one such choice, but it is not necessary to use this profile.

9.1.2 Connection Authentication

The Audit Trail and Node Authentication Integration Profile requires the use of bi-directional certificate-based node authentication for connections to and from each node. The DICOM, HL7, and HTML protocols all have certificate-based authentication mechanisms defined. These authenticate the nodes, rather than the user. Connections to these machines that are not bi-directionally node-authenticated shall either be prohibited, or be designed and verified to prevent access to PHI.

Note: Communications protocols that are not specified by IHE profiles, e.g. SQL Server, must be bi-directionally authenticated if they will be used for PHI. This profile does not specify how that authentication is to be performed.

This requirement can also be met by ensuring complete physical network security with strict configuration management. This means that no untrusted machine can obtain physical access to any portion of the network. Making the connection authentication configurable enhances performance in physically secured networks. A Secure Node Actor shall be configurable to support both connection authentication and physically secured networks.

IHE does not mandate the use of encryption during transmission. Most hospital networks provide adequate security through physical and procedural mechanisms. The additional performance penalty for encryption is generally not justified for these networks. This profile mandates the use of the

TLS security negotiation mechanism for all communications between secure nodes as a means of ensuring that they only communicate with other authorized secure nodes. It permits the negotiation of encryption if both nodes are configured to request and support encryption. This allows installation of IHE secure nodes into environments where the network is not otherwise secured.

9.2 Audit Trails

User Accountability is provided through Audit Trail. The Audit Trail needs to allow a security officer in an institution to audit activities, to assess compliance with a secure domain's policies, to detect instances of non-compliant behavior, and to facilitate detection of improper creation, access, modification and deletion of Protected Health Information (PHI). PHI is considered to be the patient-identifiable information records (e.g. Registration, Order, Study/Procedure, Reports, Images, and Presentation States). PHI may be accessed by users or exchanged between the systems. This includes information exported to and imported from every secured node in the *secure domain*.

The user accountability is further enhanced through a standards based Centralized Audit Record Repository, that provides a central Audit Record repository as the simplest means to implement security requirements. A transfer of Audit Records from all the IHE actors to the Audit Record Repository reduces the opportunities for tampering and makes it easier to audit the department. Disconnected nodes may store audit data for transfer to the Audit Repository upon reconnection to the secure domain network.

The audit trail contains information so that questions can be answered such as:

- For some user: which patients' PHI was accessed?
- For some patient PHI: which users accessed it?
- What user authentication failures were reported?
- What node authentication failures were reported?

The Audit Trail and Node Authentication Profile provides tools that are useful for enterprises attempting to become compliant with privacy and security regulations (HIPAA, European, Japanese, etc.), but the profile does not itself make the enterprise compliant. For guidance on proper audit log management enterprises should look to documents such as NIST SP 800-92 – Guide to Computer Security Log Management.

9.2.1 Audit Messages

The use of auditing as part of a security and privacy process is appropriate for situations where the people involved are generally trustworthy and need a wide range of flexibility to respond rapidly to changing situations. This is the typical healthcare provider environment. Auditing tracks what takes place, and the people involved know that their actions are being audited. This means that the audit records must capture event descriptions for the entire process, not just for individual components that correspond to individual IHE actors.

The IHE audit trail is the first of several profiles that correspond to different forms of access control and authentication. Auditing is always needed independent of the access control and authentication method chosen.

The IHE-specified audit flow is illustrated in Figure 9.2-1.

1. Real world activities take place, and some of these activities involve the applications processing of a device that includes support for some IHE profiles. This product has components that may correspond to specific IHE Actors. The product may also have other capabilities that are independent of IHE recommendations.

2. A wide variety of events take place during this process. Some of these events are directly related to IHE Actor activities. Others may be indirectly related, and still others are not related to any IHE specification. The events are both extremely detailed minor events, such as keystrokes, and high level events such as analyzing a diagnostic study. Very few of these events are relevant to security and privacy auditing. Most are too low level to be useful or are otherwise irrelevant.

3. The "Security Audit and Access Accountability Message XML Data Definitions for Healthcare Applications" (RFC-3881) defines an XML schema for reporting events that are relevant to security and privacy auditing. It was defined in cooperation with the ASTM, HL7, and DICOM standards organizations and the NEMA/COCIR/JIRA Security and Privacy Committee. The IHE recommends the use of the RFC-3881 format, and recommends reporting only events that it can describe.

4. DICOM has standardized some of the audit message vocabulary. The DICOM Audit Message Vocabulary extends the basic vocabulary provided with RFC-3881, and also further specifies some optional elements in RFC-3881. An example of vocabulary extension is the addition of a coded value to indicate that a field contains a DICOM Study Instance UID. An example of optional element specification is the requirement that the UserID field in RFC-3881 messages shall be the user ID used by the local device operating system, and that the AlternateID shall be the user ID used by the enterprise authentication system (if it is different).

5. This profile defines other events that do not correspond to events defined in the DICOM vocabulary. These events are describable by RFC-3881, and this profile includes requirements for such descriptions.

6. IHE auditing specifies that when using the RFC-3881, events that can be described using the DICOM vocabulary they shall be reported using the DICOM vocabulary, even if the device is not otherwise a DICOM compliant device. Events that do not match the DICOM vocabulary shall be reported using RFC-3881 vocabulary or other extensions. Events that cannot be reported using RFC-3881 are not candidates for reporting.

7. The local site will then apply its own reporting policies. The IHE profile specifies the capabilities that should be present for audit reporting, and also that there should be controls present to allow the local site security administration to control reporting detail. The IHE profile does not specify any audit reporting functions or formats.

8. IHE specifies events that must be reported in the audit trail. There are other events related to security, which may be reported in the audit trail or by other means. This profile does not describe them and does not require that they use this reporting format or mechanism. Examples of such events are OS login, network routing and firewall logs.

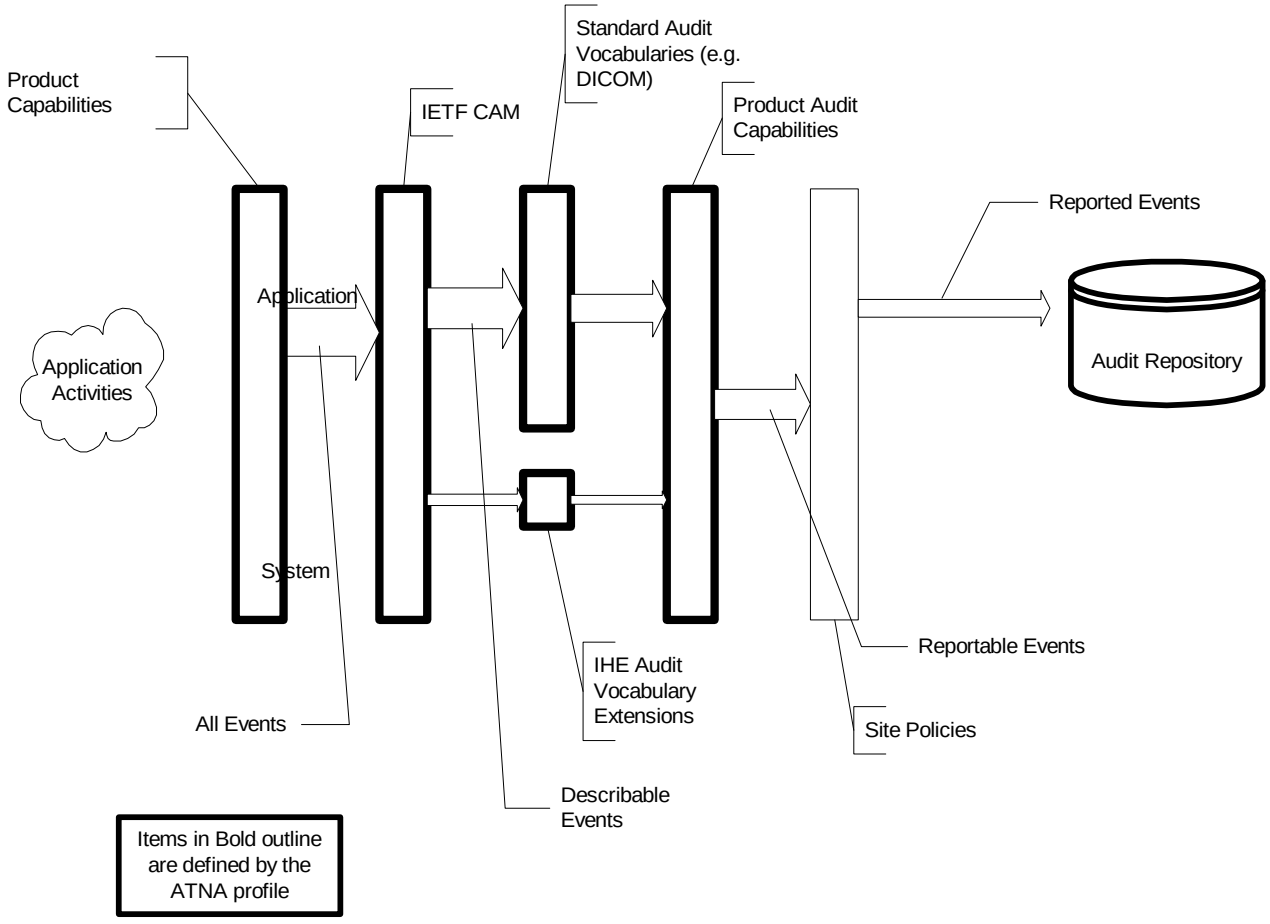


Figure 9.2-1 Flow of Events into Audit Messages

9.2.2 Backwards Compatibility

This profile also defines the continued use of messages that are formatted in accordance with the IHE Provisional Audit Message format from the deprecated Basic Security Profile in IHE Radiology TF 6.0. This older format describes events that are suitable for reporting in Radiology and other diagnostic and treatment activities. These events are a subset of the kind of events that can be described using RFC-3881 and the DICOM vocabulary.

The IHE ATNA Profile also allows for the reporting of these events using the Provisional format over either of the IHE specified transport mechanisms. The intention is that products will gradually transition from the Provisional message format to RFC-3881 format, but it is recognized that this transition will take time and that there is a significant installed base.

The Provisional format is unlikely to be of interest to other healthcare applications, which should use the RFC-3881 format and DICOM Vocabulary where appropriate.

9.3 Audit Trail Transport

The Audit Trail and Node Authentication Integration Profile specifies the use of Reliable Syslog Cooked Profile (RFC-3195, Section 4) as the mechanism for logging audit record messages to the central audit record repository. It also permits the use of BSD Syslog (RFC-3164). There are, however, several known limitations of BSD Syslog:

- There is no confirmation to the sender that the audit record message was received at the destination
- There is no option to encrypt the audit record messages
- Authentication by means of certificates of the sending nodes and the central audit repository is not possible
- Messages may be truncated or lost.

The specification of Reliable Syslog Cooked Profile messages corrects these deficiencies.

9.4 Actors/Transactions

Table 9.4-1 lists the transactions for each actor directly involved in the Audit Trail and Node Authentication Integration Profile. In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled "R"). Transactions labeled "O" are optional. A complete list of options defined by this Integration Profile that implementations may choose to support is listed in ITI-TF 1: 9.4. Their relationship is shown in Figure 9.4-1.

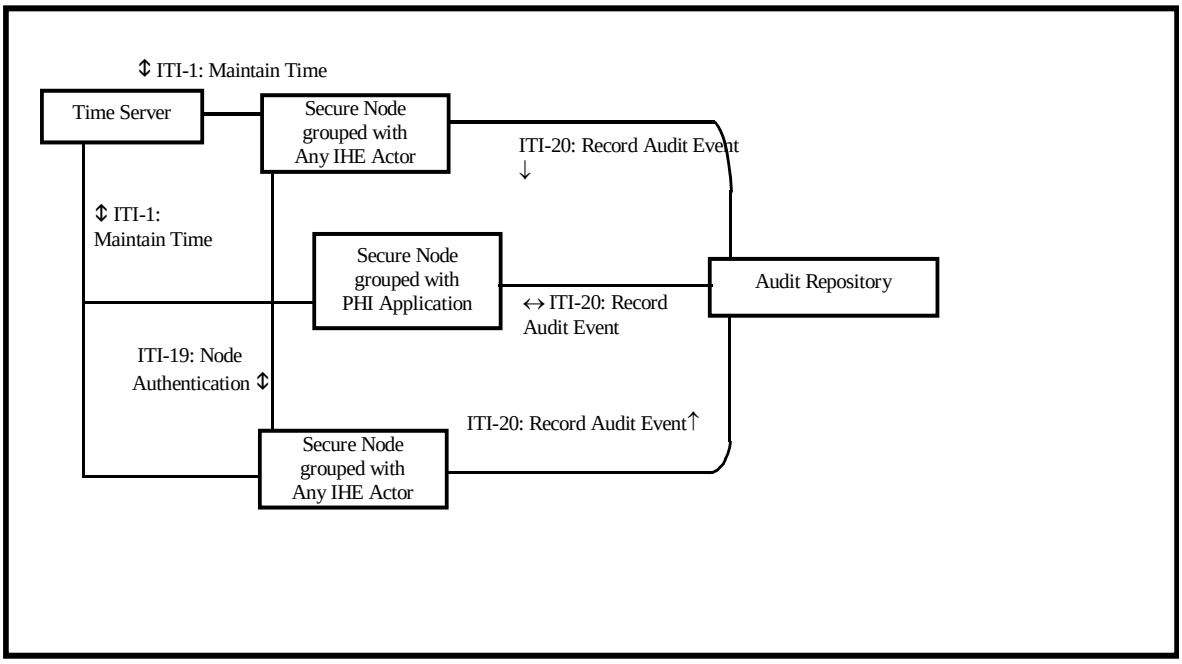


Figure 9.4-1. Audit Trail and Node Authentication Diagram

When an implementation chooses to support this Integration Profile for an actor, that actor shall be grouped with the Secure Node actor. It is required that all IHE actors and any other activities in this implementation support the Audit Trail and Node Authentication Integration Profile.

A means must be provided to upload the required certificates to the implementation, e.g. via floppy disk or file transfer via network.

Non-IHE applications that process PHI shall detect and report auditable events, and protect access.

Table 9.4-1. Audit Trail and Node Authentication Integration Profile - Actors and Transactions

Actor	Transactions	Optionality	Vol II / III Section
<any PHI application grouped with a Secure Node Actor>	Record Audit Event	R	IHE ITI-2: 3.20
<any IHE actor grouped with a Secure Node actor>	Record Audit Event	R	IHE ITI-2: 3.20
Audit Record Repository	Record Audit Event	R	IHE ITI-2: 3.20
Secure Node	Authenticate Node	R	IHE ITI-2: 3.19
	Maintain Time	R	IHE ITI-2: 3.7
Secure Application	Authenticate Node	O	IHE ITI-2: 3.19
	Maintain Time	O	IHE ITI-2: 3.7
	Record Audit Event	O	IHE ITI-2: 3.20

The Secure Node Actor shall include:

1. The Authenticate Node transaction for all network connections that may expose private information. These transactions are defined for:
    1. DICOM, using TLS
    2. HL7, using TLS
    3. HTTP, using TLS
2. All local user activity (login, logout, etc.) protected to ensure only authorized users.
3. An audit transport mechanism, either:
    a) Reliable Syslog Cooked Profile format (RFC-3195, Section 4)
    b) BSD Syslog (RFC-3164), the baseline syslog mechanism.
    c) Generation of audit messages for recommended events utilizing one of the defined alternatives for audit message formats. The audit messages formatted are:
4. The IETF common audit message format, using the DICOM and IHE vocabularies.
    a) The Provisional IHE Audit Message format

The difference between the Secure Node and the Secure Application is the extent to which the underlying operating system and other environment are secured. A Secure Node includes all aspects of user authentication, file system protections, and operating environment security. The Secure Application is a product that does not include the operating environment. The Secure Application provides security features only for the application features. See section 9.7 for the relationships among a Secure Node, Secure Application, and other actors.

1. The Audit Repository shall support:
2. Both audit transport mechanisms.

3. Any IHE-specified audit message format, when sent over one of those transport mechanisms. Note that new applications domains may have their own extended vocabularies in addition to the DICOM and IHE vocabularies. This also means that an ATNA Audit Repository is also automatically a Radiology Basic Security profile Audit Repository because it must support the IHE Provisional Message format and it must support the BSD syslog protocol.
4. Self protections and user access controls.

This profile does not specify other functions for the Audit Repository, but it is expected that most repositories will perform screening, reporting, archival, etc.

9.5 ATNA Integration Profile Options

Options that may be selected for this Integration Profile are listed in the table 8.2-1 along with the actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 9.5-1 ATNA - Actors and Options

Actor	Options	Vol & Section
Audit Record Repository	*None*	-
Secure Node	*ATNA Encryption* *Radiology Audit Trail*	ITI TF-2: 3.22 RAD TF-1: 2.2.1; TF-3: 5.1
Secure Application	*ATNA Encryption*	ITI TF-2: 3.22

9.5.1 ATNA Encryption Option

Secure Nodes may implement the ATNA Encryption Option. This option specifies the support of encryption to protect confidentiality.

9.5.2 Radiology Audit Trail Option

The Radiology Audit Trail provides specific requirements as to which audit events IHE Radiology actors are required to send. It also details the specific format of certain audit events based on the Radiology actor.

9.6 Audit Trail and Node Authentication Process Flow

The security measures in the Audit Trail and Node Authentication Integration Profile are user authentication, node authentication, and generation of audit records. Node authentication and user authentication define a number of transactions that establish the concept of a Secure Node and a collection of connected Secure Nodes in a secure domain (see Volume ITI-III: Appendix A). Generation of audit records requires a set of audit trigger events and a definition of the content of the audit records. This profile specifies two acceptable message formats:

1. Messages formatted in accordance with the IHE Audit Message format. This is a combination of the DICOM Audit Messages format and IHE extensions. The IHE extensions to RFC-3881 add event codes and information needed for uses that are not within the domain of the DICOM Standard.

2. The predecessor IHE Provisional Audit Message format. This format was defined as an interim format while the standards work to define the Common Audit Message format and vocabularies progressed through the standards organizations.

   Based on the work done in ASTM (E2147-01 Standard Specification for Audit and Disclosure Logs for Use in Health Information Systems) and HL7 (Framework for Audit Messages), IHE defined a detailed set of audit trigger events, a set of general audit messages with the content for the audit record, and a mapping for each event to a general audit message. The content of the audit record has been specified by means of an XML Schema (see Volume ITI-II: Appendix F).

In the following paragraphs three typical process flows are described for situations in which authorized users, unauthorized users, and unauthorized nodes attempt to gain access to protected health information (PHI).

9.6.1 Normal Node Process Flow

The following scenario shows how the IHE security measures operate for authorized access to PHI from an authorized node in the network:

1. Time synchronization occurs independently. These transactions may take place at any time. Correct time is needed to generate Audit Records with a correct timestamp.
2. A user logs on to Image Display/Secure Node actor.
   The user enters valid credentials and is authorized to access the node.
3. The node generates audit records.
4. The user wants to query/retrieve and view some images.
   Before image transactions can take place, an authentication process between the Image Display/Secure Node actor and the Image Manager/Image Archive/Secure Node actor takes place.
5. Following node authentication, the node initiates the query/retrieve transactions.
6. The node generates audit records.

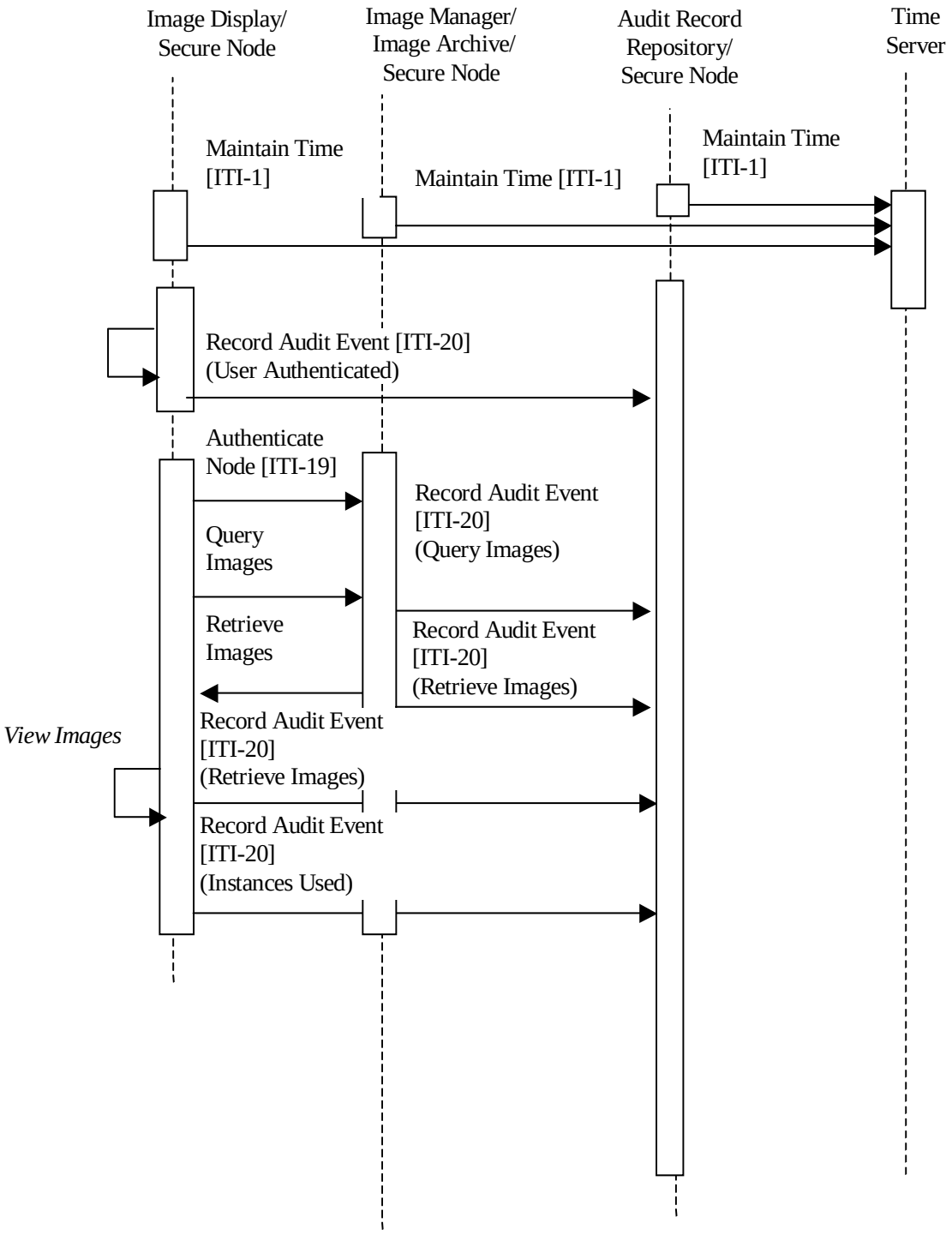


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Figure 9.6-1. Authorized Node Process Flow

9.6.2 Unauthorized Node Process Flow

The following scenario shows how the IHE security measures help to prevent unauthorized access to PHI from an unauthorized node in the network:

1. An unauthorized node tries to query the Lab Automation Manager/Secure Node actor for information. This fails because no authentication has taken place, and an audit record is generated.
2. The unauthorized node tries an authentication process with the Lab Automation Manager/Secure Node. This fails because the Lab Automation Manager/Secure Node will not trust the certificate presented by the Malicious Node, and an audit record is generated.

Note that the sequencing of the transactions is just one example; transactions from an unauthorized node are totally unpredictable and may happen in any order.

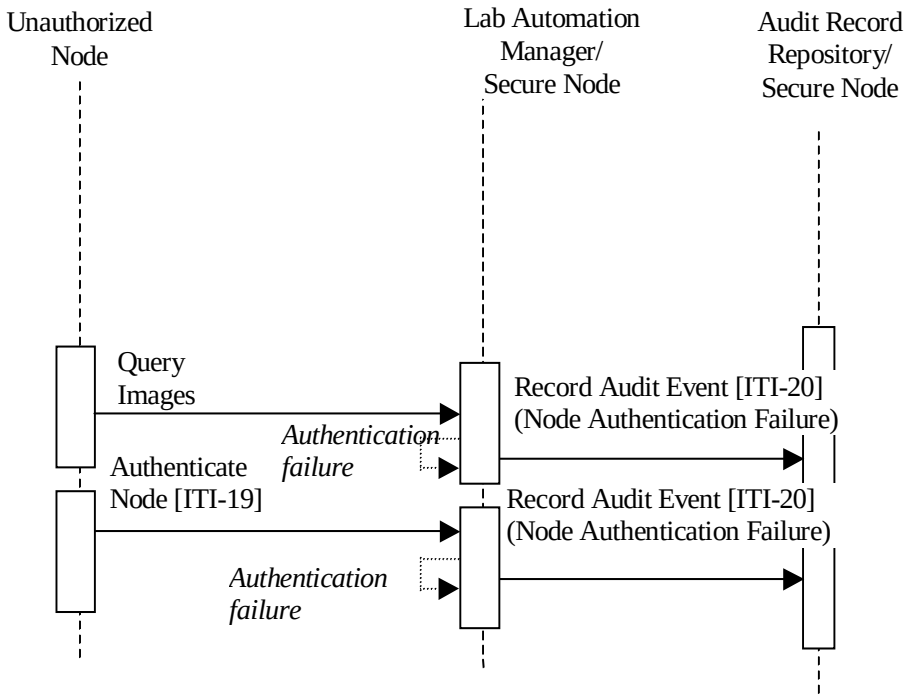


Figure 9.6-2. Unauthorized Node Process Flow

9.6.3 Unauthorized User Process Flow

The following scenario shows how the IHE security measures help to prevent unauthorized access to PHI from an unauthorized user in the healthcare enterprise:

1. An unauthorized user tries an authentication process with the ECG Display/Secure Node actor. This fails because the ECG Display/Secure Node actor detects that the user name and credentials presented are not valid at this secure node, and an audit record is generated.

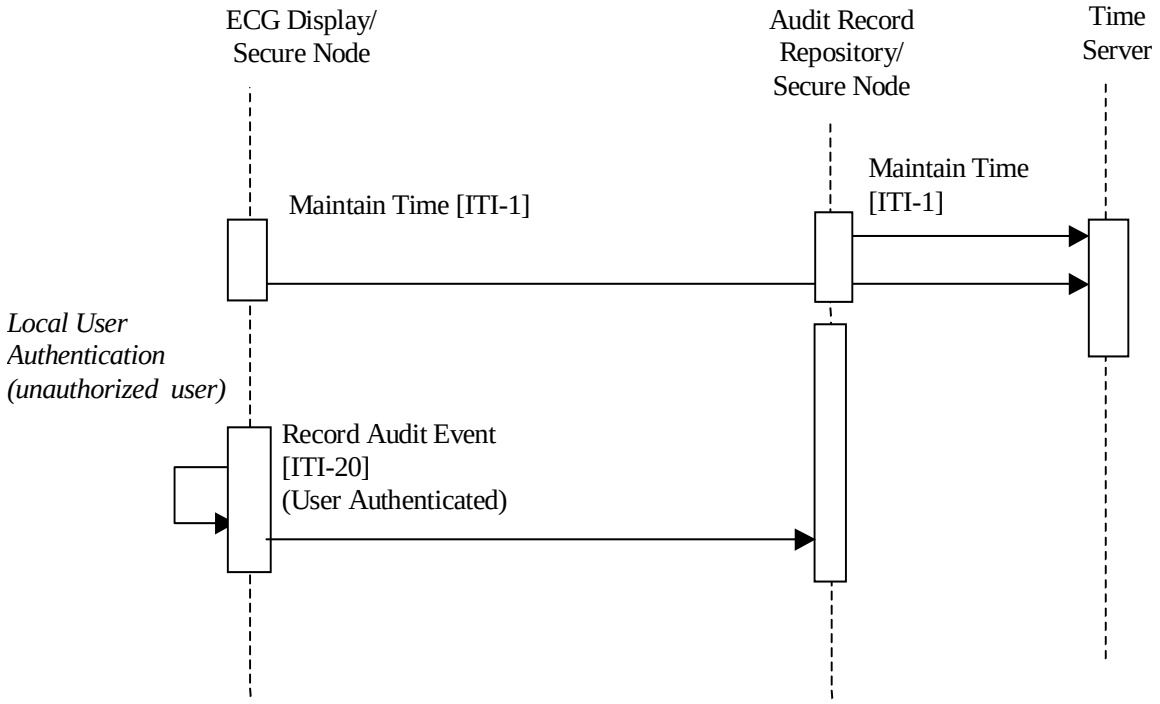


Figure 9.6-3. Unauthorized User Process Flow

9.7 Relationship between Secure Node, Secure Application, and other Actors

The allocation of responsibilities when an actor is grouped with a secure node can be complex when different parties are responsible for different parts of the system. This situation arises frequently in situations like web server applications, where there is an operating system, a web server framework, and individual web applications. These might all be from different vendors. Each of these components has a role in performing security related tasks. There is also a system integrator who is responsible for assembling these components into the final complete system. It is the responsibility of the system integrator to insure that all of the necessary security functions are implemented by the appropriate system component.

Note: The system integrator might be a product vendor, outside consultant or internal staff. IHE does not specify business relationships. The term is used here to indicate a functional role, not a business relationship.

IHE has split these into two primary categories:

- The healthcare functions. These are identified as IHE actors. IHE does not specify how functional actors are implemented. Multiple actors might be implemented by one web application, and it may take multiple web applications to implement one IHE actor. IHE allocates functions to the actors and it is the implementer's task to allocate these to individual web applications.
- The underlying operating environmental components. The IHE identifies these as the Secure Node actor. It is the system integrator that determines how the functions of the Secure Node actor are allocated to individual components.

When a product claims support for the Secure Application actor, it is claiming that it performs those functions that are appropriate to its IHE task. This will certainly include some audit responsibilities, will probably include some communications security responsibilities, and may include other security responsibilities. The specifics of these responsibilities depend upon the functions and options of that product. For example, a product that includes a user login capability will generate user related audit events and perform the user authentication. In contrast, a single function web application might only generate audit messages related to its function, and will depend upon the external secure node environment for other functions.

This means that product descriptions must be sufficient for the system integrator to determine whether all of the necessary security functions are present. If the single purpose web application is depending on the web server environment to provide node authentication, this must be clear to the system integrator. Not all web server environments provide that authentication, and the integrator will need to ensure that authentication is provided when needed.

When describing what security features have been implemented in a product, the following rules apply:

1. If the product claims to include the Secure Node actor, the product has been integrated so that all of the operating system and other environmental security features are present.
2. If the product claims only to include the Secure Application actor, that indicates that only those security features that apply to the application features are provided by the product.

Product selection can then use the IHE conformance claim for a summary view of the security features provided by the product. The system integrator can use this information to determine what additional products or integration work will be needed to establish the functionality provided by a Secure Node if the application products are only Secure Applications.

10 Cross-Enterprise Document Sharing (XDS)

The ***Cross-Enterprise Document Sharing*** IHE Integration Profile facilitates the registration, distribution and access across health enterprises of patient electronic health records. Cross-Enterprise Document Sharing (XDS) is focused on providing a standards-based specification for managing the sharing of documents between any healthcare enterprise, ranging from a private physician office to a clinic to an acute care in-patient facility.

The Cross-Enterprise Document Sharing Integration Profile specified in this Section is called XDS. The reader should be aware that it is quite often called "XDS.a" in other IHE documents to distinguish it from the new implementation choice for the Cross-Enterprise Document Sharing Integration Profile (XDS.b). New implementers of XDS are encouraged to review the XDS.b Supplement (www.IHE.net/Technical_Frameworks). The XDS Profile specified in this section (or XDS.a) employs different versions of the same standard (ebXML Registry 2.0 and 3.0) and web services standards that are no longer consistent with the current developments and best practices in the industry (e.g. moving from SOAP V1.1 to V1.2, MTOM/XOP replacing SOAP with Attachments or SwA). The XDS.b Integration Profile is now the basis for new IHE Profiles in the area of security (XUA Supplement) and Cross Community communication (XCA Supplement). The XDS.a and XDS.b Integration Profiles are equivalent in terms of functionality to facilitate migration from XDS.a to XDS.b, if desired, as well as coexistence of implementations supporting the two Integration Profiles in the same environment.

The XDS IHE Integration Profile assumes that these enterprises belong to one or more XDS Affinity Domains. An XDS Affinity Domain is a group of healthcare enterprises that have agreed to work together using a common set of policies and share a common infrastructure.

Examples of XDS Affinity Domains include:

- Community of Care supported by a regional health information organization in order to serve all patients in a given region.
- Nationwide EHR
- Specialized or Disease-oriented Care
  - Cardiology Specialists and an Acute Cardiology Center
  - Oncology network
  - Diabetes network
- Federation of enterprises
  - A regional federation made up of several local hospitals and healthcare providers
- Government sponsored facilities (e.g., VA or Military)
- Insurance Provider Supported Communities

Within an XDS Affinity Domain, certain common policies and business rules must be defined. They include how patients are identified, consent is obtained, and access is controlled, as well as the format, content, structure, organization and representation of clinical information. This Integration Profile does not define specific policies and business rules, however it has been designed to accommodate a wide range of such policies to facilitate the deployment of standards-based infrastructures for sharing patient clinical documents. This is managed through federated document

repositories and a document registry to create a longitudinal record of information about a patient within a given XDS Affinity Domain. These are distinct entities with separate responsibilities:

- A document repository is responsible for storing documents in a transparent, secure, reliable and persistent manner and responding to document retrieval requests.
- A document registry is responsible for storing information about those documents so that the documents of interest for the care of a patient may be easily found, selected and retrieved irrespective of the repository where they are actually stored.

The concept of a document in XDS is not limited to textual information. As XDS is document content neutral, any type of clinical information without regard to content and representation is supported. This makes the XDS IHE Integration Profile equally able to handle documents containing simple text, formatted text (*e.g.*, HL7 CDA Release 1), images (*e.g.*, DICOM) or structured and vocabulary coded clinical information (*e.g.*, CDA Release 2, CCR, CEN ENV 13606, DICOM SR). In order to ensure the necessary interoperability between the document sources and the document consumers, the XDS Affinity Domain must adopt policies concerning document format, structure and content.

The XDS Integration Profile is not intended to address all cross-enterprise EHR communication needs. Some scenarios may require the use of other IHE Integration profiles, such as Patient Identifier Cross-Referencing, Audit Trail and Node Authentication, Cross-Enterprise User Authentication, and Retrieve Information for Display. Other scenarios may be only partially supported, while still others may require future IHE Integration profiles, which will be defined by IHE as soon as the necessary base standards are available. Specifically:

1. The management of dynamic information such as allergy lists, medication lists, problem lists, etc is not addressed by XDS. However, the Retrieve Information for Display Integration Profile does provide some transactions (e.g., LIST-ALLERGIES, LIST-MEDS) that may be used to provide an elementary support of such capabilities. A complementary approach to managing updates and structured application access to such dynamic clinical information may be expected as a separate Integration Profile in the future.
2. The placing and tracking of orders (e.g. drug prescriptions, radiology orders, etc.) is not supported by XDS. This does not preclude the use of XDS to store and register orders and corresponding results when such artifacts need to be recorded in the patient's health record. However, XDS provides no facilities for tracking progress of an order through its workflow, and therefore is not intended for order management. A complementary approach to cross-enterprise order workflow (ePrescription, eReferral) may be expected as separate Integration Profiles in the future.
3. The operation of any XDS Affinity Domain will require that a proper security model be put in place. It is expected that a range of security models should be possible. Although the XDS Integration Profile is not intended to include nor require any specific security model, it is expected that XDS implementers will group XDS Actors with actors from the IHE Audit Trail and Node Authentication and will need an Access Control capability that operates in such a cross-enterprise environment. Specific IHE Integration Profiles complementary to XDS are available (e.g. Cross-Enterprise User Authentication, Document Digital Signature, etc).

4. The establishment of independent but consistently XDS Affinity Domains will call for their federation, as patients expect their records to follow them as they move from region to region, or country to country. IHE foresees a need for transferring information from one XDS Affinity Domain to another, or to allow access from one XDS Affinity Domain to documents managed in other XDS Affinity Domains. XDS has been designed with this extension in mind. An XDS Domains Federation Integration Profile that complements XDS may be anticipated in the future.

5. XDS does not address transactions for the management or configuration of an XDS Affinity Domain. For example, the configuration of network addresses or the definition of what type of clinical information is to be shared is specifically left up to the policies established by the XDS Affinity Domain.

10.1 Actors/Transactions

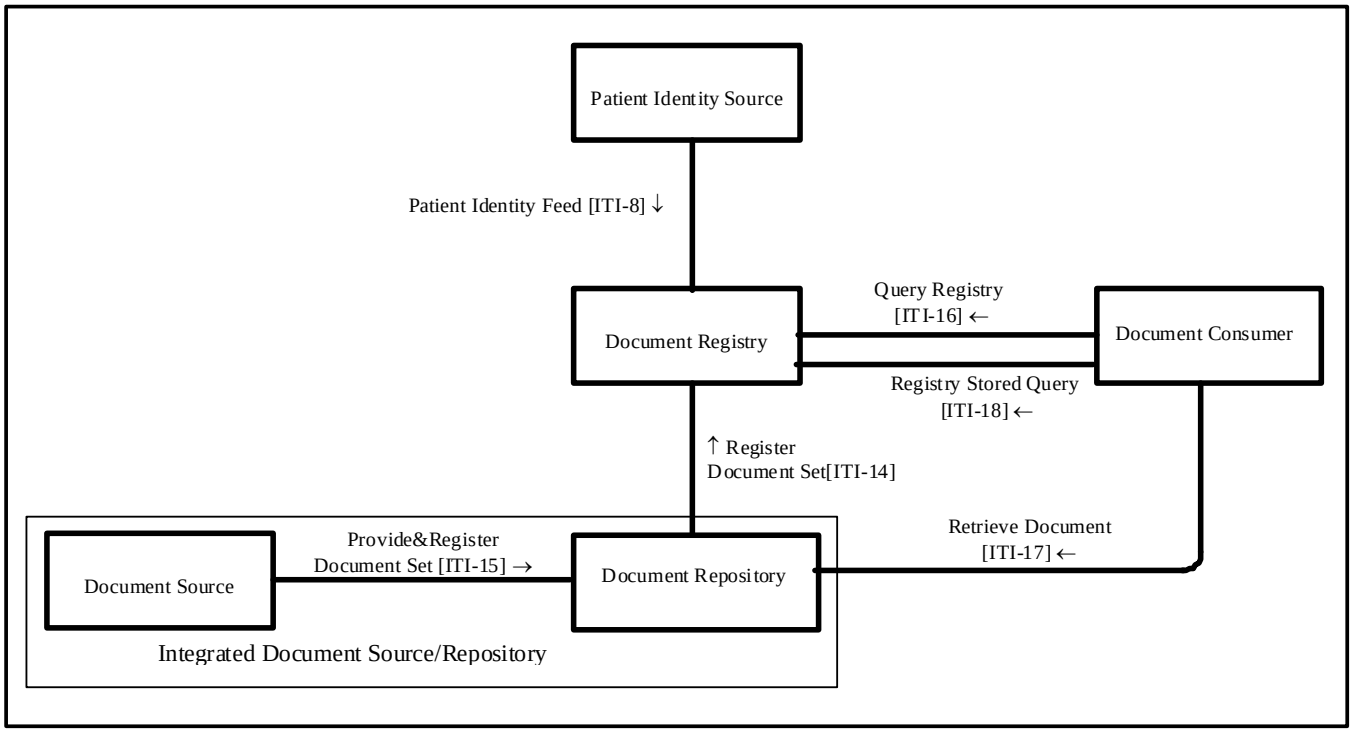


Figure 10.1-1 Cross-Enterprise Document Sharing Diagram

Table 10.1-1 XDS - Actors and Transactions

Actors	Transactions	Optionality	Section in Vol. 2
Document Consumer	Query Registry	O	ITI TF-2:3.16
	Retrieve Document	R	ITI TF-2:3.17
	Registry Stored Query	R (Note 2)	
Document Source	Provide and Register Document Set	R	ITI TF-2:3.15
Document Repository	Provide and Register Document Set	R	ITI TF-2:3.15
	Register Document Set	R	ITI TF-2:3.14
	Retrieve Document	R	ITI TF-2:3.17
Document Registry	Register Document Set	R	ITI TF-2:3.14
	Query Registry	O	ITI TF-2:3.16
	Patient Identity Feed	R	ITI TF-2:3.8
	Registry Stored Query	R (Note 2)	
Integrated Document Source/Repository	Register Document Set	R	ITI TF-2:3.14
	Retrieve Document	R	ITI TF-23.17
Patient Identity Source	Patient Identity Feed	R (Note 1)	ITI TF-2:3.8

Note 1: The Patient Identity Source is required to use an OID to identify the Assigning Authority in Transaction ITI-8. For technical details of the assigning authority information, see Transaction 8 in Technical Framework, Volume 2.

Note 2: The Document Registry actor part of the Registry Stored Query transaction shall implement all queries defined by the Registry Stored Query transaction. No such minimum requirements are placed on the Document Consumer actor.

10.1.1 Actors

10.1.1.1 Document Source

The Document Source Actor is the producer and publisher of documents. It is responsible for sending documents to a Document Repository Actor. It also supplies metadata to the Document Repository Actor for subsequent registration of the documents with the Document Registry Actor.

10.1.1.2 Document Consumer

The Document Consumer Actor queries a Document Registry Actor for documents meeting certain criteria, and retrieves selected documents from one or more Document Repository actors.

10.1.1.3 Document Registry

The Document Registry Actor maintains metadata about each registered document in a document entry. This includes a link to the Document in the Repository where it is stored. The Document Registry responds to queries from Document Consumer actors about documents meeting specific criteria. It also enforces some healthcare specific technical policies at the time of document registration.

10.1.1.4 Document Repository

The Document Repository is responsible for both the persistent storage of these documents as well as for their registration with the appropriate Document Registry. It assigns a URI to documents for subsequent retrieval by a Document Consumer.

10.1.1.5 Patient Identity Source

The Patient Identity Source Actor is a provider of unique identifier for each patient and maintains a collection of identity traits. The Patient Identify Source facilitates the validation of patient identifiers by the Registry Actor in its interactions with other actors.

10.1.1.6 Integrated Document Source/Repository

The Integrated Document Source/Repository combines the functionality of the Document Source and Document Repository actors into a single actor that does not initiate nor accept the Provide ad Register Document Set transaction. This actor may replace the Document Repository actor from the perspective of the Register Document Set or Retrieve Document transactions.

10.1.2 Transactions

10.1.2.1 Provide and Register Document Set

A Document Source Actor initiates the Provide and Register Document Set Transaction. For each document in the submitted set, the Document Source Actor provides both the documents as an opaque octet stream and the corresponding metadata to the Document Repository. The Document Repository is responsible to persistently store these documents, and to register them in the Document Registry using the Register Documents transaction by forwarding the document metadata received from the Document Source Actor.

10.1.2.2 Register Document Set

A Document Repository Actor initiates the Register Document Set transaction. This transaction allows a Document Repository Actor to register one or more documents with a Document Registry, by supplying metadata about each document to be registered. This document metadata will be used to create an XDS Document Entry in the registry. The Document Registry Actor ensures that document metadata is valid before allowing documents to be registered. If one or more documents fail the metadata validation, the Register Document Set transaction fails as a whole.

To support composite documents, an XDS Document may be a multipart document. The Document Repository must handle multi-part data sets as an “opaque entity”. The Document Repository does not need to analyze or process its multi-part structure nor the content of any parts in the context of the XDS Integration Profile.

10.1.2.3 Query Registry

The Query Registry transaction is issued by the Document Consumer Actor on behalf of a care provider (EHR-CR) to a Document Registry. The Document Registry Actor searches the registry to locate documents that meet the provider’s specified query criteria. It will return a list of document

entries that contain metadata found to meet the specified criteria including the locations and identifier of each corresponding document in one or more Document Repositories.

10.1.2.4 Registry Stored Query

The Registry Stored Query transaction is issued by the Document Consumer Actor on behalf of a care provider (EHR-CR) to a Document Registry. The Document Registry Actor searches the registry to locate documents that meet the provider's specified query criteria. It will return registry metadata containing a list of document entries found to meet the specified criteria including the locations and identifier of each corresponding document in one or more Document Repositories.

This transaction differs from Query Registry [ITI-16] by storing the query in the Document Registry actor and referencing the query in the transaction instead of passing SQL.

With the Query Registry Transaction [ITI-16], SQL language queries are transmitted to the Registry actor and results are returned. In a Stored Query, the definition of the query is stored on the Registry actor. To invoke the query, an identifier associated with the query is transmitted along with parameters defined by the query. This has the following benefits:

1. Malicious SQL transactions cannot be introduced
2. Alternate database styles and schemas can be used to implement the Document Registry actor. This is because the style of SQL query statements is directly related to the table layout in a relational database.

This profile does not define how Stored Queries are loaded into or implemented in the Document Registry actor.

10.1.2.5 Retrieve Document

A Document Consumer Actor initiates the Retrieve Document transaction. The Document Repository will return the document that was specified by the Document Consumer.

To support composite documents, an XDS Document may be a multipart document. In this case, the Document Consumer must take appropriate actions to make the multipart content accessible to the user.

10.1.2.6 Patient Identity Feed

The Patient Identity Feed Transaction conveys the patient identifier. It conveys the patient identifier and corroborating demographic data, captured when a patient's identity is established, modified or merged or in cases where the key corroborating demographic data has been modified. Its purpose in the XDS Integration Profile is to populate the registry with patient identifiers that have been registered for the XDS Affinity Domain.

The Patient Identify Feed Transaction defined in ITI TF-2: 3.8 uses standard HL7 encoding of Patient Identifiers in PID-3. As defined in ITI TF-2: 3.8, the value in PID-3 may have different components and subcomponents as long as the required values are present. This is standard encoding for HL7 applications; receiving applications are expected to extract the required data for their use.

When combined with the other XDS transactions, Document Registry actors and other actors that receive HL7 data with Patient Identifiers are required to map the data received in the HL7 message to the format specified in those other XDS transactions (ITI-14, ITI-15, ITI-16). In those transactions, the Patient ID is treated using ebXML encoding rules and not HL7 encoding rules. Specifically, the Patient ID will be treated as a string, and extra components entered in that string will cause those transactions to fail. XDS actors are required to use the specified encoding for Patient ID values in other transactions and not merely copy the value received in an HL7 transaction.

10.1.3 XDS Document Contents Support

The following table lists the document contents supported in other IHE Integration Profiles, which specify concrete content types for sharing of clinical documents in various domains. These profiles are built on the XDS profile, and may define additional constraints and semantics for cross-enterprise document sharing in their specific use cases.

Table 10.1-1: List of IHE Integration Profiles and Document Types They Support

IHE Technical Framework Domain	Integration Profile Name	Document Content Supported
Patient Care Coordination	Cross-Enterprise Sharing of Medical Summaries	Medical Summary in the HL7 CDA format
Radiology	Cross-Enterprise Document Sharing for Imaging (XDS-I)	Radiology Diagnostic Report in the plain text or PDF formats
		Reference to a collection of DICOM SOP Instances in a manifest document in the DICOM Key Object Selection format

10.2 Integration Profile Options

Options that may be selected for this Integration Profile are listed in the table 10.2-1 along with the Actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 10.2-1 XDS - Actors and Options

Actor	Options	Vol & Section
Document Source	*Off-Line transaction mode*	ITI TF-1:10.4.12 ITI TF-1:J.6
	Document Replacement	ITI TF-1:10.2.1
	Document Addendum	ITI TF-1:10.2.2
	Document Transformation	ITI TF-1:10.2.3
	Folder Management	ITI TF-1:10.2.4
	Basic Patient Privacy Enforcement	ITI TF-2:3.15.4.1.3.1

Actor	Options	Vol & Section
Document Repository	*Off-Line transaction mode*	ITI TF-1:10.4.12 ITI TF-1:J.6
Document Registry (Note 1)	*No options defined*	- -
Integrated Document Source / Repository	*Document Replacement*	ITI TF-1:10.2.1
	Document Addendum	ITI TF-1:10.2.2
	Document Transformation	ITI TF-1:10.2.3
	Folder Management	ITI TF-1:10.2.4
Document Consumer	*Basic Patient Privacy Enforcement*	ITI TF-2:3.18.4.1.3.5
	Basic Patient Privacy Proof	ITI TF-2:3.18.4.1.3.6
Patient Identity Source	*No options defined*	- -

Note 1: A XDS Document Registry has always been required to validate that documents that are registered do contain a confidentialityCode from an XDS Affinity Domain vocabulary. The BPPC profile is giving some structure to this XDS Affinity Domain defined vocabulary.

10.2.1 Document Replacement Option.

In this option the Document Source or Integrated Document Source/Repository shall offer the ability to submit a document as a replacement for another document already in the registry/repository. Grouping with Document Consumer can be used to obtain the most recent metadata and ids to be used in the replace submission.

10.2.2 Document Addendum Option

In this option the Document Source or Integrated Document Source/Repository shall offer the ability to submit a document as an addendum to another document already in the registry/repository.

10.2.3 Document Transformation Option

In this option the Document Source or Integrated Document Source/Repository shall offer the ability to submit a document as a transformation of another document already in the registry/repository.

10.2.4 Folder Management Option

In this option the Document Source offers the ability to perform the following operation:

- Create a folder[1]

[1] The term “folder” comes from the medical community which commonly places patient records in folders for specific purposes. In computer science terminology this concept is most consistent with the UNIX directory format, where a file can be simultaneously within multiple directories.

- Add one or more documents to a folder

Note: In order to support document addition to an existing folder, grouping with the Document Consumer may be necessary in order to Query the registry (e.g. for UUIDs of existing folder).

10.3 Integration Profile Process Flow

A typical patient goes through a sequence of encounters in different care settings. In each care setting, the resulting patient information is created and managed by multiple care delivery information systems (EHR-CRs). Through a sequence of care delivery activities, a number of clinical documents are created. The EHR-LR provides the means to share the relevant subset of these documents, as they are contributed by the various EHR-CRs that are part of the same XDS Affinity Domain.

Example: Cardiac Patient Management Scenario

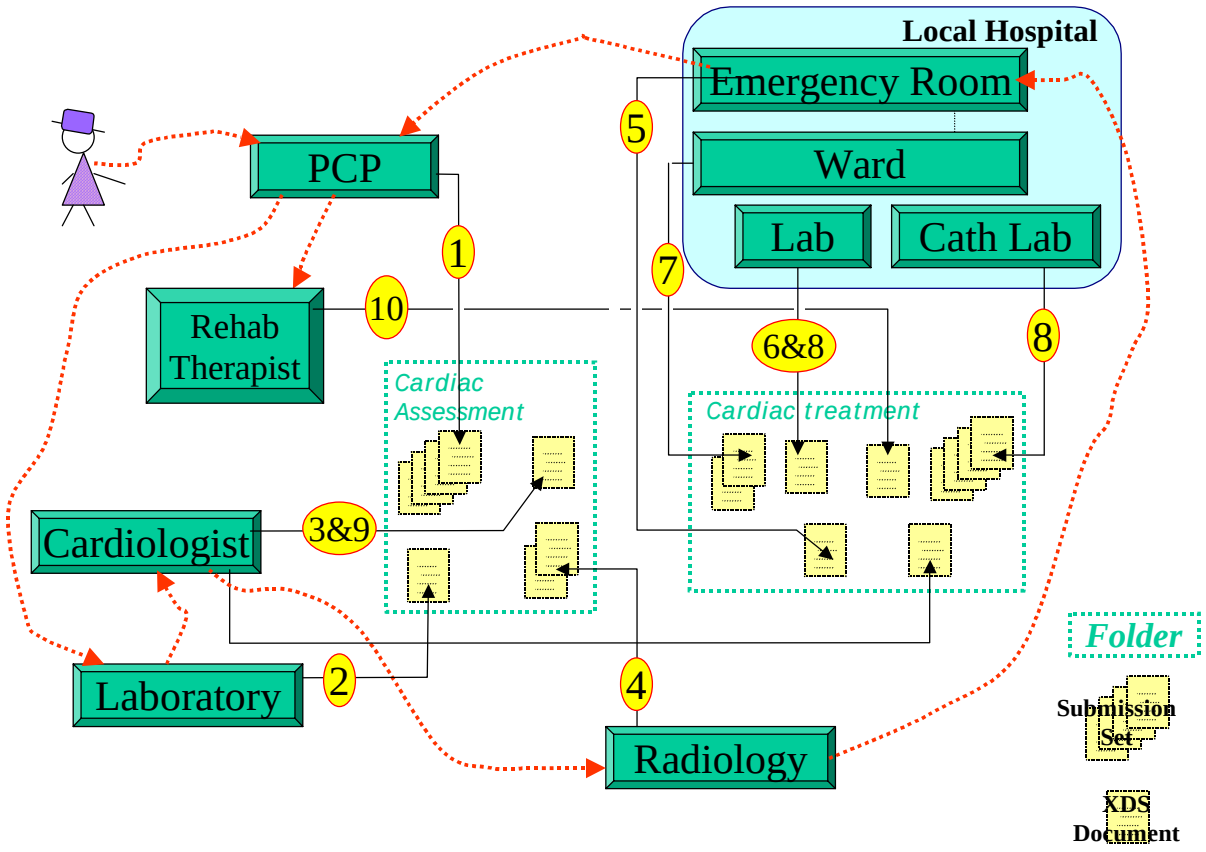


Figure 10.3-1 Cardiac Patient Management Scenario Transaction Process Flow

This scenario spans about 3 weeks of a patient's cardiac episode. The patient presents to her primary care provider (PCP) with complaints of shortness of breath, nausea, tiredness and chest pains. This doctor works closely with a local hospital that has recently established a cardiac care network that allows PCPs, cardiologists, laboratories and two local hospitals to share clinical documents to improve patient care. This cardiac network is part of a local care data exchange community that has been set-up in this community and to which the care plan to which this patient

belong has encouraged patients to subscribe. Our patient has been provided a health record account number.

1. During the patient examination, the PCP records the complaint, and determines that he should perform an ECG. He queries the cardiac care network to see if there are prior ECG reports (step 1 in Figure 10.3-2), using a coded document class "report" and a coded practice setting "cardiology" established by the cardiac care network for ECG reports. Among the matching Documents, he locates a prior ECG report that is then retrieved (step 2 in Figure 10.3-2). He compares the two results and determines that the patient should be referred to a cardiologist. He searches for additional reports in the cardiac care network (step 3 in Figure 10.3-2) for this patient, but finds none.

   Using the ambulatory EHR system, he creates a submission request onto the patients health record account number for a "PCP office visit" that includes a submission set consisting of three new documents (visit note, referral letter, new ECG report) and of one reference to the prior ECG report (step 4 in Figure 10.3-2). Following the Cardiology Network XDS Affinity Domain policy, he creates a "cardiac assessment" Folder to contain all four documents in order to facilitate collaboration with the cardiologist.

   The repository used by the ambulatory EHR system will then register the documents that are part of this submission request (step 5 in Figure 10.3-2).

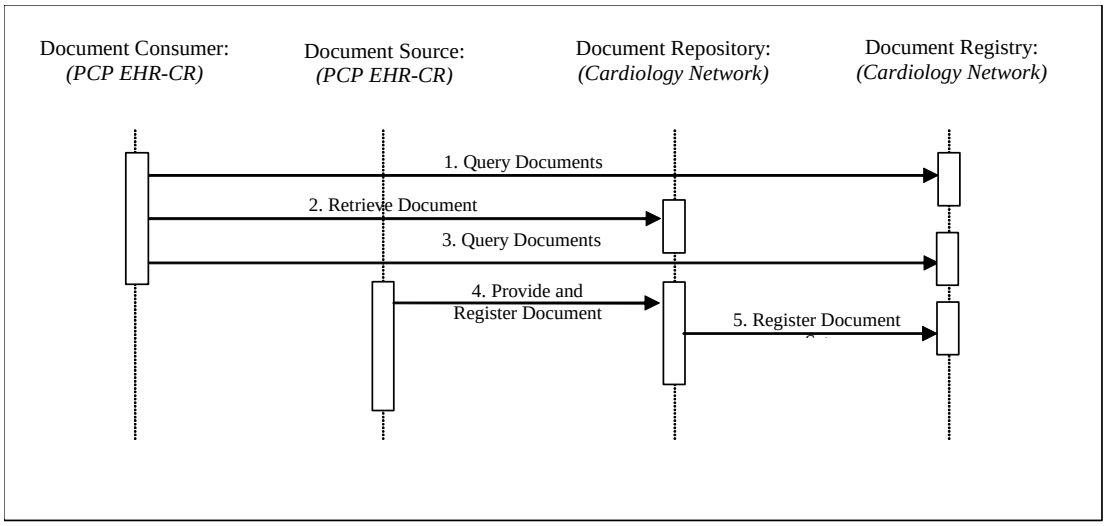


Figure 10.3-2 PCP Query Transactions Process Flow

The PCP EHR system implements the Document Consumer and Document Source actors to issue the Query, Retrieve and Provide & Register transactions as shown in Figure 10.3-2. The transactions are processed by the Document Repository and the Document Registry provided by the cardiology care network.

2. The patient appointment with the cardiologist is scheduled. The patient goes to the lab for the lab tests required before appointment. The lab creates a submission set with a clinical code of "laboratory tests" containing the lab results. The lab is not aware of the "cardiology assessment" folder.

3. The cardiologist sees the patient. He queries the repository for any patient's records in a "cardiac assessment" folder (step 1 in Figure 10.3-3). Available are the visit note from the PCP, the ECG and prior ECG, and the referral letter, which he retrieves and reviews (steps

2-5 in Figure 10.3-3). He also queries for recent lab reports, and finds the lab results (step 6 in Figure 10.3-3). This is also retrieved and reviewed (step 7 in Figure 10.3-3).

The cardiologist performs an ultrasound, dictates a visit note, and orders a nuclear stress test. The visit note and ultrasound images and report are registered as a "cardiologist office visit" submission set and placed in the "cardiac assessment" Folder. In addition, the lab report is added to the "cardiac assessment" Folder (step 8 in Figure 10.3-3).

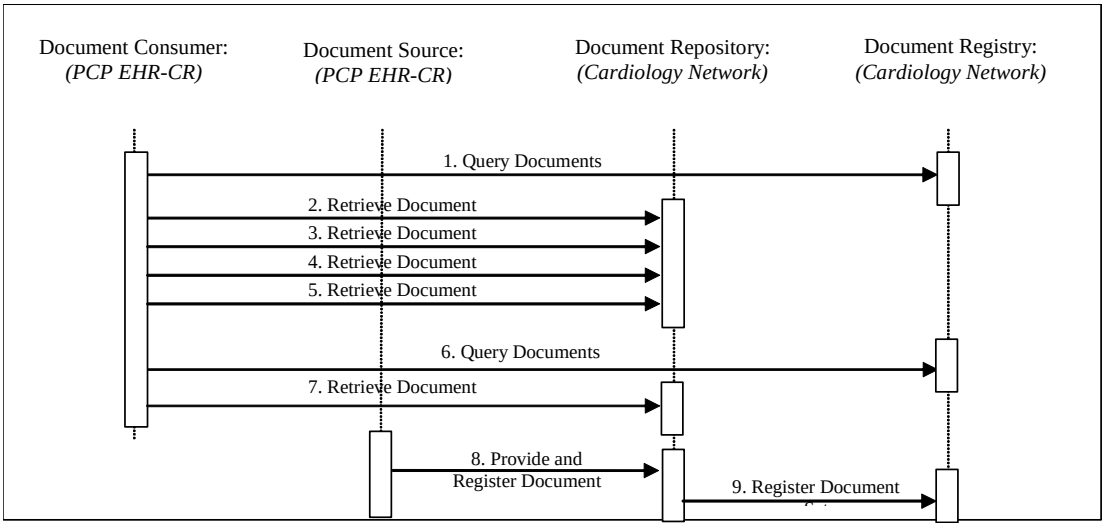


Figure 10.3-3 PCP Query Transactions Process Flow

4. The patient is seen at a radiology facility for the nuclear stress test. The test is performed, and the radiologist dictates the report. The nuclear stress test report is registered in a "radiology examination" submission set and associated with the "cardiac assessment" Folder

5. Although she has a scheduled appointment with her cardiologist in two days, she wakes up with severe chest pain. On the way to work, she decides to go to the emergency room (ER) of her local hospital.The ER doctor uses the hospital EHR system to query the cardiac care network registry and repositories for documents related to the patient in reverse chronological order (step 1 in Figure 10.3-4). Available documents from latest cardiology related Folder are the visit notes from the PCP and cardiologist, the recent and prior ECGs, the lab results, and the ultrasound images and report, and the nuclear stress test images and report.

   The ER doctor retrieves and reviews the two most relevant reports (step 2 and 3 in Figure 10.3-4).

   The ER doctor orders lab tests, ECG, and places the patient under monitoring. The lab tests and ECG are placed in the hospital EHR that acts as a Document Repository Actor for the cardiac network. Abnormal cardiac activity requires a catheterization, diagnostics and possibly intervention. The ER doctor admits the patient to the cardiology service and contacts the cardiologist.

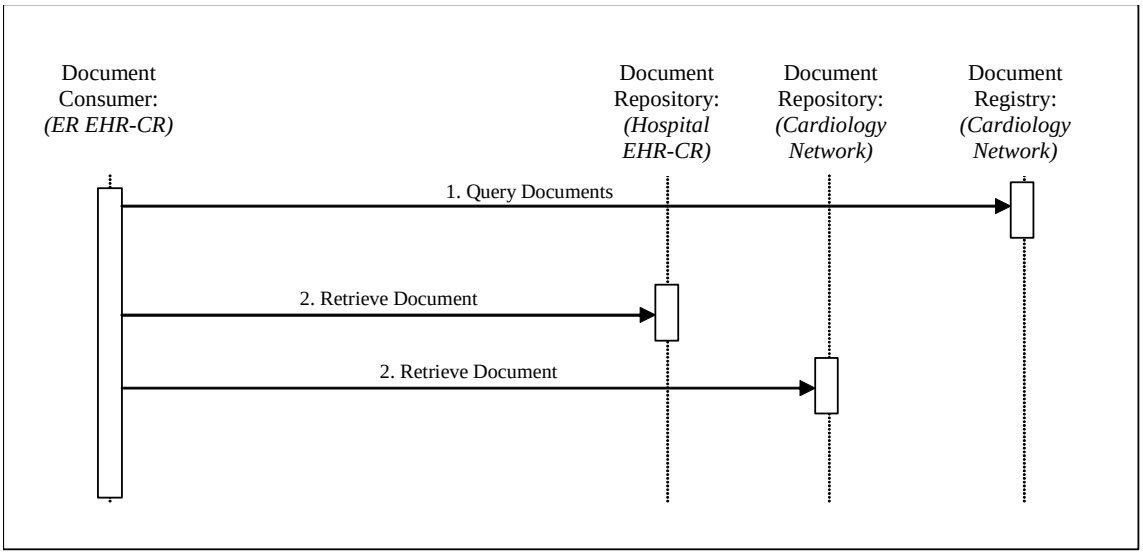


Figure 10.3-4 ER Query Transactions Process Flow

6. While talking to the ER physician, the cardiologist accesses the cardiac care network from his home office. He queries for all documents related to the patient since the last visit in his office. The nuclear stress test report that he did not previously review is available, along with lab results and ECG results from the ER. The two physicians determine a plan of care and the cardiologist makes arrangements to see the patient in the hospital.

7. As the patient is transferred from the ER, the ER visit notes are submitted as an "emergency department visit" submission set and placed in a newly created "cardiology treatment" Folder along with the earlier lab and ECG results.

8. The patient is transferred to an inpatient bed with the following sequence of events.
   - The patient is scheduled for a catheterization procedure in cath lab.
   - Additional lab tests are ordered and performed.
   - A diagnostics procedure is performed in cath lab.
   - An intervention with the placement of a stent is performed.
   - A cath intervention report is dictated.
   - Patient is returned to monitored care for recovery.
   - Education given to patient and family.
   - Discharge Summary dictated by cardiologist.
   - Cardiologist orders lab tests to be completed prior to scheduled follow-up visit.

   The admission assessment, lab results, cath intervention report and key images, and discharge summary form a "cardiology intervention" submission set, which is registered with the cardiac care network registry in the "cardiac treatment" Folder started by the ER.

9. The patient returns to the cardiologist for the post discharge follow-up visit. The resulting visit note, cardiac rehab and summary letters are placed in a "cardiology office visit" submission set and in the "cardiac treatment" Folder.

10. The patient goes to rehab sessions as scheduled by the cardiologist. The patient recovers and is seen by the PCP and cardiologist for routine visits.

10.4 General Principles

10.4.1 EDR-CR Concept

An EHR-CR or Care-delivery Record abstracts the information system or systems of a care delivery organization, which may support a broad variety of healthcare facilities: private practice, nursing home, ambulatory clinic, acute care in-patient facility, etc.

Typically a patient goes through a sequence of encounters in different care settings as depicted in the figure below.

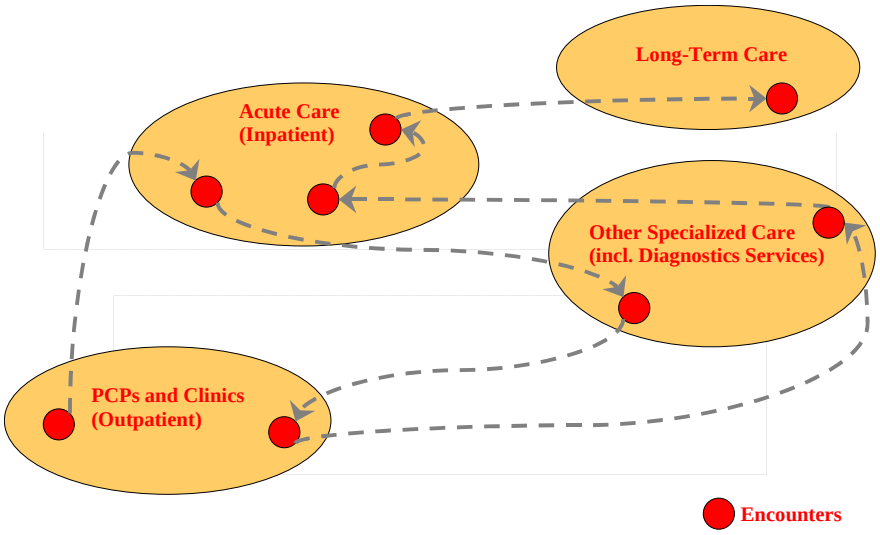


Figure 10.4.1-1 Sequence of encounters across care delivery organizations

It is out of the scope of this IHE Integration Profile to define or restrict the type of care provided, nor the internal workflow of a care delivery organization. The EHR-CR system participates only to the cross-enterprise clinical document sharing as Document Source and Document Consumer Actors according to the following principles:

1. EHR-CR as Document Source contributes documents in any one of the document formats that are supported by the XDS Affinity Domain (e.g. CDA Release 1, CDA Release 2 with specific templates, DICOM Composite SOP Classes, ASTM-CCR, CEN ENV 13606 etc).
2. This Profile does not require that the EHR-CR as Document Sources and Consumers store and manage their internal information in the form of documents as they are shared throughout the XDS Affinity Domain.
3. By grouping a Document Source with a Document Repository, an EHR-CR may leverage existing storage provide a unified access mechanism without needing to duplicate storage.
4. EHR-CRs as Document Sources and Consumers are responsible to map their local codes into the XDS Affinity Domain codes if necessary.

The XDS Documents shared by the EHR-CR and tracked by the XDS Registry form a Longitudinal Record for the patients that received care among the EHR-CRs of the XDS Affinity Domain.

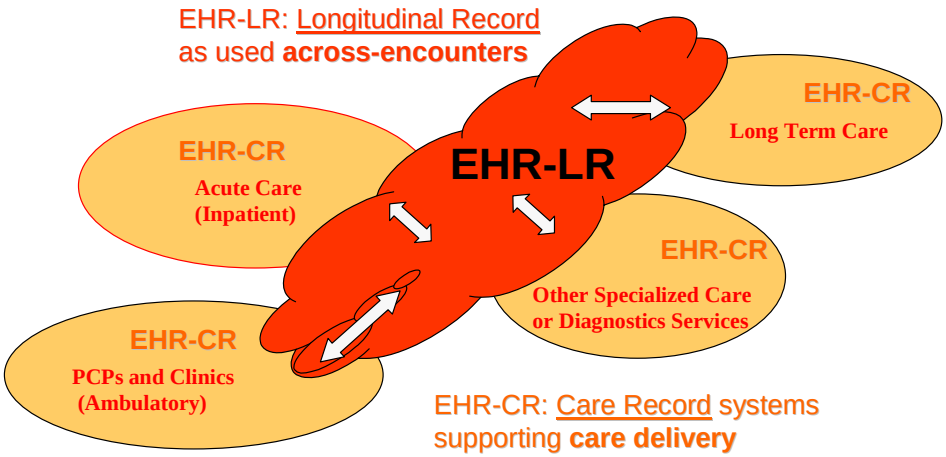


Figure 10.4.1-2 Contributing and sharing to a patients' longitudinal health record

This shared clinical record is called an EHR-LR in this Integration Profile.

10.4.2 XDS Document Concept

An XDS Document is the smallest unit of information that may be provided to a Document Repository Actor and be registered as an entry in the Document Registry Actor.

An XDS Document is a composition of clinical information that contains observations and services for the purpose of exchange with the following characteristics: Persistence, Stewardship, Potential for Authentication, and Wholeness. These characteristics are defined in the HL7 Clinical Document Architecture Release 1 specification. An XDS Document may be human readable (with the appropriate application). In any case, it should comply with a published standard defining its structure, content and encoding. IHE intends to define content-oriented Integration Profiles relying on such content standards to be used in conjunction with XDS.

The XDS Integration Profile manages XDS Documents as a single unit of information; it does not provide mechanisms to access portions of an XDS Document. Only the Document Sources or Document Consumers have access to the internal information of the XDS Document. When submitted for sharing, an XDS Document is provided to the Document Repository Actor as an octet stream. When retrieved through the Retrieve Document transaction, it shall be unchanged from the octet stream that was submitted.

The Document Source Actor is responsible to produce the metadata that will be submitted to the Document Registry Actor to form the XDS Document Entry that will be used for query purposes by XDS Consumer Actors. The Document Source maintains responsibilities over the XDS Documents it has registered. It shall replace XDS Documents that may have been submitted in error. See ITI TF-1: Appendix K for a more detailed discussion of the concept of XDS Document.

XDS Documents are required to be globally uniquely identified. See ITI TF-2: Appendix B for a definition of globally unique identifiers.

10.4.3 Submission Request

An XDS Submission Request is a means to share XDS Documents. It may be conveyed:

- by a Document Source Actor in a *Provide and Register Document Set Transaction* to the Document Repository Actor, or
- by a Document Repository Actor in a *Register Document Set Transaction* to the Document Registry Actor

An XDS Submission Request contains elements of information that will ensure the proper registration of XDS Documents. These are:

1. Metadata to be placed in Document Entries for new XDS Documents being submitted,
2. A Submission Set that includes the list of all new XDS Documents and Folders being submitted and optionally a list of previously submitted XDS Documents,
3. If desired, Folders to be created with the list of included XDS Documents (new document being submitted as well as previously submitted),
4. If desired, addition to previously created Folders of lists of XDS Documents (new document being submitted as well as previously submitted), and
5. Zero or more XDS Document octet streams for the new XDS Documents being submitted.

Following a successful Submission Request, new XDS Documents, Submission Set, and Folders included in the Submission Request are available for sharing in an XDS Affinity Domain. In case of failure to process a Submission Request, the Submission Set and any XDS Documents and Folders shall not be registered.

10.4.4 Submission Set Concept

An XDS Submission Set is related to care event(s) of a single patient provided by the care delivery organization EHR-CR performing the submission request. It creates a permanent record of new XDS Documents as well as pre-existing (i.e. already registered) XDS Documents that have a relationship with the same care event(s). It also includes the record of new XDS Folders creation.

An XDS Submission Set shall be created for each submission request. It is related to a single Document Source Actor and is conveyed by a single Provide & Register Document Set Transaction or a Register Document Set Transaction.

The Document Registry may be queried to find all documents registered in the same XDS Submission Set.

The same XDS Document, initially registered as part of a Submission Set, may also be referenced by later XDS Submission Set. This allows older documents relevant to the present care of a patient to be associated with more recent Submission Sets.

XDS provides complete flexibility to EHR-CRs to relate Documents and Submission Sets to an encounter, a visit, an episode of care, or various workflow processes within EHR-CRs.

10.4.5 Concept of Folder

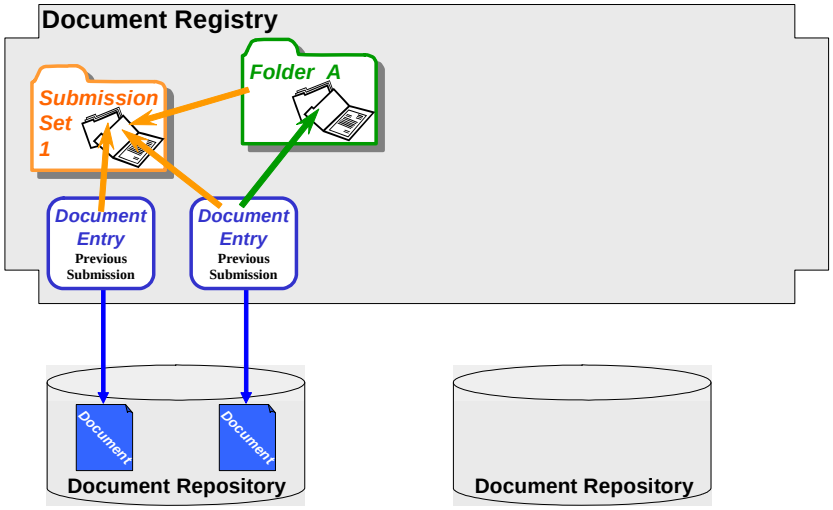
The purpose of an XDS Folder is to provide a collaborative mechanism for several XDS Document Sources to group XDS Documents for a variety of reasons (e.g. a period of care, a problem, immunizations, etc.) and to offer the Document Consumers a means to find all Document Entries placed in the same Folder. The following principles apply to an XDS Folder:

1. A Folder groups a set of XDS Documents related to the care of a single patient,
2. One or more Document Source Actors may submit documents in a given Folder,
3. A Folder may be created by a Document Source and/or predefined in an XDS Affinity Domain,
4. The content of a Folder is qualified by a list of codes/meaning,
5. Document Source Actors may find existing Folders by querying the Document Registry or by means outside the scope of XDS (e.g. Cross-enterprise workflow, such ePrescription, eReferral, etc),
6. Once created a Folder is permanently known by the Document Registry,
7. Placing previously existing Documents in Folders is not recorded as part of the Submission Set,
8. Folders in XDS may not be nested,
9. The same documents can appear in more than one Folder, and
10. Folders have a globally unique identifier.

10.4.6 Example of use of Submission Request, Submission Set and Folder

The sequence of figures below shows an example of a submission request that includes two new documents, a reference to a pre-existing document and the use of two folders. The first figure depicts the initial state of a Document Registry in which two Documents have been submitted where one is associated with a Folder A. The second figure depicts a submission request that adds two new documents, placing one of them into a pre-existing folder and the other one into a new Folder B.

Document Repository and Registry – Initial State



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Document Repository and Registry – Submission Request

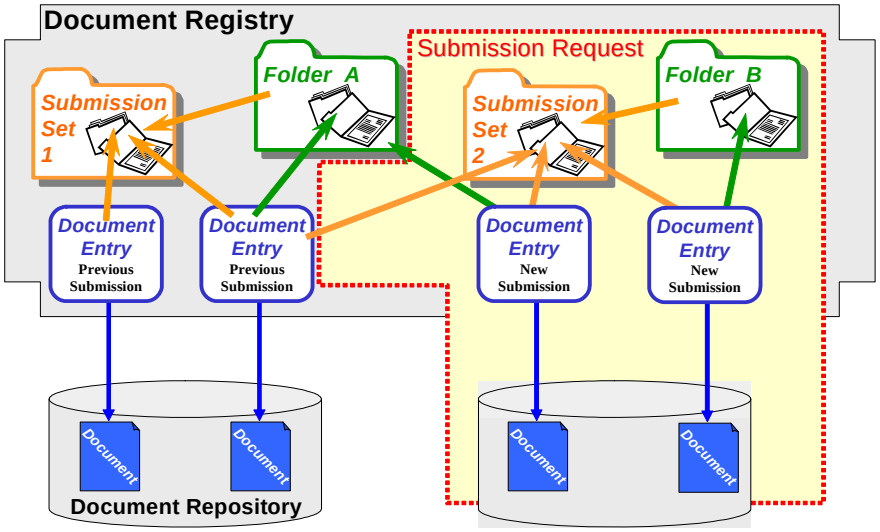

Figure 10.4.6-1 Example of a submission flow to an XDS Registry

From the above example, the contents of a Submission Set are shown by the figure below. The Document Entries associated with the Submission Set are logical part of the Submission Set.

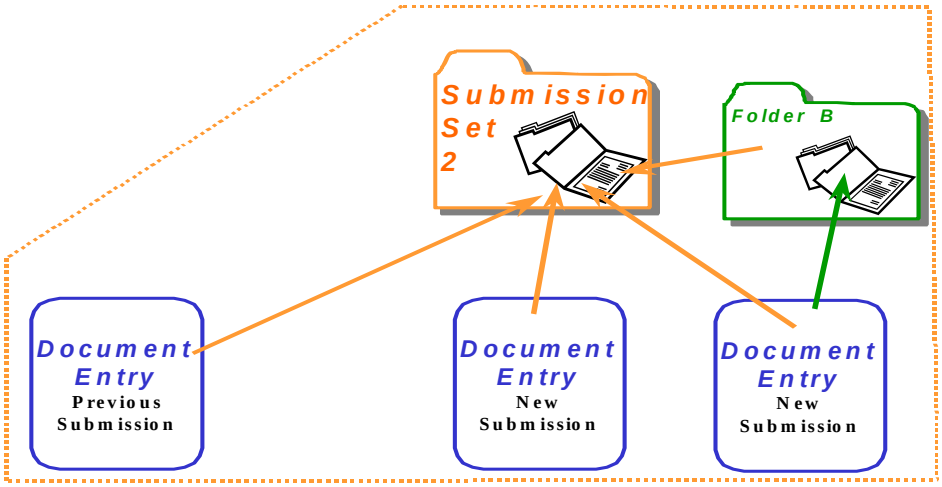

Figure 10.4.6-2 The logical content of a Submission Set

10.4.7 XDS Registry Data Model and Attributes

The XDS Integration Profile provides a means to place documents in a repository chosen by the Document Source, and also to place information about this document (or metadata) in an entry of the Document Registry that manages the XDS Affinity Domain.

The term metadata reflects that this information is "about" the documents. The purpose of well-specified document metadata is to enable a uniform mechanism for Document Consumers to locate clinical documents of interest much in the way a card catalog in a library helps readers find the book they want.

This section addresses the high-level data model in which the metadata is registered and against which queries of the XDS registry are performed. Then it presents the specific attributes that may be registered and used to filter the document entries of the registry.

10.4.7.1 XDS Document Registry Data Model

The following entities are used in the XDS Document Registry Data Model:

XDS Document Entry: Information entity managed by a Document Registry Actor that contains a set of metadata describing the major characteristics of an XDS Document along with a link to the Document Repository Actor where the actual XDS Document may be retrieved.

XDS Document: A stream of bytes stored in a Document Repository Actor and pointed to by an XDS Document Entry.

XDS Folder: A logical container that groups one or more XDS Document Entries in any way required (e.g. by source care delivery activities, by episode, care team, clinical specialty or clinical condition). This kind of organizing structure is used variably: in some centers and systems the Folder is treated as an informal compartmentalization of the overall health record; in others it might represent a significant legal portion of the EHR relating to the originating enterprise or team. The Folder is a means of providing organization of XDS Documents (or Composition in EHRCOM). The same XDS Document Entry may belong to zero or more Folders.

XDS Submission Set: When XDS Documents are registered by a Document Source Actor, they shall be included in one and exactly one Submission Set. An XDS Submission Set groups zero or more new XDS Documents and references to already registered XDS Documents to ensure a persistent record of their submission.

XDS Submission Request: A Submission Request includes one and only one Submission Set, zero or more new XDS Folders and assignment of XDS Documents into new or existing Folders. A Submission Request is processed in an atomic manner by the Document Repository and the Document Registry (i.e. all XDS Documents included or referenced in a Submission Set as well as the Folders and inclusion of Folders references are registered or none will). This ensures that they are all made available to Document Consumer Actors at the same time.

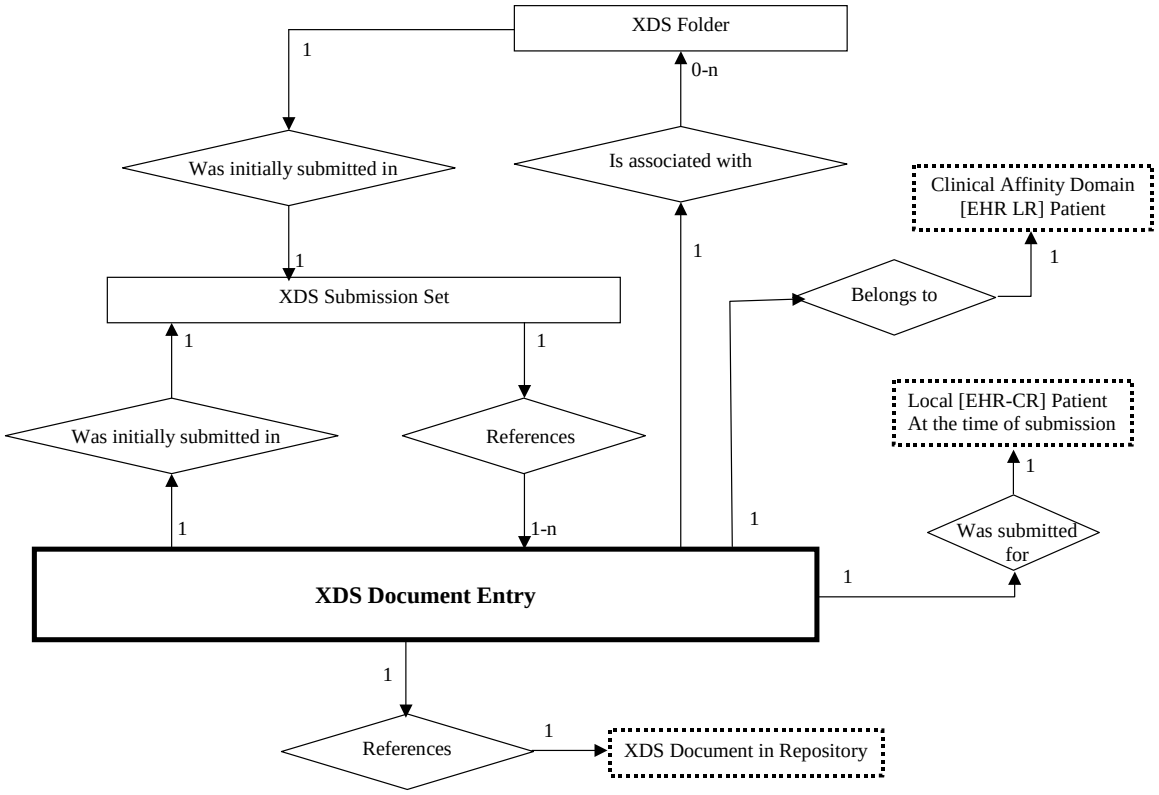

Figure 10.4.7-1 XDS Document Registry Data Model

10.4.7.2 Attributes of the XDS Document Entries

The specific attributes of each entity in the above registry data model have been selected from document header attributes from several standards (see ITI TF-2: Appendix L), including:

- ANSI/HL7 CDA R1-2000
- HL7 CDA Release 2 (draft) Document header definition (Dec 2003 Committee Ballot)
- Composition attributes from EHR ENV 13606 (draft).

XDS defines a well focused set of primary attributes that support the most common use cases to search the most relevant documents. These include:

Patient Id
Service Start and Stop Time
Document Creation Time
Document Class Code and Display Name
Practice Setting Code and Display Name
Healthcare Facility Type Code and Display Name
Availability Status (Available, Deprecated)
Document Unique Id

The three codes (Document Class, Practice Setting and Healthcare facility Type) are code set that are expected to generally include a limited number of values (between 10 and 100), thus ensuring a reasonably easy search capability.

A number of additional query attributes or attributes used to perform a secondary selection in order to decide to retrieve a specific document are also defined by this Integration Profile. At the Document Level, these include a fine grained Document Type (e.g. LOINC classification), a list of Event Code that can be used as key word, the document author and associated institution, the document relationship to manage replacement addendum and a variety of transformations, a confidentiality code, language code, etc.

The complete list of attributes and their definition is documented in the IHE ITI Register Transaction (see Volume II section 3.12).

10.4.8 Concept of an XDS Affinity Domain

An XDS Affinity Domain is an administrative structure made of a well-defined set of Document Source Actors, set of Document Repositories, set of Document Consumers organized around a single Document Registry Actor that have agreed to share clinical documents.

Note: Document Sources, Repositories and Consumers may belong to more than one XDS Affinity Domain and share the same or different documents. This is an implementation strategy and will not be further described.

Note: the XDS Integration Profile does not support the federation of XDS Affinity Domains. It is expected that a future IHE Integration Profile will address the cooperation of multiple Document Registry Actors serving different XDS Affinity Domains.

A number of policies will need to be established in an XDS Affinity Domain in order to ensure effective interoperability between Document Sources and Consumers. Some of the key technical policies include (A more extensive list of policy agreements that need to be made by XDS Affinity Domains is discussed in ITI TF-1: Appendix L):

1. The document formats that will be accepted for registration
2. The various vocabulary value sets and coding schemes to be used for the submission of metadata of document, submission set and folders registration.
3. The Patient Identification Domain (Assigning Authority) used by the Document Registry.

See ITI TF-1: Appendix K for a detailed discussion of the concepts of XDS Affinity Domain.

10.4.9 Patient Identification Management

Since the central focus of the DS Integration Profile is "sharing documents", it is critical that each document be reliably associated with the corresponding patient (Patient Id).

The XDS Document Registry is not intended to be an authority for patient identification and demographics information. This Integration Profile uses a Patient Identity Source Actor as the authoritative source of Patient Identifiers (master patient ID) for the XDS Affinity Domain.

Note: This Integration Profile can be easily extended to support a scenario where no master patient ID is defined (i.e. no Patient Identity Source for the XDS Affinity Domain). Such option, would requiring the use of federated patient identities at the time of query of the XDS Document Registry, may be expected as a future addition to this Integration Profile.

The following principles are defined:

1. The Patient Identifier Domain managed by the Patient Identity Source Actor in the XDS Affinity Domain is the source of patient identifiers (and merge operations) used by the XDS Document Registry to link Documents to a specific Patient. This Patient Identifier Domain is called the XDS Affinity Domain Patient Identification Domain (XAD-Pid Domain).
2. Submission Requests for Documents related to Patients with IDs not registered in the XDS Affinity Domain Patient Identifier Domain shall be rejected by the XDS Document Registry.
3. The XDS Document Registry will contain certain patient information (e.g. source patient ID, Surname, Given Name, Sex, Birthdate) for the purpose of audits and potential verification by Document Consumers. As this Integration Profile does not make any assumptions about the referential integrity and update of this information, these fields[2] shall not be used as query matching keys.
4. As XDS Document Sources and Consumers may belong to different Patient Identification Domains, these systems need to cross-reference their own local Patient ID to the corresponding patient ID in the XAD-Pid Domain of the Registry. Preferably, these systems may choose to use the IHE Patient Identifier Cross-referencing Integration Profile (See Appendix E.3) for this purpose.
5. The XDS Document Registry is responsible for validating Document metadata in accordance with the XDS Affinity Domain's policies. The Document Registry should reject submissions Requests that do not conform to these policies.

[2] It is possible to submit a new document to replace a previously submitted one, with a new document entry created in the registry to correct for errors in the submitted document in the original submission request. However this is not a mechanism that updates only the metadata, as the replaced document is only deprecated and remains pointed by the original metadata.

The figure below depicts an example of an XDS Affinity Domain with its Patient Identifier Domain (called XAD) and two EHR-CRs where the cross-referencing is performed internally to the Document Source and the Document Consumer Domains (Domain C and Domain D2 respectively).

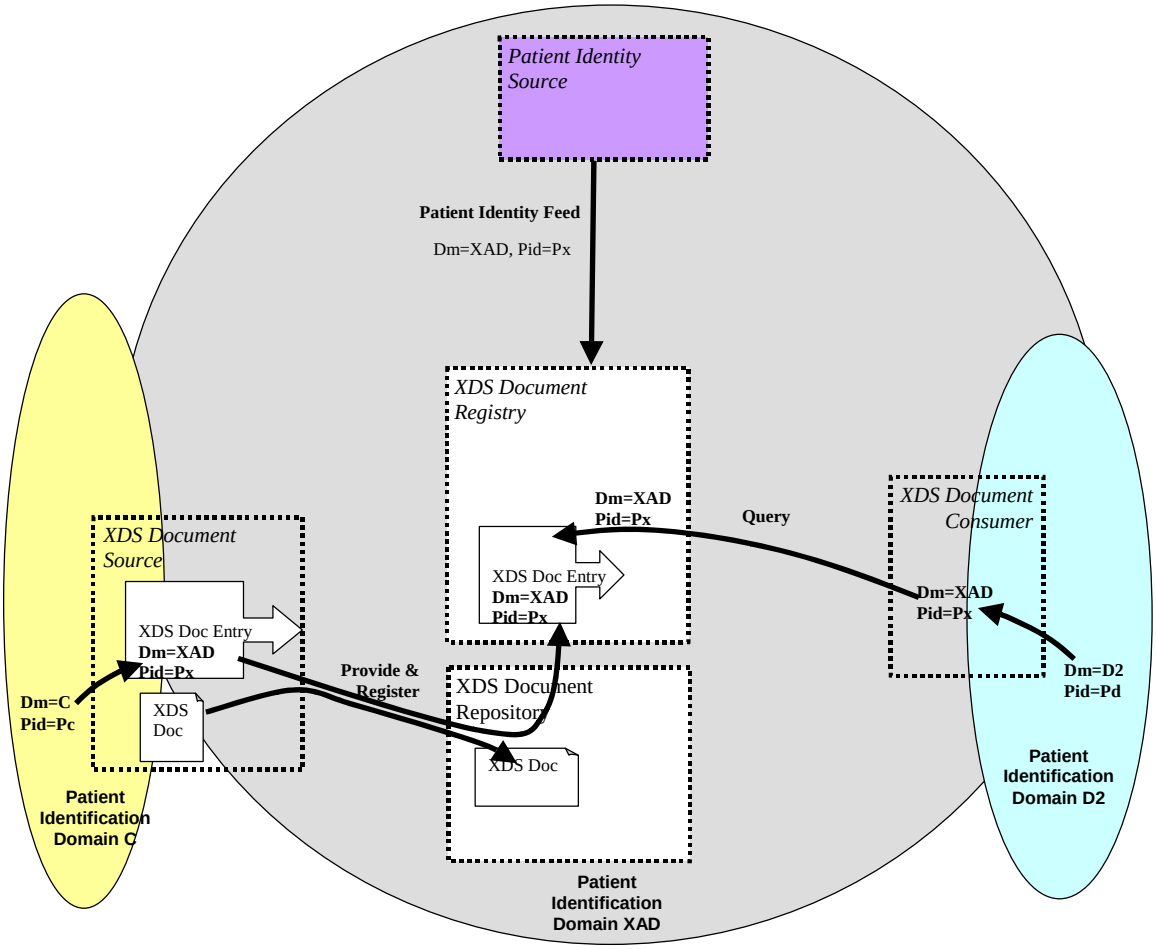

Figure 10.4.9-1 XDS Affinity Domain with patient ID cross-referencing internal to the EHR-CRs

10.4.10 Document Lifecycle

10.4.10.1 Document Availibility Status

Each XDS Document contained in a XDS Document Registry will be assigned one of the following Availability Status codes:

Approved: Available for patient care (assumes that it is authenticated, if applicable)

Deprecated: Obsolete, but may still be queried and retrieved

The XDS Document availability status is set to "approved" after the XDS Document Repository and the XDS Document Registry have successfully processed a submission request.

Note: ebXML Registry Services defines a Status of Submitted, which is used in a transient manner to provide an atomic submission. It is not significant to make this specific status externally visible.

An "approved" XDS Document may be changed to "deprecated" under the primary responsibility of its original Document Source with possible patient supervision. It is part of security policies that are beyond the scope of the XDS Integration Profile to have the XDS Repository/Registry enforce this ownership. The reason and responsible party for deprecating a document are tracked as part of the XDS Document Registry audit trail, which is a required capability. A "deprecated" Document remains available for Document Consumer queries. Except for the status change, a "deprecated" Document Entry metadata remains the same as when it was in the "approved" status.

An "approved" or "deprecated" XDS Document Entry may be deleted. This change is associated with the decision to completely remove a Document from an XDS Document Repository and the corresponding Document Entry from the XDS Document Registry. The XDS Affinity Domain shall establish the security policies associated with Document deletion. There are no transactions defined by this Integration Profile to support such operation.

See ITI TF-1: Appendix K for a detailed discussion of the concepts of XDS Document life cycle.

10.4.10.2 Document Relationships

XDS Documents may be related to predecessor documents by one of three methods:

- Replacement,
- Addendum
- Transformation
- Transformation-Replacement

These relationships between XDS Documents are tracked in the XDS Document Registry. The parent relationship attribute contained in the metadata of such Documents is a coded value that describes the type of relationship. An original Document has no parent and consequently its parent Id and parent relationship are absent. XDS Document Registry shall reject submissions that contain relationships to documents that are not registered or have been "deprecated". Document stubs are supported by XDS to allow for a valid relationship to a known but not registered Document.

A replacement document is a new version of an existing document. The replacement document has a new document Id; its parent Id attribute contains the document Id of the Document Entry associated with the previous version of the XDS Document, and parent relationship contains the code "RPLC". The Document Entry for the previous version shall have its Availability Status changed to "deprecated".

An addendum is a separate XDS Document that references a prior document, and may extend or alter the observations in the prior document. It modifies the parent document, but the parent document remains a valid component of the patient record and shall remain in the state "approved" or available for care. The addendum XDS Document metadata contains the identifier of the previous XDS Document version in parent Id, and its parent relationship contains the code "APND".

A transformed document is derived by a machine translation from some other format. Examples of transformed documents could be CDA documents converted from DICOM Structured Reporting (SR) reports, or a rendering of a report into a presentation format such as PDF. The transform XDS Document contains the document Id of the previous version in parentId, and its parent relationship contains the code "XFRM". XDS Affinity Domains may define rules that determine whether or not a transformed XDS Document replaces the source, but typically this would not be the case. If it is, an additional parent relationship of type "RPLC" is to be used.

10.4.11 Document Query

Query return info shall be either:

- a list of Registry Objects Values (e.g. XDS Document Entries)
- a list of Registry Objects UUIDs. This allows an XDS Document Consumer to receive a potentially long list of matching entries and to request them by subsets.

10.4.12 Transport Modes

The XDS Integration Profile defines an on-line mode of transport for all transactions except for the Provide & Register transactions where an off-line mode option is supported both for the Document Source and the Document Repository. In the "on-line mode" the transaction between two actors (computer applications) requires their simultaneous presence (e.g. an HTTP GET). In the "off-line mode" the transaction between the two actors (computer applications) does not require their simultaneous presence (e.g. a store and forward e-mail exchange).

1. An HTTP-based protocol (SOAP with Attachments) will be used for on-line operation.
2. The SMTP protocol will be used for off-line operation.

10.5 Implementation Strategies

The XDS Integration profile addresses the requirements of three major implementation strategies reflecting different groupings of actors within an EHR-CR as well as different configurations of the EHR-LR. This range of implementation strategies reflects the need to accommodate a variety of workflows and configurations. These implementation strategies may coexist in some environments. Other implementation strategies are possible.

- Strategy 1: Repository at the Source. A single information system acts as both the Document Source and Document Repository for the documents it creates and registers with the Document Registry
- Upon completion of a phase of care, an EHR-CR will register a submission-set of documents in a Document Repository Actor with which it is grouped (same system). Then it registers this set of documents (newly created and priors documents of interest) with the Document Registry Actor [2].
- Any other Document Consumer Actor in the XDS Affinity Domain may query the Document Registry Actor to find documents related to all phases of care for the patient [3]. It may choose to retrieve some of these documents from any Document Repository Actor [4].

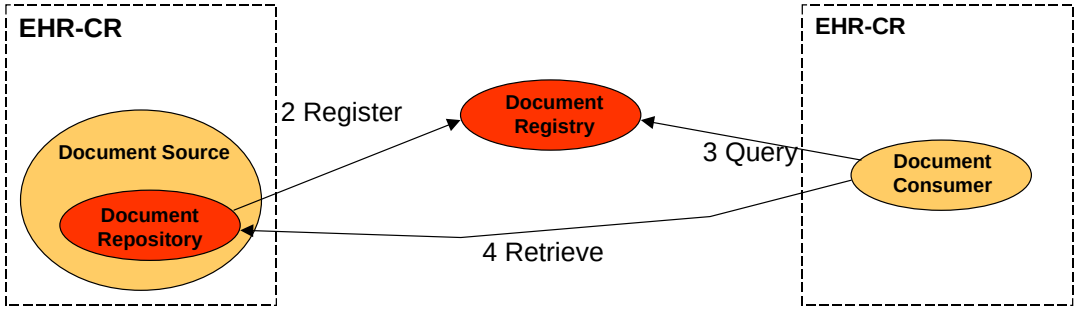

Figure 10.5-1 Implementation Strategy with Repository at the Source

- Strategy 2: Third Party Repository. The EHR-CR does not wish to be a Document Repository Actor, but rather uses the services of a third party Document Repository Actor to which it entrusts the documents it creates. First it provides both the metadata and the set of documents to this Document Repository Actor [1], which in turn forwards the registration request for the set of documents (newly created and prior documents of interest) to the Document Registry Actor [2].
- Any other Document Consumer Actor may query the Document Registry Actor to find out about documents related to all phases of care for the patient [3]. It may choose to retrieve some of these documents from any Document Repository Actor [4].

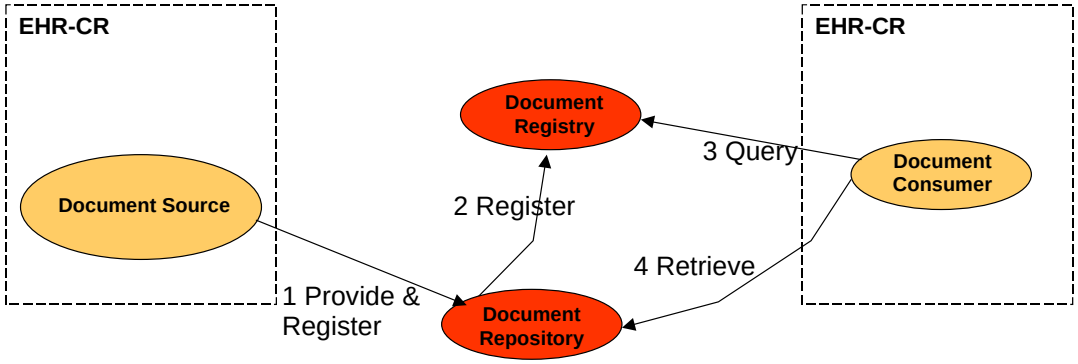

Figure 10.5-2 Implementation Strategy with 3rd party repository

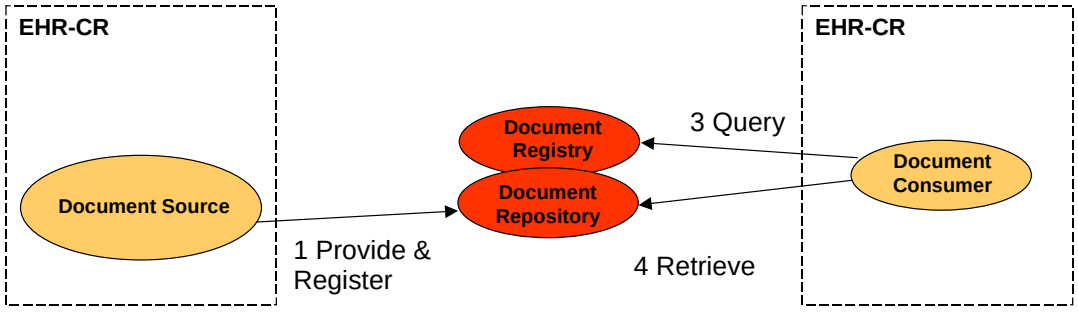

Table 10.5-3 Implementation Strategy with 3rd party central repository and registry

- Strategy 3: Direct Patient Transfer-Referral. The Document Source Actor completes a phase of care for a patient. It decides to directly provide and register [1] the set of documents (newly created and prior documents of interest) with a Document Repository [2] that has been grouped along with the Document Registry with the EHR-CR Document Consumer (Grouped Actors).
- In this case the span of the XDS Affinity Domain may be quite limited as it could be defined to cover only the two EHR-CRs. However the same transaction [1] applies. Note that, in this implementation strategy the other transactions, although supported by the actors, are not used by the Document Consumer since the Document Registry and Document Repository reside within the Document Consumer.

Figure 10.5-4 Direct patient referral with registry and repository at consumer

Patient access to an EHR-LR may be supported by a specialized EHR-CR (i.e. a portal) implementing the Document Source and Document Consumer Actors.

10.6 Patient Identifier Communication Requirements

ITI Transaction 8 described in ITI TF-2: 3.8 defines the format requirements for the patient identifier in PID-3. Specifically, the value for PID-3.4, Assigning Authority can be omitted, expressed using the first subcomponent (namespace ID) or the second and third subcomponents (universal ID and universal ID type). These rules shall apply in this profile:

1. If the Patient Identity Source does not include a value for PID-3.4, Assigning Authority, then
    a. PID-3, Patient Identifier List, is constrained to include one entry referring to one identifier.
    b. The Patient Identity Source and Document Registry shall agree that all messages from this source shall refer to a single assigning authority.
2. If PID-3.4 does contain a value for PID-3.4, Assigning Authority, then
    a. The Patient Identifier Source may send multiple patient identifiers with properly formatted components. The Document Registry shall be responsible for selecting the one identifier from the Patient Identifier List (not necessarily in the first position) that is too used to register the selected patient.
    b. As specified in ITI TF-2: 3.8, the value for PID-3.4, Assigning Authority, can be expressed using the first subcomponent (namespace ID) or the second and third

subcomponents (universal ID and universal ID type). Both methods shall be accepted by the Document Registry and shall be considered as equivalent.

ITI Transactions 14, 15 and 16 express patient ID as a string that is not parsed using typical HL7 parsing logic; please refer to requirements for Patient ID in those transactions. Document Registry actors will have to map between the Patient ID feed provided in ITI-8 as described above and the PID provided by those transactions in this profile.

11 Personnel White Pages (PWP)

The Personnel White Pages (PWP) Profile provides access to basic directory information on human workforce members to other workforce members within the enterprise. This information has broad use among many clinical and non-clinical applications across the healthcare enterprise. The information will be used to

1. enhance the clinical workflow
   a) contact information,
   b) phone numbers,
   c) email address
2. enhance the user interface
   d) displayable names,
   e) titles

This Personnel White Pages Profile specifies a method of finding directory information on the User Identities (user@realm) supplied by the Enterprise User Authentication (EUA) Integration Profile. This Profile assumes but does not define access controls, and audit trails. The use of the PWP Profile is intended for use within a healthcare enterprise. Extension to support sharing of the PWP between healthcare enterprises is possible but not fully addressed by this profile. The PWP profile is the first step on an IHE roadmap that includes Digital Certificates, Encryption, Digital Signatures, Medical Credentials, and Roles.

The directory need not support use cases beyond healthcare operations (e.g. Human Resource Operations), but does not forbid a properly designed overlap with other use cases. This profile does not intend for patients or other individuals that are not acting as part of the human healthcare workforce.

11.1 Actors/ Transactions

Figure 11.1-1 shows the actors directly involved in the PWP Integration Profile and the relevant transactions between them. Other actors that may be indirectly involved due to their participation in EUA profile are not necessarily shown.

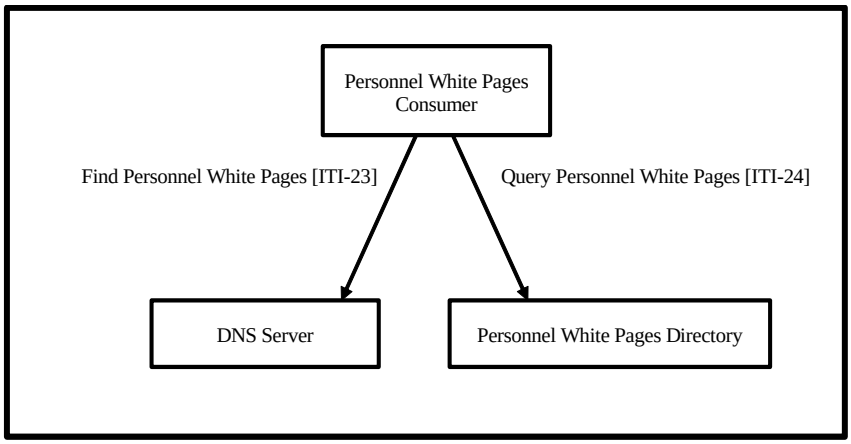

Figure 11.1-1: Personnel White Pages Profile Actor Diagram

Table 11.1-1 lists the transaction for each actor directly involved in the PWP Profile. In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled "R"). Transactions labeled "O" are optional. A complete list of options defined by this Integration Profile and that implementations may choose to support is listed in Section 11.2.

Table 11.1-1: PWP Integration Profile - Actors and Transactions

Actors	Transactions	Optionality	Section in Vol. 2
Personnel White Pages Consumer	Find Personnel White Pages	O	ITI TF-2: 3.23
	Query Personnel White Pages	R	ITI TF-2: 3.24
DNS Server	Find Personnel White Pages	R	ITI TF-2: 3.23
Personnel White Pages Directory	Query Personnel White Pages	R	ITI TF-2: 3.24

11.2 PWP Integration Profile Options

Options that may be selected for this Integration Profile are listed in the table 11.2-1 along with the Actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 11.2-1 PWP Integration Profile - Actors and Options

Actor	Options	Vol & Section
Personnel White Pages Consumer	*no option*	
DNS Server	*no option*	
Personnel White Pages Directory	*no option*	

11.3 PWP Integration Profile Process Flow

The Personnel White Pages Profile addresses the following use cases:

- A Clinical user logs into an acquisition device that is acting as a Personnel White Pages Consumer. The clinical application queries the DNS Server Actor using [ITI-23] to find the Personnel White Pages Directory. The clinical application then queries [ITI-24] the Personnel White Pages Directory using the user's username and displays the user's full name with First Name, Middle, and Last. There are information fields to support both European and Asian naming conventions.

- The Clinical user acquires clinical data. The application queries [ITI-24] the Personnel White Pages Directory for the user's demographics to include the user's organization identification to embed in the data record.
- The User then needs to send this report by means of email to a colleague. The application allows the user to search [ITI-24] the Personnel White Pages Directory for the destination user, and selects the destination user's email address.
- The User reviews an existing clinical report and finds initials have been recorded in the report. The user system does a query [ITI-24] of the Personnel White Pages Directory for the initials found in the report and the system displays the displayable name(s).

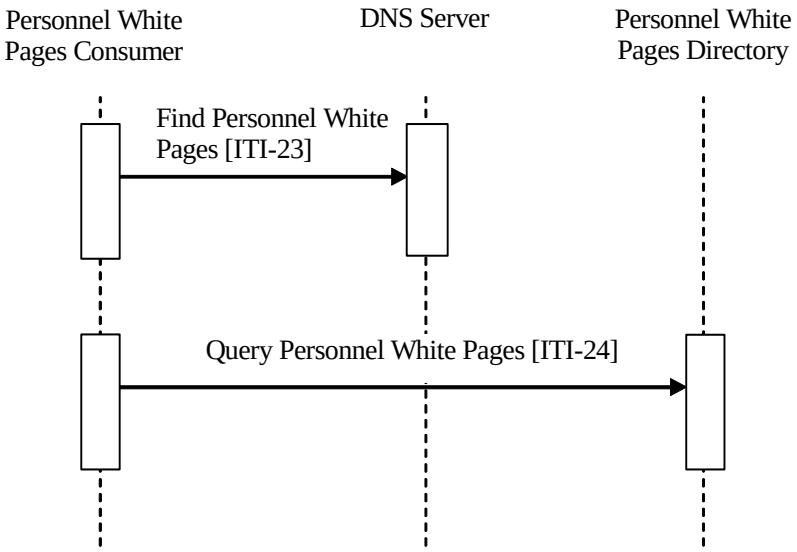

Figure 11.2-1: Basic Process Flow in PWP Profile

12 This is reserved for Notification of Document Availability (NAV)

13 Cross Enterprise User Assertion (XUA) Integration Profile

The Cross-Enterprise User Assertion Profile (XUA) provides a means to communicate claims about an authenticated principal (user, application, system...) in transactions that cross enterprise boundaries. To provide accountability in these cross-enterprise transactions there is a need to identify the requesting user in a way that enables the receiver to make access decisions and proper audit entries. The XUA Profile supports many solutions including enterprises that have chosen to have their own user directory with their own unique method of authenticating the users, and others that have chosen to use a third party to perform the authentication.

There are transactions defined by IHE that cross enterprise boundaries and are web-services based on ITI TF-2:Appendix V. The existing IHE profiles for an authenticated user identity (IHE Enterprise User Authentication Profile [EUA]) are not intended to function in cross-enterprise transactions. In a cross-enterprise environment it is more likely that the transactions will be going between two enterprises that maintain their own independent user directories (IHE Personnel White Pages [PWP]). This type of requirement is the focus of the Identity Federation standards. Identity Federation has received much attention by the security and the platforms industry. Identity Federation is agnostic to the type of user directory; it allows for a centralized user directory, but also supports the more powerful federation of user directories. Identity Federation supports:

- A Country that delegates the provisioning of all users into a single assigning authority domain (e.g., France) and provides a common service that handles all user authentication requests
- Support for centralized user directories
- A Region that knits together a network of cooperating hospitals and clinics where each hospital/clinic manages its own users.
- Support for distributed user directories
- Patients who wish to use an identity provider of their choosing (e.g. ISP, email provider).
- Support for non-healthcare specific user directories
- A Hospital that provisions users by issuing identity badges with picture and name printed, RFID for building access, and smart-card for strong authentication
- Support for claims about the method used to authenticate the user (e.g. strong authentication methods such as smart-cards)
- A Small clinic in a rural setting that supports a dozen users.
- Support for small scale systems (e.g., user at a kiosk, system using simple passwords)
- A General practice doctor retrieving results of a test performed by an outpatient clinic, where the outpatient clinic wants to have an audit trail specific to the user requesting the information.
- Support for the service provider to get a user identity for audit log purposes
- An automated System, based on a scheduled procedure, that is capable of being a delegate for a doctor pre-fetches the available documents so that it can determine a relevant few documents to offer to the doctor when the patient arrives

The XUA Profile leverages Web-Services Security, SAML 2.0 Token Profile and the various profiles from W3C, OASIS, and WS-I to support identity federation. In this way we will be able to take advantage of the vast experience of the communities outside of healthcare standards. This

profile leverages the experience of programs around the globe that have started work with SAML in healthcare.

13.1 Use Cases

The XUA profile supports complex environments, for example one where two different trust domains are operating under different technology, procedures, role-models, etc. They are cooperating in the XDS Affinity domain under an overarching trust relationship policy (See Vol 2, Appendix L) that indicates that these differences can be rationalized. The XDS transactions are transferring control from one entity to another, for example, when using XDS to exchange data between a single doctor practice and large multi-site hospital. It is not likely that they will all agree to the same access control model (organizational roles, functional roles, workflows, permissions, etc). It is not necessary to have the same access control across these entities, but it is reasonable that at the policy level they will agree to a set of processing rules. This illustrates an important fact that the XUA is useful for security audit logging, but is to a lesser extent useful for access controls.

The following is a list of use-cases that have been proposed for XUA. Some of these use-cases will not be supported due to lack of standards or sufficient guidance on the proper solution.

1. Country that provisions users into a single assigning authority domain (e.g., Germany) and handles all user authentication requests
    - Support for centralized user directories
2. Region that knits together many competing hospitals and clinics where each hospital/clinic manages its own users.
    - Support for distributed user directories
3. Patients who wish to use their email provider as their authentication authority uses a PHR-like application to access their own information in an XDS Affinity Domain.
    - Support for non-healthcare specific user directories
4. Hospital that issues identity badges with picture and name printed, RFID for building access, and smart-card for strong authentication
    - Support for claims about the method used to authenticate the user (e.g. strong authentication methods such as smart-cards)
5. Small clinic in a rural setting that supports a dozen users.
    - Support for small scale systems (e.g., user at a kiosk, system using simple passwords)
6. General practice doctor who retrieving results of a test performed by an outpatient clinic, where the outpatient clinic wants to have an audit trail specific to the user requesting.
    - Support for the service provider to get a user identity for audit log purposes
7. System, based on a scheduled procedure, pre-fetches the available documents so that it can determine a relevant few documents to offer to the doctor when the patient arrives.
    - Support for identifying the user as the system for tasks that are not initiated by a human user

8. User using Registry or Repository where the service provider wants to be assured that the user has been authenticated to a specific assurance level. This is not a case of not trusting the system, but recognition that the requester supports different levels of authentication. For example the system supports a proximity card as a form of authentication, as well as Smart-Card with PIN. This is not a replacement for ATNA access controls which give distributed access controls.

    - User Identity with level of assurance of that identity is needed.

9. Specialized XDS Affinity Domain for Emergency Dataset. In this case the transfer of information to the XDS Consumer is not critical to fully control, and thus the administration is willing to accept requests from any system as long as they can provide a user-assertion from a trusted source. This trusted-source may be a specialized identity provider for First Responders. (See RSA Pilot)

    - In this case only a user identity with proper linkage to a trusted identity provider is needed. No specific attributes are needed.

10. User acting in an identified clinical role accesses the Registry where the Registry wants to know the user identity and the role they are acting in to record the identity and role in the audit log.

    - Support inclusion of functional roles as named vocabulary
    - The Role of the user as the data subject (patient)

11. Service provider wants to enforce some form of access controls based on the user identity and/or functional role.

    - Support for the service provider to augment access controls based on some non-specified rules that are applied to the user and/or functional role

12. Access to a document by an individual that can't be identified because the Assertion Provider is not accessible

13.2 XUA Development

The vast majority of the use-cases (items 1-11) rely on claims about an authenticated identity, which a SAML 2.0 Identity Assertion can provide. This is a mature standard produced by OASIS. XUA Profile is focused on Web-Services transactions that follow ITI TF-2:Appendix V. XUA specifies that when a Cross-Enterprise User Assertion is needed, these Web-Services transactions will additionally use the Web-Services Security header with a SAML 2.0 Token containing the identity Assertion. As with any IHE profile, the applications are not forbidden to use other methods of providing the principal (user) identity, providing that interoperability has been assured through some policy.

A very clear need on all the use-cases is the recording of the user identity in any security audit logs. The XUA profile does not define these auditable events. The need to record a security audit event is driven by the grouped transactions (e.g., XDS.b Registry Stored Query, and XDS.b Retrieve Document Set). XUA does specify how to reference the Identity Assertion in an ATNA Audit Message.

The method of authenticating the principal (user) and the method that the X-Service User Actor (e.g. XDS.b Document Consumer) uses to get the Identity Assertion are outside the scope of this profile.

There are principal (user) attributes that appear to be needed in the use-cases: Doctor, Patient, Guardian, Emergency-Access. The Identity Assertion can contain attributes about the principal (user). At this time it is not clear what standards to use to identify these attributes and their values, so this is left to specific implementations that have defined a local vocabulary or vocabulary translation.

The method used by the X-Service User (e.g. XDS.b Document Consumer) Actor to determine the contents of the Identity Assertion is outside the scope of this profile. This might be accomplished using the SAML Metadata and WS-Policy.

It is expected that extending this solution to HL7 and DICOM will be supported in the future.

13.4 Actors/Transaction

Figure 13.4-1 shows the actors directly (Bold and Solid Boxes) involved in the XUA Integration Profile and the relevant transactions between them (Bold and Solid Line). The diagram also shows ancillary actors (Dashed and Grey Boxes) that are not profiled but include interactions (Dashed and Grey Lines). Actors grouped with are shown as the dashed line between the X-Service User and the X-Service Provider.

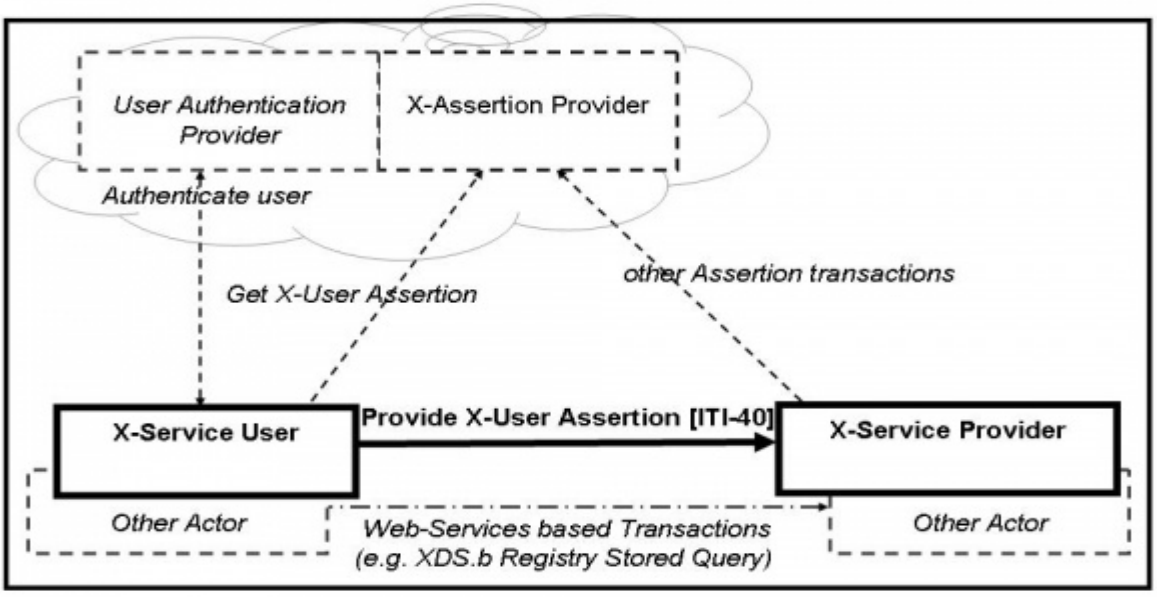

Figure 13.4-1 Cross-Enterprise User Assertion Actor Diagram

Table 13.4-1 lists the transactions for each actor directly involved in the XUA Profile. The ancillary actors and associated transactions may be supported by various technologies and system configurations varying from internal shared services to infrastructures for identity management.

In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled "R"). Transactions labeled "O" are optional. A complete list of options defined by this Integration Profile and that implementations may choose to support is listed in Section 13.5.

Table 13.4-1 XUA - Actors and Transactions

Actor	Transaction	Optionality	Section
X-Service User	Provide X-User Assertion	R	ITI-40
X-Service Provider	Provide X-User Assertion	R	ITI-40

13.5 Options

Options that may be selected for this Integration Profile are listed in the table 13.5-1 along with the Actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 13.5-1 XUA - Actors and Options

Actor	Option	Section
X-Service User	*None*	
X-Service Provider	*None*	

13.6 Grouping

13.6.1 Audit Trail and Node Authentication (ATNA)

The X-Identity Assertion is valuable and must be protected against confidentiality risks. In some Profiles (e.g. XDS), there is already an inherited requirement to group with IHE-ATNA Secure Node Actor. This grouping forces the network transactions to utilize mutually authenticated and encrypted TLS. This is leveraged by XUA to support the protection of the X-User Assertion to some risks to confidentiality and integrity. When ATNA Secure Node grouping is not required, there will need to be some other mechanism to protect the Provide X-User Assertion.

Volume 2 includes encoding rules for representing an X-User Assertion in an ATNA Audit Message.

13.6.2 Cross-Enterprise Document Sharing (XDS)

When an XDS.b Document Consumer is grouped with X-Service User Actor, the XDS.b Document Consumer shall conform to all the requirements in the Provide X-User Assertion Transaction. The Document Consumer will obtain a properly scoped XUA Assertion targeted for the XDS.b Document Registry or XDS.b Document Repository. The method used may be through internal means, SAML 2.0 Core protocols, WS-Trust, or any other means.

The XDS.b Document Registry and XDS.b Document Repository when grouped with the XUA X-Service Provider Actor shall conform to all the requirements in the Provide X-User Assertion Transaction. The XUA Profile does not constrain how the Assertion can be used (e.g. ignored, access control, etc).

13.6.3 Enterprise User Authentication (EUA)

An application that groups EUA and XUA Actors may use WS-Trust to get the X-User Assertion from the Security Token Service (STS). In this case the AuthnContextClassRef element of the SAML assertion shall be:

```
urn:oasis:names:tc:SAML:2.0:ac:classes:Kerberos
```

This conversion from one security token format to another is documented in the WS-Trust standard, and not further profiled by IHE.

13.6.4 Any Web-Services Transaction that leverages ITI TF-2: Appendix V

Any Actor that uses Web-Services according to ITI TF-2:Appendix V may be grouped with the appropriate XUA Actors. The Actor grouped with X-Service User Actor, the Requesting Actor, shall conform to all the requirements in the Provide X-User Assertion Transaction. The method used may be through internal means, SAML 2.0 Core protocols, WS-Trust, or any other means. The actor grouped with the X-Service Provider Actor shall conform to all the requirements in the Provide X-User Assertion Transaction. The XUA Profile does not constrain how the Assertion can be used (e.g. ignored, access control, etc).

13.7 Process Flow

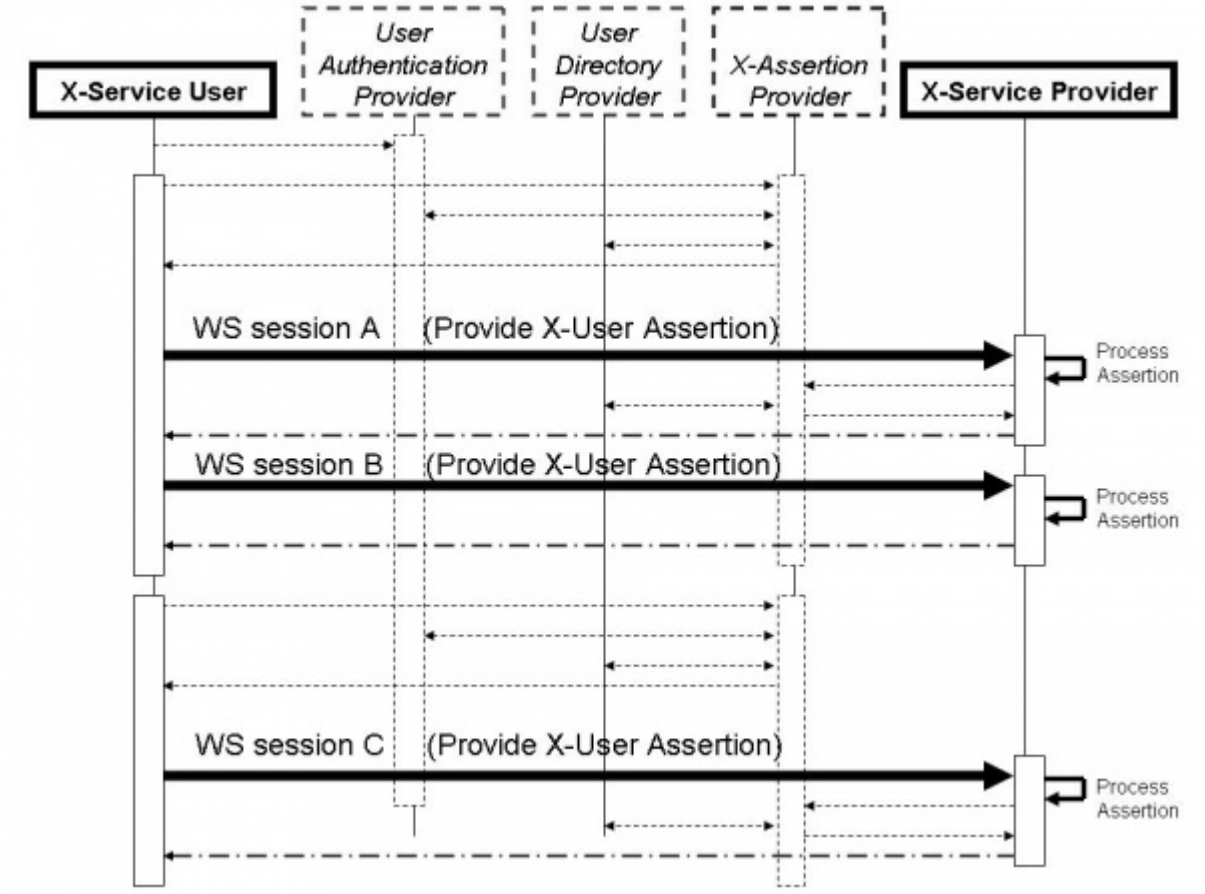

Figure 13.6-1 Cross-Enterprise User Assertion Process Flow

In the above flow we are showing more actors than are specified in this profile. This is a diagram showing a possible grouping with IHE-EUA (User Authentication Provider), IHE-PWP (User Directory Provider), and a SAML Identity Provider (X-Assertion Provider). The User Authentication Provider, User Directory Provider and X-Assertion Provider are not profiled here, but rather are shown to give a context to the XUA transactions.

In this figure the dark lines represent the X-User Assertion transaction. The dashed lines represent other standards based transactions that may be used. Web-Services session A and B show an example where one X-User Assertion is used to cover two Web-Services transactions, where Web-Services Session C is using a different X-User Assertion. This may be due to a different user, timeout of the previous X-User Assertion, or some other reason.

13.8 Security Considerations

The security risk assessment for XUA enumerates assets, threats, and mitigations. The security risk assessment for the Actors that are grouped (e.g. Registry Stored Query and Retrieve Document Set) with the XUA Actors are out of scope of the XUA profile, please look at those transactions for the Security Considerations. The complete risk data are stored and available from IHE. The purpose of this risk assessment is to notify vendors and healthcare providers of some of the risks that they are advised to consider in implementing XUA Actors. For general IHE risks and threats, please see ITI TF-1: Appendix L. The vendor is also advised that many risks can not be mitigated by the IHE profile and instead responsibility for mitigation is transferred to the vendor, and occasionally to the affinity domains, individual enterprises and implementers. In these instances, IHE fulfills its responsibility to notify affected parties through the use of the following sections.

14 Patient Administration Management (PAM) Integration Profile

14.1 Patient Administration Management Use Cases

The Patient Administration Management Integration Profile defines transactions based on message exchanges to support patient identity and encounter information, as well as movements within an acute care encounter. These can be represented by the following use cases.

14.2 Patient Identity Management Use Case

A Patient Registration application decides to create a new patient John Smith, based on patient information input from Hospital Sun. At this time, however, there is a limited set of personal information traits of John Smith available. His date of birth, home address, and home phone number are unknown. The registration application creates the patient identity and sends a Patient Creation message to its downstream applications with the set of known personal information traits.

The next day, detailed personal information about John Smith becomes available. The registration application updates its patient identity record, and sends out a Patient Update message.

After a week, the registration application creates a temporary patient identity John Doe based on input from Imaging Center Moon. After reconciliation of the temporary patient, it updates John Doe’s demographics to (a new instance of) John Smith, and changes the temporary Patient Identifier originally assigned to a permanent identifier.

After human inspection, it turns out that these two identities of John Smith represent the same person. The operator decides to merge the second identity to the previously established identity John Smith. A Patient Merge is communicated downstream.

14.2.1 Patient Encounter Management Use Case

Patient Alan Alpha arrives for an annual exam at a clinic. The registration system sends the patient registration information to the local ancillary systems, and the affiliated hospital's ADT system.

The exam of Alan Alpha reveals a serious condition, and an immediate hospital admission is recommended. Alan Alpha is referred to the affiliated hospital for admission. He is pre-admitted in the hospital for relevant diagnostic tests. The tests confirm the condition, and the patient is admitted in the hospital's ICU. During the stay in the ICU, the patient's insurance is verified, and the updated information is sent from the hospital’s ADT system to the hospital’s ancillary systems.

After a day in the ICU, Alan Alpha’s condition has improved, and he is transferred to a regular bed. The nurse recording the transfer makes a mistake, and enters the wrong room and bed. After discovering the error, the transfer is canceled, and the correct transfer is recorded. The patient is now recovered and about to leave the hospital. According to the hospital's procedures, he is transferred to an outpatient unit for administering follow-up tests. The patient is registered in the Hospital Outpatient Registration System.

The outpatient encounter of Alan Alpha is completed; based on satisfactory test results, he is discharged from the hospital and the Outpatient Registration system.

In this use case, two patient encounter management systems (the hospital ADT system and the hospital Outpatient Registration system) cooperate as peers.

14.2.2 Actors/ Transactions

Figure 14.1-1 shows the actors directly involved in the Patient Administration Management Integration Profile and the relevant transactions between them. Other actors that may be indirectly involved because of their participation in other IHE Integration Profiles, such as Radiology Scheduled Workflow, Patient Identity Cross-Referencing Integration Profiles, etc., are not shown.

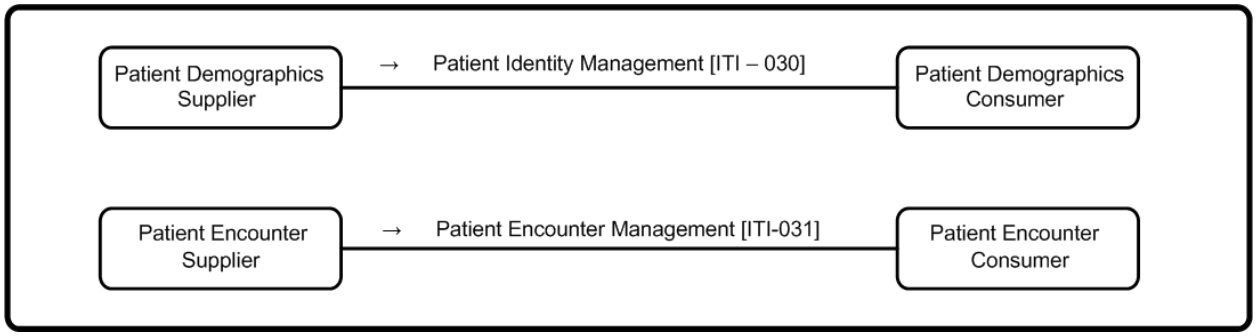

Figure 14.1-1 Patient Administration Management Actor Diagram

Table 14.2-1 lists the transactions for each actor directly involved in the Patient Management Integration Profile. In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled "R"). A complete list of options defined by this Integration Profile that implementations may choose to support is listed in Table 14.2-1.

Table 14.2-1. Patient Administration Management - Actors and Transactions

Actors	Transactions	Optionality	Section in Vol. 2
Patient Demographics Supplier	Patient Identity Management	R	ITI TF-2 : 3.30
Patient Demographics Consumer	Patient Identity Management	R	ITI TF-2 : 3.30
Patient Encounter Supplier	Patient Encounter Management	R	ITI TF-2: 3.31
Patient Encounter Consumer	Patient Encounter Management	R	ITI TF-2: 3.31

14.3 Patient Administration Management Integration Profile Options

Options that may be selected for this Integration Profile are listed in the table 14.3-1 along with the Actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 14.3-1 Patient Administration Management - Actors and Options

Actor	Options	Vol & Section
Patient Demographics Supplier	*Merge (Note 1)*	ITI TF-2: 3.30
	Link / Unlink (Note 1)	ITI TF-2: 3.30
Patient Demographics Consumer	*Merge (Note 1)*	ITI TF-2: 3.30
	Link / Unlink (Note 1)	ITI TF-2: 3.30
Patient Encounter Supplier	*Inpatient / Outpatient Encounter Management*	ITI TF-2: 3.31
	Pending Event Management (Note 2)	ITI TF-2: 3.31
	Advanced Encounter Management	ITI TF-2: 3.31
	Temporary Patient Transfer Tracking	ITI TF-2: 3.31
	Historic Movement	ITI TF-2: 3.31
Patient Encounter Consumer	*Inpatient / Outpatient Encounter Management*	ITI TF-2: 3.31
	Pending Event Management (Note 2)	ITI TF-2: 3.31
	Advanced Encounter Management	ITI TF-2: 3.31
	Temporary Patient Transfer Tracking	ITI TF-2: 3.31
	Historic Movement	ITI TF-2: 3.31

Note 1: An IHE National Extension shall select at least one of the Merge and Link / Unlink Options, and shall mandate the same option for both the Patient Demographics Supplier and the Patient Demographics Consumer implementations in its realm to ensure interoperability.

Note 2: The Pending Event Management Option depends on the Inpatient / Outpatient Encounter Management Option. An implementation supporting the Pending Event Management Option must also support the Inpatient / Outpatient Encounter Management Option.

The PAM profile offers a large number of options to support the exchange of patient demographic and encounter data in a wide variety of environments. Particularly, this profile addresses both acute care settings and ambulatory healthcare organizations. It is unlikely that one particular environment will need all the options.

On one hand, an ambulatory care community might need only the pair of actors Patient Demographics Supplier/Patient Demographics Consumer, using transaction ITI-30. On the other hand, the exchange of patient demographic and encounter data between a hospital patient administration system and its ancillary systems (laboratory, radiology, cardiology, etc.) might be fully satisfied with the pair of actors Patient Encounter Supplier/Patient Encounter Consumer, using transaction ITI-31 with the only option “Inpatient/Outpatient Encounter Management”.

Hence, the first decision that must be made by a healthcare organization for the deployment of this profile is to select the proper actors and the appropriate set of options to cover its needs, ensuring that each selected option will be supported by the actors on both ends of the transactions.

Furthermore, as an IT Infrastructure profile, the PAM profile may not be used standalone. Rather, its actors and transactions will be leveraged by other domain integration profiles (in radiology, cardiology, laboratory, or in cross enterprise document sharing). Here again, the first decision that will be taken by the IHE committee that wishes to leverage PAM for its domain, will be to select the proper set of options and to ascertain the consistent use of these options in its domain.

Thus, during the building process of IHE domain technical frameworks, as well as in the deployment process, the PAM profile will be constrained to reduce its original number of options.

However, to accommodate situations in which a consumer application would not support an option implemented by a supplier application, the PAM profile states that the consumer application shall application-reject a message that it does not support (see ITI TF-2: Appendix C.2.3).

14.3.1 Merge Option

The Merge Option defines the information exchange needed to manage the merging of patient identifiers.

14.3.2 Link / Unlink Option

The Link / Unlink Option defines the information exchanges needed to manage the linking and unlinking of patient identifiers, respectively.

14.3.3 Inpatient / Outpatient Encounter Management Option

The Inpatient / Outpatient Encounter Management Option extends the basic patient encounter management functions by defining the information exchanges needed for pre-admitting a patient and for transferring a patient from one location to another location in the enterprise, as well as for changing patient class.

14.3.4 Pending Event Management Option

The Pending Event Management Option extends the basic patient encounter management functions by defining the information exchanges needed for supporting pending events, e.g., admission, transfer, and discharge.

14.3.5 Advanced Encounter Management Option

The Advanced Encounter Management Option extends the basic patient encounter management functions by defining a set of messages for handling patient temporary absence, changing attending doctor in an encounter, and moving accounts among different patient identities.

14.3.6 Temporary Patient Transfer Tracking Option

The Temporary Patient Transfer Tracking Option defines the information exchange needed for tracking a temporary leave / return of a patient from / to a care facility.

14.3.7 Historic Movement Option

The Historic Movement Option extends the basic patient encounter management functions, as well as the following Options:

- Inpatient / Outpatient Encounter Management
- Pending Event Management
- Advanced Encounter Management Options.

The Historic Movement Option provides a means to uniquely identify any movement event conveyed in the underlying information exchange. This enables updates of such events at any later time point after they were initially reported.

14.4 Patient Administration Management Integration Profile Actor Grouping

14.4.1 Actor Grouping of Patient Encounter Supplier

In order to obtain patient identity and demographics information to serve its patient encounter message functions in transaction ITI-31, a Patient Encounter Supplier is required to be grouped with either a Patient Demographics Supplier or a Patient Demographics Consumer, as shown in Figure 14.4-1.

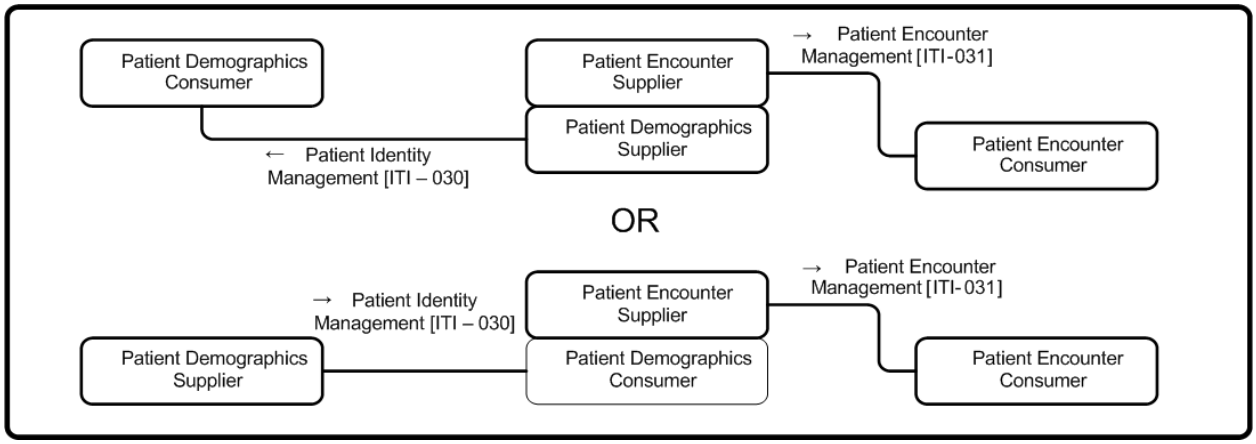

Figure 14.4-1 Patient Encounter Supplier Grouping Requirements

On the other hand, transaction ITI-31 is self-contained in a sense that the Patient Encounter Supplier sends both patient encounter information and patient identity and demographics information (in the context of the encounter data) to the Patient Encounter Consumer. In addition, transaction ITI-31 also allows the Patient Encounter Supplier to send messages to the Patient Encounter Consumer for patient identity maintenance in the encounter context, including patient update and identity merge. There is no required grouping for the Patient Encounter Consumer.

14.4.2 Actor Grouping with other IHE Actors

The PAM profile provides an infrastructure in a healthcare enterprise or across a number of enterprises to distribute the patient identity, demographics, and encounter information, in order to enable various clinical functions in clinical settings. The PAM actors can be grouped with actors in other IHE Integration Profiles.

One possible grouping is between the Patient Demographics Supplier actor in the PDQ profile and either the Patient Demographics Supplier actor or the Patient Demographics Consumer actor in this profile, to add query support defined in the Patient Demographics Query transaction to the same set of patient information managed in the PAM profile.

Furthermore, the Patient Demographics Supplier actor in the PDQ profile can be grouped with the Patient Encounter Supplier actor of this profile. Due to the required grouping of the Patient Encounter Supplier actor (see section 14.4.1), such a grouping can provide query support defined in both the Patient Demographics Query and Patient Demographics and Visit Query transactions to the same set of patient and encounter information that is managed in the PAM profile.

These are some examples of possible grouping of the PAM actors with other IHE actors. Many other possibilities may be useful (either to provide additional values or to allow profile structure simplification). For example, in the radiology scheduled workflow (SWF) profile, the Order Placer and Order Filler actors can be grouped with the Patient Encounter Consumer actor.

14.5 Patient Administration Management Process Flow

14.5.1 Patient Identity Management

The Patient Identity Management incorporates the following process flows. This refines the use case shown in section 14.1.1.

14.5.1.1 Patient Identity Creation and Maintenance

- ***Create Patient***. The Patient Demographics Supplier decides to create a new patient John Smith, based on patient information input from Hospital Sun. At this time, however, there is a limited set of personal information traits of John Smith available. His date of birth, home address, and home phone number, *e.g.*, are unknown. The Patient Demographics Supplier creates the patient identity and sends a Patient Creation message to the Patient Demographics Consumer with the set of known personal information traits.
- ***Update Patient Demographics***. The next day, detailed personal information about John Smith becomes available. The Patient Demographics Supplier updates its patient identity record, and sends out a Patient Update message, including date of birth, home address and home phone number.
- ***Create Temporary Patient***. After a week, the Patient Demographics Supplier creates a temporary patient identity John Doe based on input from Imaging Center Moon.
- ***Update Patient Demographics and Change Patient Identifiers*** After reconciliation of the temporary patient, the Patient Demographics Supplier updates John Doe's demographics to (a new instance of) John Smith, and changes the temporary Patient Identifier originally assigned to a permanent identifier
- ***Merge Patient Identifiers***. After human inspection, it turns out that the two patients named John Smith in the Patient Demographics Supplier actually represent the same real-world patient. The operator decides to merge the two patient identities. The Patient Demographics Supplier sends a Patient Merge message to the Patient Demographics Consumer.

The following diagram shows the process flow:

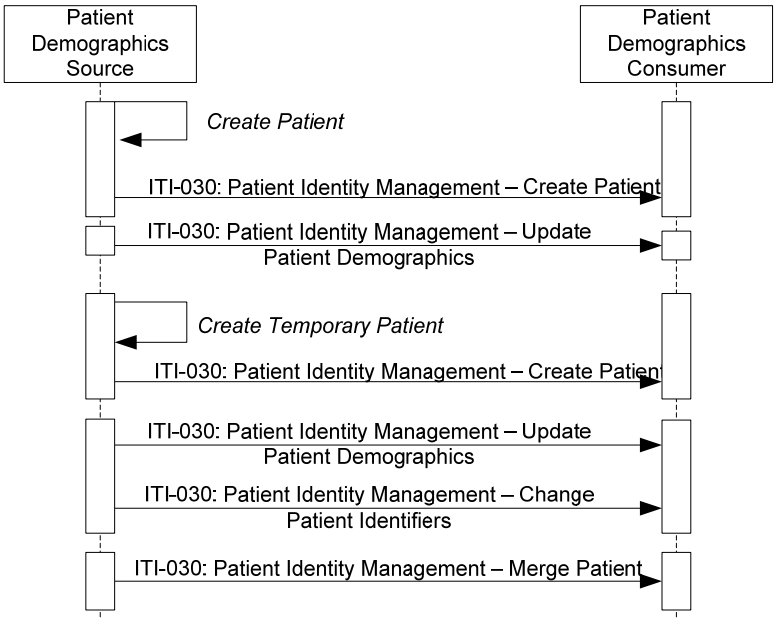

Figure 14.5-1 Patient Identity Management Process Flow in PAM Profile

14.5.1.2 Alternative Process Flow

- ***Link Patient Identifiers***. A similar situation as that mentioned above, except that the local procedures request the Patient Demographics Supplier to link these two duplicated patient records instead of merging them. The operator performs the link function. The Patient Demographics Supplier sends a Patient Identifiers Link message to the Patient Demographics Consumer.

The following diagram shows the alternate portion of the process flow:

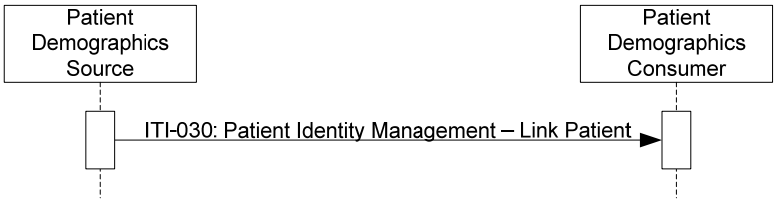

Figure 14.5-2 Patient Identity Management Alternate Process Flow in PAM Profile

14.5.2 Patient Encounter Management

The Patient Encounter Management incorporates the following process flows:

14.5.2.1 Inpatient/Outpatient Encounter and Pending Event Management

In this section, inpatient/outpatient encounter management process flow is described in an environment that involves a number of instances of Patient Encounter Supplier and Patient Encounter Consumer. This refines the use case shown in section 14.1.2

In some institutions, there may be one central Patient Encounter Supplier, while others may have multiple Patient Encounter Suppliers serving patient encounter management functions in different clinical settings (*e.g.*, hospital inpatient, hospital outpatient, clinics). It is the responsibility of a healthcare institution to define the actor roles of its systems, as well as to configure the relationship of a Patient Encounter Supplier and its Patient Encounter Consumers, to satisfy their business process models.

As shown in Figure 14.5-3, in the healthcare institution of this process flow, there are three Patient Encounter Suppliers, each of which serves a number of Patient Encounter Consumers in a specific clinical setting of the institution.

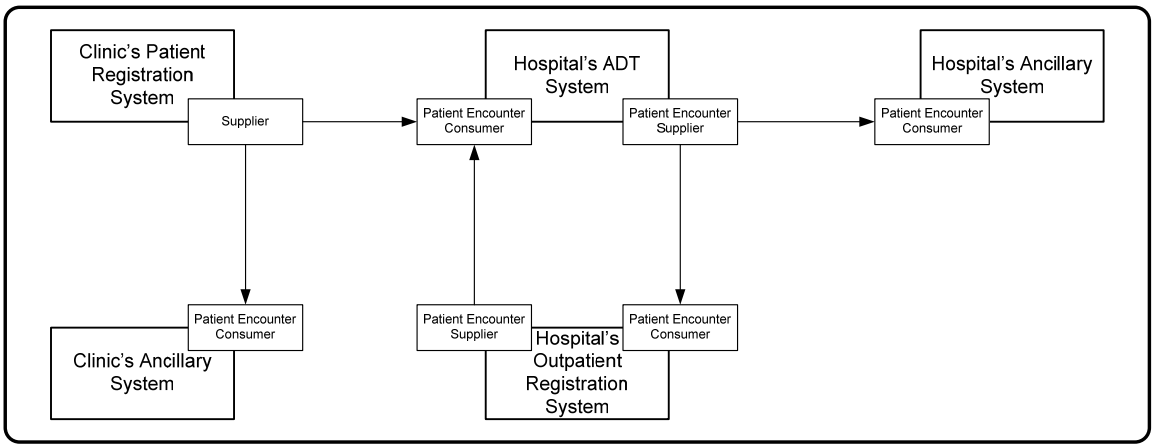

Figure 14.5-3 System and PAM Actor Role Configuration

The systems involved in this process flow implement the following PAM roles:

- Clinic Registration System as Patient Encounter Supplier
- Clinic Ancillary System as Patient Encounter Consumer
- Hospital ADT system as both Patient Encounter Supplier and Patient Encounter Consumer
- Hospital Ancillary system as Patient Encounter Consumer
- Hospital Outpatient Registration System as both Patient Encounter Supplier and Patient Encounter Consumer

Note that the Hospital ADT and Outpatient Registration Systems play both the roles of Patient Encounter Supplier and Patient Encounter Consumer, and cooperate as peers. The relationship between the Patient Encounter Supplier and Patient Encounter Consumer in the same system is dependent on the clinical application logic implemented in the institution, and the definition of this relationship is beyond the scope of the PAM Integration Profile.

The process flow in Figure 14.5-4 is described in the following:

- ***Patient Registration***: A patient arrives for an annual exam at a clinic. The patient record has been created previously by a Patient Demographics Supplier, and exists in the clinic's registration system through its grouping with the Patient Demographics Supplier actor. The clinic's registration system sends the Patient Registration message to the local ancillary systems, and the affiliated hospital's ADT system.
- ***Change Outpatient to Inpatient***: The exam reveals a serious condition of the patient, and an immediate hospital admission is recommended. The patient is referred to the affiliated

hospital for admission. A Change Outpatient to Inpatient message is sent to the hospital's ADT System.

- ***Pre-admit Patient for Hospitalization***: The patient is pre-admitted in the hospital for relevant diagnostic tests. The hospital ADT system sends Patient Pre-Admit message to the Hospital Ancillary System.
- ***Patient Admitted Notification***: The tests confirm the condition, and the patient is admitted to the hospital's ICU. The hospital ADT system sends an Admission Notification message to the Ancillary System.
- ***Patient Insurance Information Update***: During the stay in the ICU, the patient's insurance is verified, and the updated information is sent from the hospital ADT to the Hospital Ancillary System.
- ***Patient Location Transfer***: After a day in the ICU, the patient's condition has improved, and the patient is transferred to a regular bed. The hospital ADT system sends a Patient Transfer message to the Hospital Ancillary System.
- ***Patient Location Transfer Error Reconciliation***: The nurse recording the transfer makes a mistake, and enters the wrong room and bed. After discovering the error, the hospital ADT system sends a Cancel Patient Transfer message to the Hospital Ancillary System, followed by a new Patient Transfer message.
- ***Patient Pending Discharge***: The patient is now recovered and about to leave the hospital. The ADT system sends a Patient Pending Discharge message to the Hospital Ancillary System.
- ***Change Inpatient to Outpatient***: According to the hospital's procedures, the patient is transferred to an outpatient unit for administration of follow-up tests. The ADT system sends a Change Inpatient to Outpatient message to the Hospital Outpatient Registration System.
- ***Register Patient as Outpatient***: The patient is registered in the Hospital Outpatient Registration System, which sends a Patient Registration message to the Hospital ADT system and the Hospital Ancillary System.
- ***Patient Discharged from Outpatient System***: The outpatient encounter is completed. A Patient Discharge message is sent to the Hospital ADT System and to the Hospital Ancillary System.
- ***Patient discharged from Hospital ADT System***: Based on satisfactory test results, the patient is discharged. The hospital ADT system sends a Patient Discharge message to the Hospital Ancillary System.

The following diagram shows the process flows of the discussed use cases:

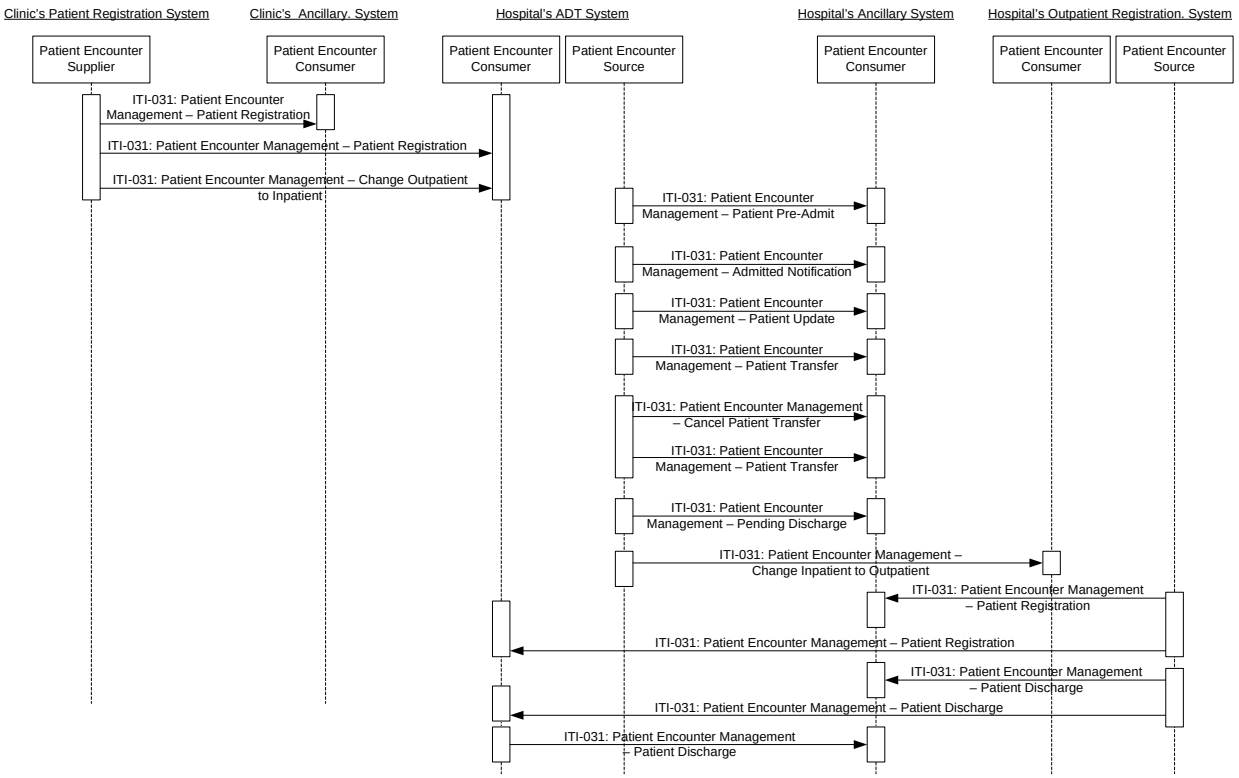

Figure 14.5-4 Inpatient / Outpatient Encounter Management Process Flow in PAM Profile

14.5.2.2 Advanced Encounter Management

- ***Attending Physician Change***: A patient's attending physician changes during an inpatient stay. The Patient Encounter Supplier sends a notification message that contains the name of the new attending doctor to the Patient Encounter Consumer.
- ***Cancellation of Attending Physician Change***: A notification of change of a patient's attending physician was sent in error. The Patient Encounter Supplier sends a cancellation message that contains the name of the old attending doctor to the Patient Encounter Consumer.
- ***Leave of Absence***: An inpatient is authorized a weekend leave of absence from the medical center. The Patient Encounter Supplier sends a notification message to the Patient Encounter Consumer that contains the date and time of the leave of absence and of the expected return.
- ***Cancellation of Leave of Absence***: A notification that an inpatient was authorized a weekend leave of absence was sent in error. The Patient Encounter Supplier sends a cancellation message to the Patient Encounter Consumer.
- ***Return from Leave of Absence***: An inpatient returns to the medical center from a weekend leave of absence. The Patient Encounter Supplier sends a notification message to the Patient Encounter Consumer that contains the date and time of the expected return and of the actual return.

- ***Cancellation of Return from Leave of Absence***: A notification that an inpatient returned from a weekend leave of absence was sent in error. The Patient Encounter Supplier sends a cancellation message to the Patient Encounter Consumer.
- ***Move Account***: The Patient Encounter Supplier sends a message that incorrectly associates Account 12345 with Patient A; in fact, Account 12345 should be associated with Patient B. To effect a correction, the Patient Encounter Supplier sends a message to the Patient Encounter Consumer that contains the account identifier and the identifiers of the patient records between which the account association is to be moved.

The following diagram shows these discussed use cases:

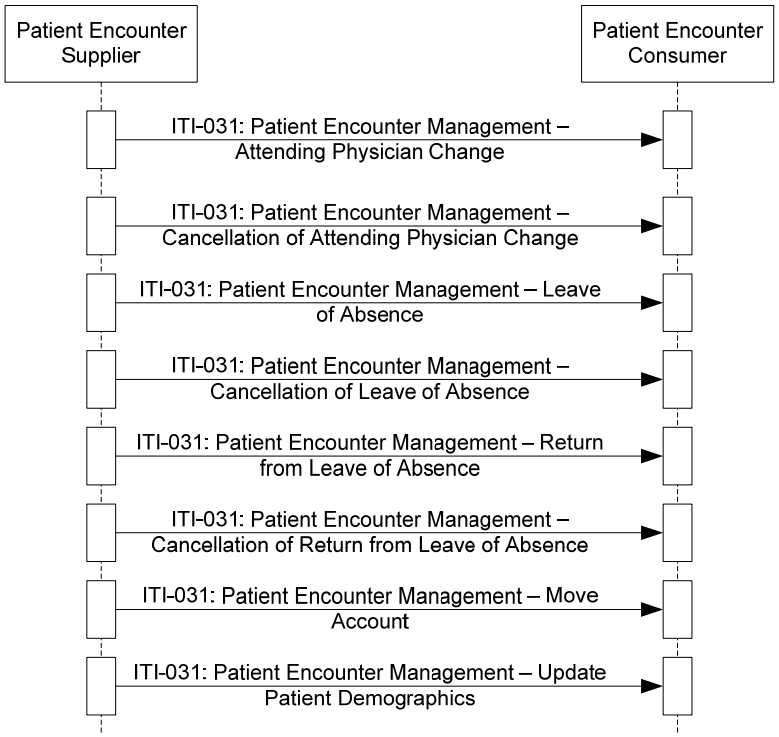

Figure 14.5-5 Advanced Encounter Management Process Flow in PAM Profile

14.5.2.3 Historic Movement Management

Historic tracking of patient admissions, discharges, and other movements may be needed in some healthcare institutions. Such historic events may need to be tracked even beyond the boundary of an episode of care. In order to facilitate this tracking, the Patient Encounter Supplier may send the messages in 14.5.2.1 and 14.5.2.2 to the Patient Encounter Consumer, with the addition of an identifier for the particular encounter with which the patient admission, discharge, or movement is associated.

- ***Patient Location Transfer***: A patient is transferred to bed 23 of Room B after a few days of stay in ICU. The hospital ADT system sends a Patient Transfer message (including the elements provided in the Historic Movement Management Option) to the downstream applications.
- ***Update Previous Transfer Event***. After two days, the operator of the ADT system detects that the transfer destination and time in the previously sent Patient Transfer message were

wrong. He corrects the errors and an Update Historic Patient Transfer message is sent out, to communicate the true room / bed information and the true transfer time.

The following diagram shows these use cases:

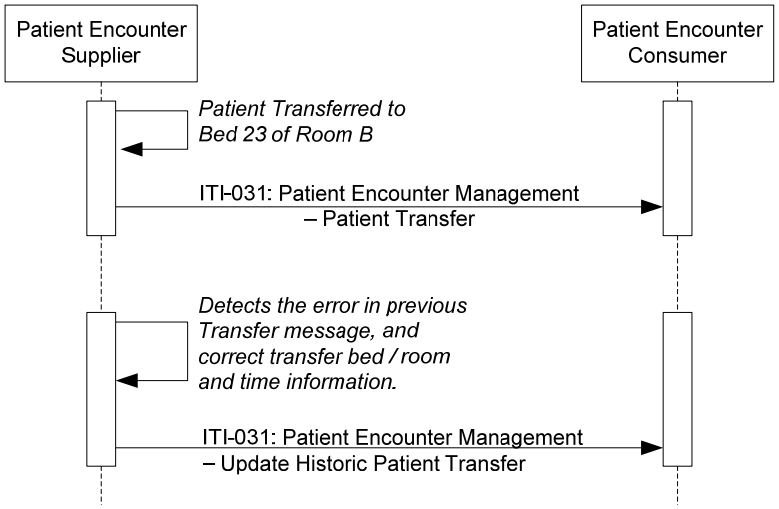

Figure 14.5-6 Historic Movement Management Process Flow in PAM Profile

14.5.2.4 Temporary Patient Transfer Tracking

- ***Departure to Temporary Location***: A chest X-ray is scheduled for an inpatient. To perform this service, the patient needs to be moved from her inpatient bed in the medical service to the Radiology department. When the patient departs from her inpatient bed, the Patient Encounter Supplier sends a notification message to the Patient Encounter Consumer that contains the temporary location to which the patient is being moved.
- ***Arrival at Temporary Location***: When the patient arrives at the Radiology department, the Patient Encounter Supplier sends a notification message to the Patient Encounter Consumer that contains the temporary location to which the patient has been moved.
- ***Cancellation of Departure to Temporary Location***: It is incorrectly communicated that a patient left her inpatient bed to move to the Cardiology department for treatment. The Patient Encounter Supplier sends a cancellation message to the Patient Encounter Consumer that contains the patient's location(s) (permanent and / or temporary) prior to the time of the erroneously communicated departure.
- ***Cancellation of Arrival at Temporary Location***: It is incorrectly communicated that a patient, having left her inpatient bed, arrived in the Surgery department for treatment. The Patient Encounter Supplier sends a cancellation message to the Patient Encounter Consumer that contains the patient's location(s) (permanent and / or temporary) prior to the time of the erroneously communicated arrival.

The following diagram shows these discussed use cases:

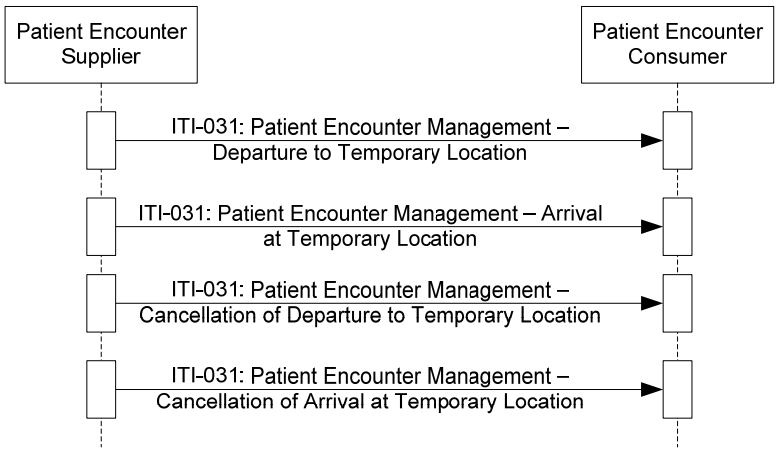

Figure 14.5-7 Temporary Patient Transfer Tracking Process Flow in PAM Profile

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16 Cross-Enterprise Document Media Interchange (XDM) Integration Profile

Cross-Enterprise Document Media Interchange (XDM) provides document interchange using a common file and directory structure over several standard media types. This permits the patient to use physical media to carry medical documents. This also permits the use of person-to-person email to convey medical documents. XDM supports the transfer of data about multiple patients within one data exchange.

Physician to patient to physician - Bob has an MRI and cancer is diagnosed. He is given a CD-R with his MRI results and referral information on it to give to the specialist of his choice.

Patient visiting ED - In addition, Bob, the informed patient, maintains a copy of his EHR record at home and can bring the CD-R with him when he visits the ED for an unrelated emergency.

Physician to physician - Dr. Primary refers his aging patient Mr. Robinson to his first appointment with a gastroenterology specialist. He transfers relevant documents in a zip file attached to an email to the specialist.

The common thread of these use cases is that they are person-to-person communications. The XDM solution is intended to be easy to implement with pre-existing email clients, CD burners and USB ports. XDM does not include any additional reliability enhancements. XDM requires that the recipient be able to support human intervention in order to manually control the importing of the data (patient ID reconciliation, selection of patient of interest from possibly multiple patients' documents on the media).

XDM is document format agnostic, supporting the same document content as XDS and XDR. Document content is described in XDS Document Content Profiles. Examples are XDS-MS, XPHR, XDS-SD, and XD*-LAB.

XDM defines no new metadata. It leverages XDS metadata with emphasis on patient identification, document identification, description, and relationships.

A directory and file structure is documented for populating the media. This structure maintains separate areas for each patient listed and is supported on all referenced media types. Media and the structure were selected based on experience with media interoperability in Radiology, i.e. PDI profile. The media selected are the widespread CD-R, USB removable media, and email with ZIP attachment.

16.1 Actors/ Transactions

Figure 16.1-1 shows the actors directly involved in the XDM Integration Profile and the relevant transactions between them. Other actors that may be indirectly involved due to their participation in XDS, PIX or PDI are not shown.

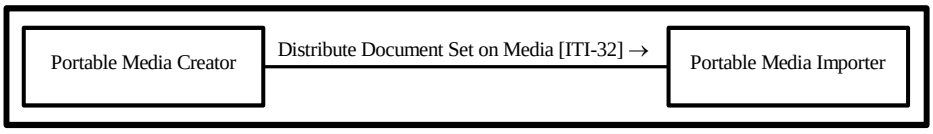

Figure 16.1-1 XDM Actor Diagram

Table 16.1-1 lists the transactions for each actor directly involved in the XDM Profile. In order to claim support of this Integration Profile with one or more actors, an implementation must perform the required transactions (labeled "R"). Transactions labeled "O" are optional. A complete list of options defined by this Integration Profile and that implementations may choose to support is listed in Section 16.2.

Table 16.1-1 XDM Integration Profile - Actors and Transactions

Actors	Transactions	Optionality	Section in Vol. 2
Portable Media Creator	Distribute Document Set on Media	R	ITI TF-2:3.32
Portable Media Importer	Distribute Document Set on Media	R	ITI TF-2:3.32

16.2 XDM Integration Profile Options

Options that may be selected for this Integration Profile are listed in the table 16.2-1 along with the Actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 16.2-1 XDM - Actors and Options

Actor	Options	Vol & Section
Portable Media Creator	*USB (Note 1)*	ITI TF-1 16.2.1
	CD-R (Note 1)	ITI TF-1 16.2.2
	ZIP over Email (Note 1)	ITI TF-1 16.2.3
	Basic Patient Privacy Enforcement	ITI TF-2:3.32.4.1.4.1
	Zip over Email Response (Note2)	ITI TF-1 16.2.4
Portable Media Importer	*USB (Note 1)*	ITI TF-1 16.2.1
	CD-R (Note 1)	ITI TF-1 16.2.2
	ZIP over Email (Note 1)	ITI TF-1 16.2.3
	Basic Patient Privacy Enforcement	ITI TF-2:3.32.4.1.4.1
	Zip over Email Response (Note2)	ITI TF-1 16.2.4

Note 1: At least one of these options is required for each Actor. In order to enable a better interoperability, is highly recommended that the actors support all the options.

Note 2: This option requires the ZIP over Email Option.

16.2.1 USB Option

In this option the Portable Media Creator writes a set of documents on USB media. The media is physically transported to the Portable Media Importer which then imports the document set.

16.2.2 CD-R Option

In this option the Portable Media Creator writes a set of documents on CD-R media. The media is physically transported to the Portable Media Importer which then imports the document set.

16.2.3 ZIP over Email

In this option the Portable Media Creator creates an ordinary ZIP file of the virtual media containing document set(s). The ZIP file is attached to an Email sent to the Portable Media Importer which then retrieves the Email and imports the ZIP file containing the document set.

16.2.4 ZIP over Email Response

In this option the Portable Media Importer sends a response (MDN Based) to the Portable Media Importer to acknowledge that the Import operation of the Document Set(s) received was successful. If this option is supported, the ZIP over Email option shall be supported.

16.3 XDM Process Flow

XDM describes the exchange of a set of a patient's documents between healthcare providers, such as: physicians, hospitals, special care networks, or other healthcare professionals.

Where XDS is not desirable or available for one of the participants in the exchange of information, XDM is a viable option.

XDM should be used in a situation where the information receiver is an individual who will manually interpret or examine the data and associated documents as though they were using physical media. XDM also allows for the exchange of documents relating to multiple patients, since the data will be interpreted manually by human intervention.

The XDM integration profile is intended only for exchange of personal medical documents and not intended to address all cross-enterprise EHR communication needs. Some use cases may require the use of other IHE integration profiles such as XDS, DSG, PIX, and ATNA. Other use cases may only be partially supported, while still others may require future IHE integration profiles.

Use Cases:

1. Dr. Primary refers his aging patient Mr. Robinson to his first appointment with a gastroenterology specialist.

   In a case where either Dr. Primary's office or Dr. Gastro's clinic was not able to handle secure email, or other sustained online point-to-point communications (eg: http over VPN), the XDM profile would provide further solutions for the simpler environment, such as the use of physical media, or email where the interchanged document set will be manually interpreted by a human intervention.

2. In a hospital that does not have an XDS infrastructure; the XDS-MS content profile discharge use case can also be handled by XDM. For example:

In a hospital, or in the case of a family physician not using robust EHR, the patient could be handed a CD or USB media with their discharge information on it to bring with them to their follow-up visit with their family physician.

3. Mabel is transferred from a hospital setting to her retirement home for long-term care.

   If the hospital does not have an EHR application that automatically interprets her medical data and shares it with the necessary members of her health team, the information can be transferred manually directly to the file clerk, intake coordinator, records manager, or primary physician depending on the organization's resource model.

4. Stanley's recent MRI has generated unusual results that Stanley's primary physician would like to consult with another specialist in a specialized cancer facility located across the state. Since there is not likely to be an affinity domain between the remote health environments, XDM can be used instead.

5. Bob, the informed patient, maintains a copy of his Personal Health Record (PHR) at home. In this situation, Bob can be given a copy of his medical information on physical media such as a CD-ROM to take home with him. Bob now has an advantage that he can continue to have his complete medical record available with him on sudden emergency department visits, even when he is on an out-of-state trip where the new ED would have no access to the repository of his home affinity domain.

This profile is only defining the digital transport mechanism used for such use cases. Content transported will be detailed by Content Profiles such as the ones defined by the IHE PCC (Patient Care Coordination) domain.

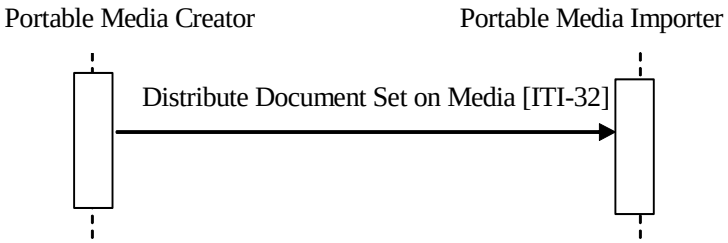

Figure 16.3-1 Process Flow in XDM Profile

16.4 Digital communication

16.4.1 Actual Media Type

The media can be either CD-R or a USB media device, because these are the most common media types in other industries for the portable transport of electronic information. This supplement requires using one of these media types, depending on the use case. The benefit and risks of the reusability of the media deployed should be taken into account, especially when the media is under the control of the patient.

Note: 1. Because the size of documents to be exchanged rarely requires more than the capacity of a CD, and the format for storing data on various different recordable DVD media is not totally stable yet, this profile is following the restriction defined in the IHE RAD PDI Profile, to not use recordable DVD media at this time.

2. CD-RW is excluded from this profile because field experiences with CD-RW in radiology with this media showed significant interoperability problems and significant accidental damage levels.

3. The CD-R media is limited to the 74 minute blanks because the long playing CD-R format gains the larger capacity by eliminating one level of error correction and detection. The resulting much higher undetected error rate is considered unacceptable for medical data.

16.4.2 Virtual Media over a Network

The media can be a ZIP file containing the document set and sent via a secure email message.

16.4.3 Media Content

The requirements for media content are intended to promote the simple transfer of medical documents, including patient summaries, lab results, discharge letters and reports, and to allow for the viewing of such documents on general purpose computers by care providers or patients.

Created media are required to contain documents and the relevant associated metadata.

The media contains one or more Submission Sets including the documents and the associated metadata, organized in a well-defined directory structure starting at the root level.

The media content can be made web viewable by a web browser by providing optional files containing HTML content. This content must be based on the original documents in order to ensure consistency. Any ordinary web browser can be used to read these files. The Portable Media Importer ignores these files. They are just intended for the human recipient.

Additional content may be present (files, directories), but can be ignored by the Portable Media Importer.

To summarize, the Portable Media Importer has two complementary ways to access the media and its content through a basic web browser:

- By inspecting in the directory dedicated to XDM all the subdirectories that contain a specifically named metadata file compatible with XDM
- By presenting to the user the HTML index file that lists the submission sets and documents contained in the media.

Access to the content of an individual document is outside the scope of this Integration Profile and shall be addressed in specific IHE document content Integration Profiles.

16.5 Security considerations

The Profile assumes that the Healthcare delivery organizations that are using Portable Media Creator and Importer have an agreement defining when they can interchange PHI. This may require an explicit patient consent (depending on existing regulations) and an agreement on how to manage the potential inconsistency between the security policies. The main aspects that should be covered by this agreement are similar to XDS – See Appendix L. In addition, the following aspects should be covered:

- Management of Patient identification in order to perform patient reconciliation correctly upon importation of the documents.
- Measures taken to avoid or limit loss of media or email, and detect that which occurs.

In the case of physical media, security responsibilities for confidentiality and integrity are transferred to the patient by providing the media to the patient. In this case it is the patient's responsibility to protect the media, and the patient has the authority to disclose the contents of the media as they choose. They disclose the contents by providing the media.

The Portable Media Creator in most cases does not know who the ultimate Importer will be, thus rendering encryption impractical.

In the case of transfer over email using a ZIP attachment, the transaction is secured by the use of S/MIME.

Both Actors for this Profile require a grouping with an ATNA Secure Node or Secure Application.

17 Basic Patient Privacy Consents Integration Profile

The document sharing infrastructure provided by XD* allow for the publication and use of clinical documents associated with a patient. This profile allows for an XDS Affinity Domain to have a number of privacy consents. This allows for more flexibility to support some patient concerns, while providing an important and useful dataset to the healthcare provider. Without BPPC, the XDS profile requires that the administrators of an XDS Affinity Domain creates and agrees to a single document publication and use policy (See ITI TF-1: Appendix L). Such a single XDS Affinity Domain Policy is enforced in a distributed way through the inherent access controls of the systems involved in the XDS Affinity Domain.

This profile will use terms consistent with ISO 22600 - Privilege Management and Access Control (PMAC), but is not restricted to systems that implement PMAC. The systems involved in XDS are expected to support sufficient Access Controls to carry out the Policy of the XDS Affinity Domain3.

Healthcare providers utilize many different sets of data to carry out treatment, billing, and normal operations. This information may include patient demographics, contacts, insurance information, dietary requirements, general clinical information and sensitive clinical information. With BPPC, this information may be published to XDS as independent documents under different privacy consent policies.

Healthcare providers in different functional roles will have different needs to access these documents. For example, administrators may need to be able to access the patient demographics, billing and contact documents. Dietary staff will need access to the dietary documents but would not need access to insurance documents. General care providers will want access to most clinical documents, and direct care providers should have access to all clinical documents.

This profile provides a mechanism by which an XDS Affinity Domain can create a basic vocabulary of codes that identify XDS Affinity Domain privacy consent policies with respect to document sharing. Each privacy consent policy should identify in legal text what the acceptable use, re-disclosure uses, which functional roles may access which document and under which conditions, etc. The administration of the XDS Affinity Domain will assign each privacy consent policy a unique identifier (or code) for use within the XDS Affinity Domain. Future profiles may include in addition to the legal text, a structured and coded expression of the consent policy that can be used to support even more dynamic understanding of the patient's directives (see HL7 and OASIS).

17.1 Basic Patient Privacy Consent Use-Cases

This section gives examples of some possible patient privacy consent policies and how the systems publishing documents and using documents might act. This is an informative section and should not be interpreted as the only way to implement the BPPC profile. Its purpose is to allow implementers of BPPC to more easily understand the principle of operation of BPPC.

[3] See the IHE white paper "HIE Security and Privacy through IHE" published on the IHE web site http://www.ihe.net/Technical_Framework/upload/IHE_ITI_Whitepaper_Security_and_Privacy_2007_07_18.pdf

17.1.1 Implied Consent vs Explicit Consent

This profile supports both Implied Consent as well as Explicit Consent environments. In order to provide a profile with global appeal we have supported both environments. In an implied consent environment it would be normal for a Document Consumer to find no instance of a patient specific acknowledgement of a privacy consent policy in the XDS Affinity Domain, as capturing the act of acknowledging a privacy consent policy would not be required. Note: this may also be true in an Explicit Consent environment, where obtaining the acknowledgement is delayed due to medical reasons (e.g. emergency).

An XDS Affinity Domain might have a paper document that describes their Privacy Consent Policy. In our example this Privacy Consent Policy will be given a local XDS Affinity Domain policy unique identifier (e.g. an OID such as: 9.8.7.6.5.4.3.2.1). The example in Figure 19.1-1 is ridiculous (i.e., chicken costume) but is provided to emphasize that IHE doesn't write these policies, and to make clear that the BPPC profile could be used to enforce any policy that could be written in human readable form, provided that all actors can be configured to enforce that policy. This example also points out that the content of the policy is human readable text, and that we provide no structured or coded way to interpret. This example policy might looks like:

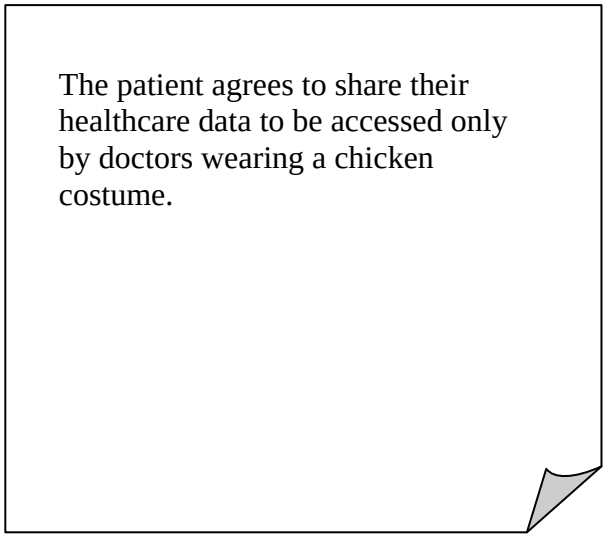

Figure 17.1-1 Policy Example

17.1.1.1 Opt-In

A common structure for sharing clinical documents requires that the patient first acknowledge that they want this sharing to happen before any documents are actually shared. In this case the XDS Affinity Domain administrators would write a policy that indicates what should be shared, when it should be shared, when it can be used, etc. There would also be an overriding XDS Affinity Domain policy that indicates that no document will be shared until the patient has explicitly chosen to participate.

17.1.1.2 Opt-Out

Equally as common is a structure for sharing documents that presumes that when the patient chooses to get care within a care setting, that they are implicitly agreeing to the normal sharing of their documents for treatment purposes. In this environment, there is usually a control that allows a patient to choose to NOT participate in this sharing. This is commonly referred to as “opt-out”.

In this case the existence an acknowledgement to an opt-out policy would mean that documents should no longer be shared, and any documents that might appear should not be used. Clearly the XDS Affinity Domain administrators need to make the actual behavior clear in their policies.

17.1.2 Wet Signature

An XDS Affinity Domain might have the patient acknowledge the consent through ink on paper. For Example:

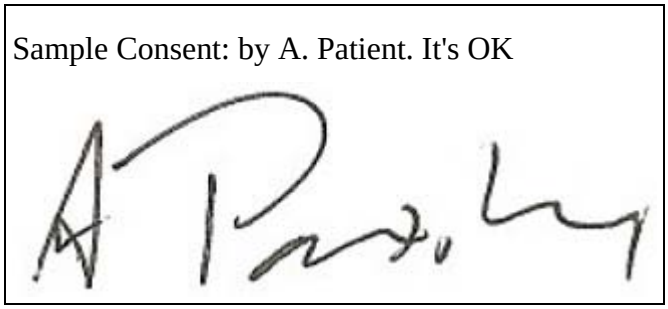

Figure 17-2 Simplistic Consent Example

This acknowledgement is captured according to the XDS Scanned Document Content Profile (XDS-SD), with the additional parameters specified in the BPPC Content Profile also applied. This is submitted into the XDS Affinity Domain as proof that the patient has acknowledged policy 9.8.7.6.5.4.3.2.1.

The following shows this graphically:

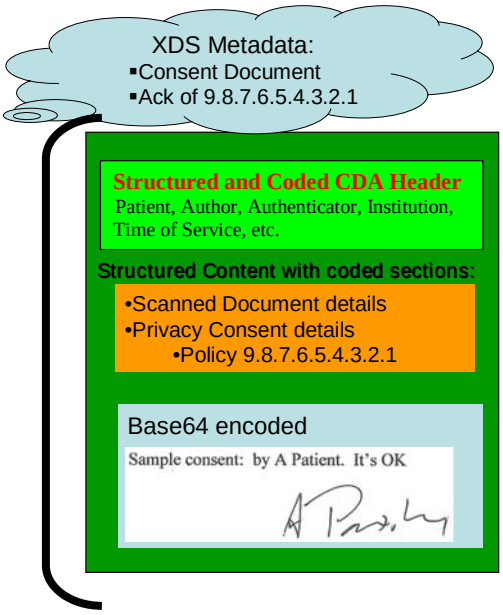

Figure 17.1.2-1 Graphical representation of consent with wet signature

If an XDS Affinity Domain wants to further provide non-repudiation protections it may choose to apply a digital signature using the IHE-DSG Content Profile to the whole package with the appropriate purpose and signed by an appropriate signing system/person.

17.1.3 Advanced Patient Privacy Consents

An XDS Affinity Domain may have jurisdictional or organizational policies that require support for more complex patient privacy consent policies. These privacy policies may require that a patient explicitly consent to disclosure of protected or sensitive health information to specific entities. The BPPC profile provides a starting point for implementing these types of privacy consent policies, but does not explicitly specify how additional information needed to enforce the policy would be conveyed. In these cases, the capability of BPPC may not be enough to support all types of needs. An example of an Advanced Patient Privacy Consent would be when a patient wants to name individuals that can access their documents.

17.2 Creating Privacy Consent Policies

The administrators of the XDS Affinity Domain will need to develop and publish an overall XDS Affinity Domain Policy that clearly defines the overall appropriate use of the XDS Affinity Domain. This is the subject of Appendix L and is not further defined here.

Within this XDS Affinity Domain Policy is a defined set of acceptable use Patient Privacy Consent Policies. A Patient Privacy Consent Policy further explains appropriate use of the XDS Affinity Domain in a way that provides choices to the patient. The BPPC profile places no requirements on the content of these policies nor the method used to develop these policies (See Appendix P for some guidance on developing these policies). BPPC only assumes that the overall XDS Affinity

Domain Privacy Policy can be structured as a set of specific policies (A, B, C, D in the example below), where each one may be used independently or combined in relationship to publication and access of a specific type(s) of document.

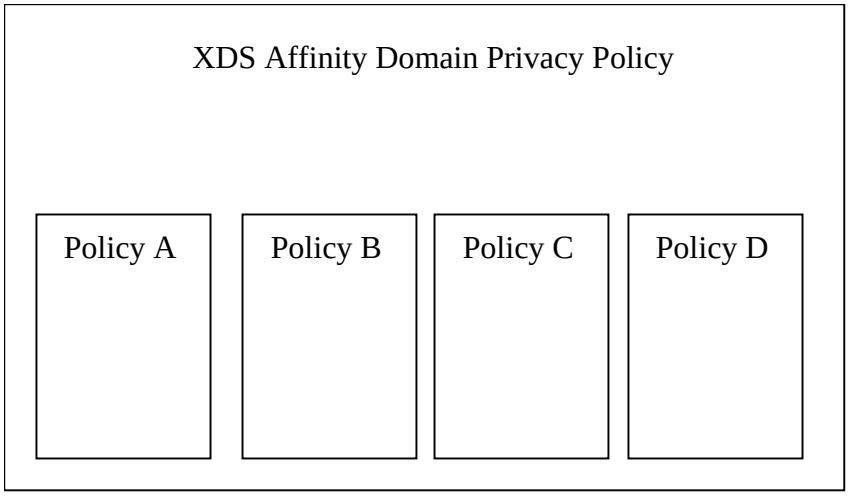

Figure 17.2-1: Privacy Policy Hierarchy

A Privacy Consent Policy will identify who has access to information, and what information is governed by the policy (e.g., under what conditions will a document be marked as containing that type of information). The mechanism for publishing these policies is not described by this profile. The set of Privacy Consent Policies written by the XDS Affinity Domain must be able to be implemented by the technologies in all of the systems that have access to the XDS Affinity Domain. This means that the Privacy Consent Policies must be created with great care to ensure they are enforceable.

Each Patient Privacy Consent Policy will be given a unique identifier (OID) known as a Patient Privacy Consent Identifier. It is this identifier that is used to label documents published into the XDS Affinity Domain. This label provides the control linkage back to the appropriate Patient Privacy Consent Policy. This label is additionally used when capturing a patient's acknowledgement of a specific Patient Privacy Consent Policy.

An XDS Affinity Domain may have legacy documents that were published prior to all systems supporting the BPPC Profile, and thus will have confidentiality codes not defined under the BPPC Profile (e.g. For example, the HL7 confidentialityCode for "N" [normal]). The XDS Affinity Domains will need to provide Privacy Consent Policies for granting access to documents that use these non-BPPC confidentiality code values.

XDS Affinity domains should also determine their strategy for addressing the changing over time of Privacy Consent Policies.

Finally, Privacy Consent Policies used within an XDS Affinity Domain will very likely be different than those used with the XDM or XDR Profiles as these profiles often are used to transfer documents in ad-hoc ways. The patient may provide a consent given to share information on media to the provider creating the media for specific use, rather than for more general sharing within an XDS Affinity Domain. When transferring information that originated in an XDS Affinity Domain to media (XDM), the Privacy Consent Policies found in the XDS Affinity Domain might be changed during the publication process. There are also differences in the sensitivity that should be

considered for consents shared on media or transmitted through XDR and those shared in an XDS Affinity Domain. See the section Security Considerations later in this volume for more details.

Consumers of documents that implement this profile are required to enforce access control based on the policies described by the XDS Affinity Domain. This is because the consumers of the documents are best aware of the functional role, how the data will be used, the relationship between provider and patient, the urgency of access, etc. The mechanism that Document Consumers use to associate individual users with functional roles is not within the scope of this profile.

17.2.1 Summary of the creation and publication of the policies

1. The XDS Affinity Domain will write and agree to the Affinity Domain Policy (lots of lawyers involved).
2. The XDS Affinity Domain Policy will include a small set of Privacy Consent Policies (more lawyers). These are text documents very similar to the privacy consent documents used today.
3. Each Privacy Consent Policy will be given an XDS Affinity Domain unique identifier (OID) called the Privacy Consent Policy Identifier
4. The XDS Affinity Domain Policy and all of the Privacy Consent Policies will be published in a way consistent with the XDS Affinity Domain's Policy. It is expected that this will be sufficiently public to support local regulation.

17.3 Actors/Transactions

The BPPC Integration Profiles Actors are represented in the figure below.

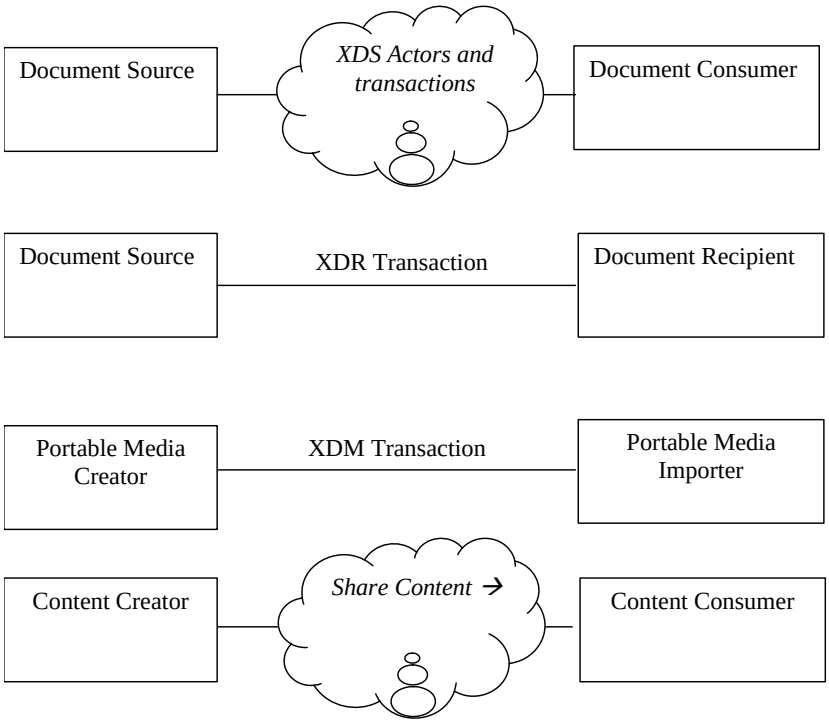

Figure 17.3-1 BPPC Actor Diagram

For details on Content Creator and Content Consumer Actors see PCC TF 2:4.

Table 17.3-1 lists the transactions for each actor directly involved in the BPPC Profile. In order to claim support of this Integration Profile, an implementation must perform the required transactions (labeled "R"). Transactions labeled "O" are optional. A complete list of options defined by this Integration Profile and that implementations may choose to support is listed in Section 19.4.

Table 17.3-1. BPPC Integration Profile - Actors and Transactions

Actors	Transactions	Optionality	Section
Document Source	Provide and Register Document Set	R (note 1)	ITI TF-1:19.4.1
	Provide and Register Document Set-b	R (note 1)	ITI TF-1:19.4.1
Document Consumer	Retrive Document	R (note 1)	ITI TF-1:19.4.1
	Retrieve Document Set	R (note 1)	ITI TF-1:19.4.1
	Registry Stored Query	R (note 1)	ITI TF-1:19.4.1
Document Recipient	Provide and Register Document Set-b	R (note 1)	ITI TF-1:19.4.1
Portable Media Creator	Distribute Document Set on Media	R (note 1)	ITI TF-1:19.4.1
Portable Media Importer	Distribute Document Set on Media	R (note 1)	ITI TF-1:19.4.1
Content Creator	*Share Content*	R (note 2)	ITI TF-1:19.4.3 ITI TF-1:19.4.4
Content Consumer	*Share Content*	R (note 3)	ITI TF-1:19.4.5

Note 1: Actor shall implement the Basic Patient Privacy Enforcement Option.

Note 2: Content Creator shall implement the Basic Patient Privacy Acknowledgement Option, and may choose to implement the Basic Patient Privacy Acknowledgement with Scanned Document Option

Note 3: Content Consumer shall implement the Basic Patient Privacy Acknowledgement View Option.

17.4 Basic Patient Privacy Consent Profile Options

Options that may be selected for this Integration Profile are listed in Table 19.4-1 along with the IHE actors to which they apply.

Table 17.4-1 Basic Patient Privacy Consents - Actors and Options

Actors	Option	Section in Vol. 2
Document Source	*Basic Patient Privacy Enforcement*	ITI TF-2:3.15.4.1.3.1 ITI TF-2:3.41.4.1.3.1
Document Consumer	*Basic Patient Privacy Proof*	ITI TF-2:3.18.4.1.3.6
	Basic Patient Privacy Enforcement	ITI TF-2:3.18.4.1.3.5 ITI TF-2:3.17.4.1.3.1 ITI TF-2:3.43.4.2.3

Document Recipient	*Basic Patient Privacy Enforcement*	ITI TF-2:3.41.5.1
Portable Media Creator	*Basic Patient Privacy Enforcement*	ITI TF-2:3.32.4.1.4.1
Portable Media Importer	*Basic Patient Privacy Enforcement*	ITI TF-2:3.324.1.4.1
Content Creator	*Basic Patient Privacy Acknowledgement*	ITI TF-2:5.1.2
	Basic Patient Privacy Acknowledgement with Scanned Document	ITI TF-2:5.1.3
Content Consumer	*Basic Patient Privacy Acknowledgement View*	ITI TF-2:5.1.2 ITI TF-2:5.1.3

17.4.1 Basic Patient Privacy Enforcement Option

All documents managed in an XDS Affinity Domain, or transferred using XDM/XDR, are labeled with a confidentialityCode. The BPPC Profile provides a way for the administrators of an XDS Affinity Domain to define a vocabulary and meaning to that vocabulary. Actors which support this option in XDS, XDM, and XDR have requirements detailed in those profiles.

17.4.2 Basic Patient Privacy Proof Option

To know if a specific patient has acknowledged a specific Patient Privacy Consent Policy, a Document Consumer Actor can use the Registry Stored Query Transaction to query for Patient Privacy Consent Acknowledgment Documents. This should be done by document class (consent). The XDS Metadata in the query response is enough to determine the Patient Privacy Consent Policies that have been acknowledged (EventCodeList), and the effective timeframes if specified. The details for this are in ITI TF-2:3.18.4.1.3.5.

17.4.3 Basic Patient Privacy Acknowledgement Option

The Content Creator that claims to support the Basic Private Privacy Acknowledgement option shall be able to create Patient Privacy Consent Acknowledgement Document Content as specified in ITI TF-2:5.1.

A Patient Privacy Consent Acknowledgement Document is a kind of medical document. The content of a Patient Privacy Consent Acknowledgement Document shall include the effective time of the consent and XDS Affinity Domain defined coded vocabulary identifying the Patient Privacy Consent Policy Identifier (OID) acknowledged by the patient. The content of the Patient Privacy Consent Acknowledgement Document may include a text description of what the patient has acknowledged.

The Patient Privacy Consent Acknowledgement Document may be signed. There are cases, as seen in the use-cases, where the Content Creator would need to be grouped with a DSG Content Creator. The BPPC profile does not require this grouping. This grouping can be fully specified in an IHE Integration Statement.

17.4.4 Basic Patient Privacy Consent Acknowledgement with Scanned Document Option

A Basic Patient Privacy Consent Acknowledgement Document may include a scanned document. For example of the scanned document could be a wet signature by the patient on the text. The Content Creator that claims to support Basic Patient Privacy Consent Acknowledgement with Scanned Document Option shall be able to create a Patient Privacy Consent Acknowledgement with Scanned Document Content as specified in ITI TF-2:5.1.3.

17.4.5 Patient Privacy Consent Acknowledgement View Option

The Content Consumer that claims to support the Patient Privacy Consent Acknowledgement View Option shall be able to display the Patient Privacy Consent Acknowledgement Document Content as specified in IHE ITI TF-2:5.1.2 and IHE ITI TF-2:5.1.3.

The BPPC Content Consumer shall be grouped with a XDS-SD Content Consumer. This means that a Content Consumer for BPPC Content must also be able to display XDS-SD content. This is required due to the the common practice of capturing Wet Signatures.

17.5 Basic Patient Privacy Documents Bindings to XDS, XDR, XDM

Actors from the ITI XDS, XDM and XDR profiles embody the Content Creator and Content Consumer sharing function of this profile. A Content Creator or Content Consumer may be grouped with appropriate actors from the XDS, XDM or XDR profiles to exchange the content described therein. The metadata sent in the document sharing or interchange messages has specific relationships or dependencies (which we call bindings) to the content of the clinical document described in the content profile.

The Patient Care Coordination Technical Framework (PCC-TF) defines the bindings to use when grouping the Content Creator of this Profile with actors from the IHE ITI XDS, XDM or XDR Integration Profiles.

Scanned Documents Bindings			
Content	**Binding**	**Actor**	**Optionality**
Basic Patient Privacy Documents	Medical Document Binding to XD*	Content Creator	R
		Content Consumer	R

17.6 BPPC Process Flow

This flow shows how an XDS Affinity Domain would use the BPPC Profile. Only a basic flow is shown, the profile supports many alternative flows.

17.6.1 Checking for a patient's acknowledgement of a privacy consent policy

A Document Consumer Actor that supports the BPPC Profile – Basic Patient Privacy Consent Proof Option can query an XDS Affinity Domain for instances of Patient Privacy Consent Acknowledgement Documents that have been acknowledged by a specific patient. Through the XDS Metadata the Document Consumer can determine which Patient Privacy Consent Policies have been acknowledged.

Note if the local regulations allow, some XDS Affinity Domains may not publish the consent documents, so systems should be able to handle the configurations where no Patient Privacy Consent Acknowledgment Document is in the XDS Affinity Domain for a specific patient.

Note if the local regulations allow, some patients may have documents shared before informed consent can be captured. In this case the XDS Affinity Domain policy needs to explain the default behavior, that behavior for the absence of a consent document.

17.6.2 Recording a patient's acknowledgement of a privacy consent policy

The Content Consumer Actor creates Patient Privacy Consent Acknowledgement Documents with or without a scanned document part. This document records the patient's acknowledgement of a specified policy.

17.6.3 Publishing documents against a consent policy

All documents managed in an XDS Affinity Domain, or transferred using XDM/XDR, are labeled with a confidentialityCode. The BPPC Profile provides a way for the administrators of an XDS Affinity Domain to define a vocabulary and meaning to that vocabulary.

The Document Source Actor determines which of the XDS Affinity Domain – Privacy Consent Policies would allow the documents to be published. In some XDS Affinity domains this may require that the system check that a patient has indeed acknowledged to the specific policy.

The Document Source Actor will set the XDS Metadata – confidentialityCode - to the OIDs of the Privacy Consent Policy Identifiers that indicate the appropriate use/constraint (determined by the XDS Affinity Domain Policy)

The XDS Registry validates that each of the confidentialityCode(s) are from the approved list of confidentialityCode for use within the XDS Affinity Domain.

17.6.4 Using published documents

When a Document Consumer queries the XDS Affinity Domain it may utilize the confidentialityCode filter in the Registry Stored Query to restrict the documents returned to those that the Document Consumer can utilize.

The Document Consumer can set the confidentialityCode in the Registry Stored Query Transaction to the list of XDS Affinity Domain Policy Identifiers (OIDs) that would allow for the documents to be used. This way the Document Consumer will receive only information on documents with the specified confidentialityCode(s).

The Document Consumer will enforce access controls based on the returned XDS metadata-confidentialityCode, system type, user, context, and any number of other factors that the system is capable of enforcing.

The Document Consumer may be capable of querying for 'Approved' consent acknowledgement documents and using the resulting XDS Metadata as the list of currently Approved Patient Privacy Consent Acknowledgement Documents. There is no requirement for the Document Consumer system to retrieve the Patient Privacy Consent Acknowledgement Document content.

17.7 Security Considerations

Consents stored in an XDS Affinity Domain are also governed by privacy policies. The content of a Patient Privacy Consent Acknowledgement Document may itself contain sensitive information. For example, a terminally ill patient may decide that his prognosis should not be shared with his family members, but that other information may be. Sharing the Patient Privacy Consent Acknowledgement Document with family members would potentially inform them of a negative prognosis.

However, Patient Privacy Consent Acknowledgement Documents stored in the clear on media (XDM), or transmitted through XDR, should not contain sensitive information. The rationale is that the receiver of the information must be able to read the consent that was used to share this information in order to understand how they must treat the information with respect to their own Privacy Consent Policies.

Implementation of Patient Privacy Consent Policies within a healthcare environment has different considerations and risks than implementing similar access control policies within other non-treatment environments. This is for the simple reason that failing to provide access to critical healthcare information has the risk of causing serious injury or death to a patient. This risk must be balanced against the risk of prosecution or lawsuit due to accidental or malicious disclosure of private information. The XDS Affinity Domain should take care in writing their Privacy Consent Policies to avoid this.

One mitigation strategy that is often adopted in healthcare provides accountability through audit controls. That is to say that the healthcare providers are trusted not to abuse their access to private information, but that this is followed up by a policy of monitoring healthcare provider accesses to private information to ensure that abuse does not occur. This strategy reduces the risk of serious death or injury due to lack of access to critical healthcare information, at the increased risk of disclosure of private information. This is why the ITI Technical Committee created the Audit Trail and Node Authentication (ATNA) Integration profile, and furthermore, why that profile is a requirement of XDS and related profiles.

Another risk that must be resolved by an affinity domain is how to address the issues of sharing truly sensitive information in a registry (e.g., psychology documents). One strategy that might be recommended is that truly sensitive data not be shared within the XDS Affinity Domain; directed communications using XDR or XDM may be more appropriate.

18 Cross-Enterprise Sharing of Scanned Documents Content Integration Profile

A variety of legacy paper, film, electronic and scanner outputted formats are used to store and exchange clinical documents. These formats are not designed for healthcare documentation, and furthermore, do not have a uniform mechanism to store healthcare metadata associated with the documents, including patient identifiers, demographics, encounter, order or service information. The association of structured, healthcare metadata with this kind of document is important to maintain the integrity of the patient health record as managed by the source system. It is necessary to provide a mechanism that allows such source metadata to be stored with the document.

This profile defines how to couple such information, represented within a structured HL7 CDA R2 header, with a PDF or plaintext formatted document containing clinical information. Furthermore, this profile defines elements of the CDA R2 header necessary to minimally annotate these documents. Such header elements include information regarding patient identity, patient demographics, scanner operator identity, scanning technology, scan time as well as best available authoring information. Portions of CDA R2 header, along with supplemental document registration information, are then used to populate XDS Document Entry metadata.

The content of this profile is intended for use in XDS, XDR and XDM. Content is created by a Content Creator and is to be consumed by a Content Consumer. The Content Creator can be embodied by a Document Source Actor or a Portable Media Creator, and the Content Consumer by a Document Consumer, a Document Recipient or a Portable Media Importer. Obligations imposed on the Content Creator and the Content Consumer by this profile are understood to be fulfilled by the software that creates the final document for submission and/or consumes profile conformant documents rather than any particular scanning technology.

18.1 Use Cases

18.1.1 Content Use Cases

Text Chart Notes

Examples of this content include handwritten, typed or word processed clinical documents and/or chart notes. These documents are typically multi-page, narrative text. They include preprinted forms with handwritten responses, printed documents, and typed and/or word processed documents, and documents saved in various word processing formats. Appropriate formats are PDF, derived from the word processing format, or plaintext, if the text structure is all that needs to be conveyed. PDF is desirable because it most faithfully renders word processed document content and it preserves meaning embodied in non-textual annotations.

Graphs, Charts and/or Line Drawings

Examples of this content include Growth Charts, Fetal Monitoring Graphs. Line drawings such as those described above are best rendered using PDF versus an image based compression, such as JPEG. However, when computer generated PDFs include lines or lossy compression is not acceptable for diagnostic purposes, PDF should be used.

Object Character Recognition (OCR) Scanned Documents

Clinical documents can contain text and annotations that cannot be fully processed by optical character recognition (OCR). We call attention to the fact that the OCR text content may only partially represent the document content. These are best supported by converting to PDF format, which can mix the use of OCR'd text, compressed scanned text, and scanned image areas.

Electronic Documents

Existing clinical documents that are electronically transmitted or software created (e.g. PDF, or plaintext) can be considered as actually scanned, previously scanned or virtually scanned before they are shared. In this context, "actually scanned" refers to electronic documents, newly created via some scanning technology from legacy paper or film for the purposes of sharing. "Previously scanned" refers to electronic documents that were previously produced via some scanning technology from legacy paper or film, but have existed in their own right for a period of time. "Virtually Scanned" electronic documents are existing electronic documents not derived from legacy paper or film that either are PDF/A or plaintext format or have been converted to one of these formats for the purposes of sharing. This content is covered by this profile.

18.1.2 Content Creator Use Cases

Content is created by a Content Creator. Impact on application function and workflow is implementation specific and out of scope of this content profile, though we note that they will be compliant with this content profile if they can produce CDA wrapped PDF, CDA wrapped plaintext or both. The following example use case is included to aid in the scoping of this content profile.

Legacy Clinic is a small two-physician clinic. They presently store their patient's medical records on paper. The Clinic is trying to figure out what to do with its paper and word processing documents as it converts over to an electronic system. They would like to be able to view the files over their local intranet.

Presently, most records are handwritten on preprinted paper forms that are inserted into specific sections of the patient's chart. More detailed encounter reports are dictated and sent to a transcription company that returns them in a word processing format. The medical records clerk at Legacy Clinic receives these files via e-mail, decrypts them, prints them out, and adds them to the patient's chart in the correct section.

Over the years, Legacy Clinic has used a number of different transcription companies, and the documents are stored in a variety of word processing formats. Several years ago, they began to require that returned documents be in RTF format in an attempt to reduce frustrations induced by dealing with discrepant word processing formats. Only in some cases was patient and encounter metadata stored within the word processing document in a regular format, depending upon the transcription company used at the time. A third party presently handles labs for the clinic. These are usually returned to the Clinic as printed documents. The clerk inserts these into the labs section in the patient's chart.

In the case of Legacy Clinic, the link between the word processing documents and the patient has been maintained for many of its documents, since the existing manual process maintains that association, and some of the files also contain the encounter metadata. However, the link to the specific encounter will need to be reestablished by interpreting the document content, which will require a great deal of manual effort for some of their documents which do not have it, and will still require custom handling depending upon the format used to store this metadata.

Legacy Clinic uses a transcription provider that can generate PDF documents, wrapped in a CDA Release 2.0 header. These are sent to Legacy Clinic via e-mail. While the same manual process is used, these documents are now in a format that is ready to be used by their new EHR system.

18.1.3 Content Consumer Use Cases

Content is consumed by a Content Consumer. Impact on application function and workflow is implementation specific and out of scope of this content profile. However, we note that adoption of this profile will necessitate the Content Consumer, upon document receipt, support the processing of both CDA wrapped PDF and CDA wrapped plaintext.

18.2 Actors/ Transactions

There are two actors in the XDS-SD profile, the Content Creator and the Content Consumer. Content is created by a Content Creator and is to be consumed by a Content Consumer. The sharing or transmission of content from one actor to the other is addressed by the appropriate use of IHE profiles described below, and is out of scope of this profile. A Document Source or a Portable Media Creator may embody the Content Creator Actor. A Document Consumer, a Document Recipient or a Portable Media Importer may embody the Content Consumer Actor. The sharing or transmission of content or updates from one actor to the other is addressed by the use of appropriate IHE profiles described in the section on Content Bindings with XDS, XDM and XDR.

Figure 20.2-1 shows the actors directly involved in the Scanned Documents Content Integration Profile and the relevant transactions between them. Other actors that may be indirectly involved due to their participation in other profiles are not necessarily shown.

Figure 3.18.2-1. Scanned Documents Actor Diagram

18.3 Scanned Documents Content Integration Profile Options

Options for Scanned Documents leverage those in the Patient Care Coordination (PCC) Technical Framework (TF). Options that may be selected for this Integration Profile are listed in the table 20.3-1 along with the Actors to which they apply. Dependencies between options when applicable are specified in notes.

Table 18.3-1: XDS-SD - Actors and Options

Actor	Options	Vol & Section
Content Creator	*No options defined*	
Content Consumer	*View Option*[1]	PCC TF- 2:4.0.1
	Document Import Option[1]	PCC TF-2:4.0.2
	Document Import Option[1]	PCC TF-2:4.0.2
	Discrete Document Import Option[1]	PCC TF-2:4.0.4

Note 1: The Actor shall support at least one of these options.

18.4 Scanned Documents Bindings to XDS, XDR, XDM

Actors from the ITI XDS, XDM and XDR profiles embody the Content Creator and Content Consumer sharing function of this profile. A Content Creator or Content Consumer may be grouped with appropriate actors from the XDS, XDM or XDR profiles to exchange the content described therein. The metadata sent in the document sharing or interchange messages has specific relationships or dependencies (which we call bindings) to the content of the clinical document described in the content profile. The Patient Care Coordination Technical Framework (PCC-TF) defines the bindings to use when grouping the Content Creator of this Profile with actors from the IHE ITI XDS, XDM or XDR Integration Profiles. See PCC TF-2:5.

18.5 Scanned Documents Content Process Flow

This profile assumes the following sequence of events in creation of an XDS-SD document.

1. A legacy paper document is scanned and a PDF/A is rendered. Alternatively, an electronic document is converted, if necessary, to PDF/A or plaintext format (see ITI TF-2: 5.2.1 and 5.2.1.1).
2. Software, conformant to this profile and most likely with the aid of user input (eg. to provide document title, confidentiality code, original author), renders the CDA R2 header pertaining to the PDF or plaintext produced. The document is wrapped and the XDS-SD document is completed (see ITI TF-2: 5.2.3).
3. XDS metadata is produced from data contained in the CDA header and supplemental information (see ITI TF-1: 5.2.2).
4. The completed XDS-SD document and corresponding metadata is sent via the Provide an Register Document Set Transaction [ITI-15] or [ITI-41] of XDS/XDR, or the Distribute Document Set on Media Transaction [ITI-32] of XDM.

Appendix A: Actor Descriptions

Actors are information systems or components of information systems that produce, manage, or act on information associated with operational activities in the enterprise. The following are definitions of actors used in the IHE IT Infrastructure Integration Profiles:

Audit Repository – This actor provides a repository for audit events. IHE does not specify what analysis and reporting features should be implemented for an audit repository.

Client Authentication Agent – Provides local management of user authentication.

Context Manager – This actor serves as a broker for the communication between two or more context participant actors (either Patient Context Participant or User Context Participant). It supports the passing of the user and patient subjects.

Display – A system that can request specific information or documents from an Information Source and display them.

Document Source - The Document Source Actor is the producer and publisher of documents. It is responsible for sending documents to a Document Repository Actor. It also supplies metadata to the Document Repository Actor for subsequent registration of the documents with the Document Registry Actor.

Document Consumer - The Document Consumer Actor queries a Document Registry Actor for documents meeting certain criteria, and retrieves selected documents from one or more Document Repository actors.

Document Registry - The Document Registry Actor maintains metadata about each registered document in a document entry. This includes a link to the Document in the Repository where it is stored. The Document Registry responds to queries from Document Consumer actors about documents meeting specific criteria. It also enforces some healthcare specific technical policies at the time of document registration.

Document Repository - The Document Repository is responsible for both the persistent storage of these documents as well as for their registration with the appropriate Document Registry. It assigns a URI to documents for subsequent retrieval by a Document Consumer.

DNS Server – This actor has authoritative location information.

Information Source – A system that responds to requests for specific information or documents and returns ready for presentation information to be displays on the requesting actor.

Kerberos Authentication Server – Provides central authentication of enterprise users.

Kerberized Server – Receives user authentication information for further use by the service that contains this actor

Patient Context Participant – This actor participates in a shared context environment by both setting the patient context and responding to context changes as communicated by the Context Manager Actor. This actor shall respond to all patient context changes. This actor shall set the patient context, if the application containing this actor has patient selection capability.

Patient Demographics Consumer – A system that uses demographic information provided by the Patient Demographics Supplier about a patient.

Patient Demographics Supplier – A system responsible for adding, updating and maintaining demographics about a patient, and additional information such as related persons (primary caregiver, guarantor, next of kin, etc.). It supplies new and updated information to the Patient Demographics Consumer.

Patient Encounter Source – A system responsible for adding, updating and maintaining encounter information about a patient. It supplies new and updated information to the Patient Encounter Consumer.

Patient Encounter Consumer – A system that uses patient encounter information provided by the Patient Encounter Source about a patient.

Patient Identifier Cross-reference Consumer – This actor allows a system in a Patient Identifier Domain to determine the identification of a patient in a different Patient Identifier Domain by using the services of a Patient Identifier Cross-Reference Manager Actor.

Patient Identifier Cross-reference Manager – Serves a well-defined set of Patient Identifier Domains. Based on information provided in each Patient Identifier Domain by a Patient Identification Source Actor, it manages the cross-referencing of patient identifiers across Patient Identifier Domains.

Patient Identity Source – - The Patient Identity Source Actor is a provider of unique identifier for each patient and maintains a collection of identity traits. Each Patient Identifier Domain requires this Actor to assign patient identities and to notify other Actors (e.g. a Patient Identifier Cross-reference Manager or a Document Registry Actor) of all events related to patient identification (creation, update, merge, etc.).

Personnel White Pages Consumer – This actor has a use for information that can be found in the Personnel White Pages Directory.

Personnel White Pages Directory – This actor has authoritative Personnel White Pages information on the human workforce members of the enterprise.

Portable Media Creator: This actor assembles the content of the media and writes it to the physical medium. A priori this document set is extracted from an Electronic Healthcare Record (EHR) or a Personal Health Record (PHR) system.

Portable Media Importer: This actor reads and displays the information contained on the media, allows the user to select information, and store any or all of the elements. Typically this document will be integrated into an Electronic Healthcare Record (EHR) or Personal Health Record (PHR) and can then process the instances.

Secure Node – The presence of this actor on a system means that all of the other actors and other non-IHE software comply with the IHE rules for user authentication, communications authentication, and security policies.

Time Client – Establishes time synchronization with one or more Time Servers using the NTP protocol and either the NTP or SNTP algorithms. Maintains the local computer system clock synchronization with UTC based on synchronization with the Time Servers.

Time Server – Provides NTP time services to Time Clients. It is either directly synchronized to a UTC master clock (e.g. satellite time signal) or is synchronized by being grouped with a Time Client to other Time Server(s).

User Context Participant - Receives notification of user context changes and follows them for the application that contains it.

X-Service Provider - System providing a service that needs a X-User Assertion.

X-Service User - System making a services request of an X-Service Provider.

Appendix B: Transaction Descriptions

Transactions are interactions between actors that transfer the required information through standards-based messages. The following are brief descriptions of the transactions defined by IHE.

1. **Maintain Time:** NTP transactions used to maintain time synchronization.
2. **Get User Authentication:** The Client Authentication Agent requests user authentication from the Kerberos Authentication Server. When the user is authenticated, the Kerberos Authentication Server returns a Ticket Granting Ticket (TGT) to optimize future activity.
3. **Get Service Ticket:** Obtain a ticket using Kerberos protocol for use with a service.
4. **Kerberized Communication:** The Kerberized Communication transaction is an aspect of the connection between a local client and a remote server.
5. **Join Context**: Allows a Context Participant Actor to locate and establish communication with the Context Manager Actor.
6. **Change Context:** Includes all messages required to initiate and finalize a context change transaction:
    - Initiation of a context change request from the instigating participant actor
    - Delivery of survey results to instigating actor and display of associated replies
    - Communication of context change decision to the Context Manager Actor
7. **Leave Context**: Allows Context Participant Actor to notify the Context manager Actor that it is breaking off communication.
8. **Patient Identity Feed**: Allows a Patient Identity Source Actor to notify a Patient Identifier Cross-Reference Manager Actor of all events related to patient identification (creation, update, merge, etc.).
9. **PIX Query**: This transaction allows a Patient Identifier Cross-reference Consumer to find out the identification of a patient in different Patient Identifier Domains by using the services of a Patient Identifier Cross-reference Manager Actor.
10. **PIX Update Notification:** Allows a Patient Identifier Cross-reference Consumer to be notified by the Patient Identifier Cross-reference Manager Actor of changes to the identification of all patients in Patient Identifier Domains the Consumer is interested in.
11. **Retrieve Specific Information for Display:** A request issued by a display system for specific information related to a patient returned in a ready for presentation information format.
12. **Retrieve Document for Display:** A display system requests an instance of a uniquely identified persistent document under custodianship by an information source and receives its content ready for presentation.
13. **Follow Context:** Accounts for all messages required to propagate a context change to a responding participant actor:
    - Survey of all other Context Participant Actors by the Context Manager Actor and display by the instigating Participant Actor of any associated replies

- Notification of context change result from the Context manager Actor to the Context Participant Actors
- Retrieval of the context data by the Context Participant Actors

14. Provide and Register Document Set

A Document Source Actor initiates the Provide and Register Document Set Transaction. For each document in the submitted set, the Document Source Actor provides both the documents as an opaque octet stream and the corresponding metadata to the Document Repository. The Document Repository is responsible to persistently store these documents, and to register them in the Document Registry using the Register Documents transaction by forwarding the document metadata received from the Document Source Actor.

15. Register Document Set

A Document Repository Actor initiates the Register Document Set transaction. This transaction allows a Document Repository Actor to register one or more documents with a Document Registry, by supplying metadata about each document to be registered. This document metadata will be used to create an XDS Document Entry in the registry. The Document Registry Actor ensures that document metadata is valid before allowing documents to be registered. If one or more documents fail the metadata validation, the Register Document Set transaction fails as a whole.

16. Query Registry

The Query Registry transaction is issued by the Document Consumer Actor on behalf of a care provider (EHR-CR) to a Document Registry. The Document Registry Actor searches the registry to locate documents that meet the provider's specified query criteria. It will return a list of document entries that contain metadata found to meet the specified criteria including the locations and identifier of each corresponding document in one or more Document Repositories.

17. Retrieve Document

A Document Consumer Actor initiates the Retrieve Document transaction. The Document Repository will return the document that was specified by the Document Consumer.

18. Intentionally Left Blank

19. **Node Authentication:** This transaction is embedded within all network communications activity. All DICOM, HL7, and HTML connections shall comply with the IHE specification for bi-directional authentication and authorization of communications of Protected Healthcare Information (PHI). IHE does not specify how other protocols that transfer PHI shall perform bi-directional authentication and authorization, but requires that other protocols perform such authentication and authorization.

20. **Record Audit Event**: The delivery of an audit event description from any secure node to the Audit Repository.

21. **Patient Demographics Query**: Look up and return patient demographic information in a single patient demographics source, based upon matches with full or partial demographic information entered by the user.

22. **Patient Demographics and Visit Query:** Look up and return patient demographic and visit information in a single patient demographics source, based upon matches with full or partial demographic/visit information entered by the user.
23. **Find Personnel White Pages:** This transaction will find the LDAP Directory by querying the DNS.
24. **Query Personnel White Pages:** This transaction provides for read-only access to the Personnel White Pages directory.
25. **reserved for** Send Notification
26. **reserved for** Receive Notifications
27. **reserved for** Send Acknowledgement
28. **reserved for** Receive Acknowledgement
29. **reserved for** Cross Enterprise User Authentication
30. **Patient Identity Management –** The Patient Demographics Supplier registers or updates a patient and forwards the demographic information (i.e., all information directly related to the patient, such as ID, address, next of kin, guarantor, etc.) to other systems implementing the Patient Demographics Consumer Actor.
31. **Patient Encounter Management –** The Patient Encounter Supplier registers or updates an encounter (inpatient, outpatient, pre-admit, etc.) and forwards the information to other systems implementing the Patient Encounter Consumer Actor. This information will include the patient's location and care providers for a particular (usually current) encounter.
32. **Distribute Document Set on Media** - A source actor (Portable Media Creator) writes a set of documents on an interchange media. The media is physically transported to another actor (Portable Media Importer) which then imports the document set, or sent as a ZIP attachment via Email. The media can also be provided to a patient or a referring physician for web-based viewing.
33. **placeholder**
34. placeholder
35. placeholder
36. placeholder
37. placeholder
38. placeholder
39. placeholder
40. **Provide X-User Assertion** - This transaction provides a trustable user assertion from the service user to the service provider

Appendix C: IHE Integration Statements

IHE Integration Statements are documents prepared and published by vendors to describe the conformance of their products with the IHE Technical Framework. They identify the specific IHE capabilities a given product supports in terms of IHE actors and integration profiles (described in ITI TF-1: 2).

Users familiar with these concepts can use Integration Statements to determine what level of integration a vendor asserts a product supports with complementary systems and what clinical and operational benefits such integration might provide. Integration Statements are intended to be used in conjunction with statements of conformance to specific standards (e.g. HL7, IETF, DICOM, W3C, etc.).

IHE provides a process for vendors to test their implementations of IHE actors and integration profiles. The IHE testing process, culminating in a multi-party interactive testing event called the Connect-a-thon, provides vendors with valuable feedback and provides a baseline indication of the conformance of their implementations. The process is not intended to independently evaluate, or ensure, product compliance. In publishing the results of the Connect-a-thon and facilitating access to vendors' IHE Integration Statements, IHE and its sponsoring organizations are in no way attesting to the accuracy or validity of any vendor's IHE Integration Statements or any other claims by vendors regarding their products.

IMPORTANT -- PLEASE NOTE: Vendors have sole responsibility for the accuracy and validity of their IHE Integration Statements. Vendors' Integration Statements are made available through IHE simply for consideration by parties seeking information about the integration capabilities of particular products. IHE and its sponsoring organizations have not evaluated or approved any IHE Integration Statement or any related product, and IHE and its sponsoring organizations shall have no liability or responsibility to any party for any claims or damages, whether direct, indirect, incidental or consequential, including but not limited to business interruption and loss of revenue, arising from any use of, or reliance upon, any IHE Integration Statement.

C.1 Structure and Content of an IHE Integration Statement

An IHE Integration Statement for a product shall include:

1. The Vendor Name
2. The Product Name (as used in the commercial context) to which the IHE Integration Statement applies.
3. The Product Version to which the IHE Integration Statement applies.
4. A publication date and optionally a revision designation for the IHE Integration Statement.
5. The following statement: "This product implements all transactions required in the IHE Technical Framework to support the IHE Integration Profiles, Actors and Options listed below:"
6. A list of IHE Integration Profiles supported by the product and, for each Integration Profile, a list of IHE Actors supported. For each integration profile/actor combination, one or more of the options defined in the IHE Technical Framework may also be stated.

Profiles, Actors and Options shall use the names defined by the IHE Technical Framework Volume I. (Note: The vendor may also elect to indicate the version number of the Technical Framework referenced for each Integration Profile.)

Note that implementation of the integration profile implies implementation of all required transactions for an actor as well as selected options.

The statement shall also include references and/or internet links to the following information:

1. Specific internet address (or universal resource locator [URL]) where the vendor's Integration Statements are posted
2. URL where the vendor's standards conformance statements (e.g., HL7, DICOM, etc.) relevant to the IHE transactions implemented by the product are posted.
3. URL of the IHE Initiative's web page for general IHE information www.himss.org/ihe.

An IHE Integration Statement is not intended to promote or advertise aspects of a product not directly related to its implementation of IHE capabilities.

C.2 Format of an IHE Integration Statement

Each Integration Statement shall follow the format shown below. Vendors may add a cover page and any necessary additional information in accordance with their product documentation policies.

IHE Integration Statement		**Date**	**12 Oct 2003**
Vendor	**Product Name**	**Version**	
Any Medical Systems Co.	IntegrateRecord	V2.3	
This product implements all transactions required in the IHE Technical Framework to support the IHE Integration Profiles, Actors and Options listed below:			
Integration Profiles Implemented	**Actors Implemented**	**Options Implemented**	
Retrieve Information for Display	Information Source	none	
Enterprise User Authentication	Kerberized Server	none	
Patient Identity Cross-referencing	Patient Identifier Cross-reference Consumer	PIX Update Notification	
Internet address for vendor's IHE information:	www.anymedicalsystemsco.com/ihe		
Links to Standards Conformance Statements for the Implementation			
HL7	www.anymedicalsystemsco.com/hl7		
Links to general information on IHE			
In North America: www.himss.org/ihe	In Europe: www.ihe-europe.org	In Japan: www.jira-net.or.jp/ihe-j	

Appendix D: User Authentication Techniques - Passwords, Biometrics, and Tokens

Authentication techniques are based on one or more of three factors: Something you know, something you are, or something you have. There are many different authentication techniques in use today. The technologies supporting these techniques are not well standardized. There are also excellent security reasons to avoid specifying any single set of technologies for authentication use.

The Kerberos protocol was originally defined to work with any user authentication technique. Kerberos has been shown to support a wide variety of authentication technologies. These include various forms of tokens and biometric technologies. Specific implementations of these technologies often include proprietary components. There is often a pair of proprietary components added – one at the user workstation and a matching component at the authentication server. Once the user authentication is complete, the subsequent Kerberos transactions are the same.

These extensions are not yet standardized. The IHE specification for the use of Kerberos does not prevent the use of these extensions at a specific site, nor does it ensure that the extensions will work.

The Kerberos system specified for the Enterprise User Authentication utilizes a challenge response system together with a username and password system to authenticate the user. The minimal support of passwords provides a standardized baseline for the IHE “Enterprise User Authentication”. Kerberos enables enforcement of a central password policy which facilitates stronger passwords. Such password policies are beyond the scope of IHE. Kerberos does not prevent the use of weak passwords. The password strength policy must be chosen and enforced by the site security administration.

Appendix E: Cross Profile Considerations

E.1 Combined use of RID, EUA and PIX Integration Profiles

When used alone, the Retrieve Information for Display Integration Profile assumes that the Patient Identifier Domain is the same for both the Display and the Information Source Actors. Furthermore, any user authentication on the Information Source is not addressed explicitly. This Appendix discusses combination of the Retrieve Information for Display Integration Profile with other IHE Integration Profiles to address these two problems.

When used in conjunction with the Patient Identifier Cross-referencing Integration Profile, implementations of the Retrieve Information for Display Integration Profile shall take into account that the Information Source Actor may need to map Patient IDs from different identifier domains to the one used in its own domain. The combined use of these Integration Profiles is achieved by grouping the Information Source and the Patient Identifier Cross-reference Consumer Actors. This is depicted in Figure E-1.

Similarly, the Information Source Actor may perform certain access control functions based on the requesting user authentication performed by the actors implementing the Enterprise User Authentication Integration Profile. The combined use of these Integration Profiles is achieved by grouping the Display Actor with the Client Authentication Agent Actor and the Information Source Actor with the Kerberized Server Actor. This is also shown in Figure E-1.

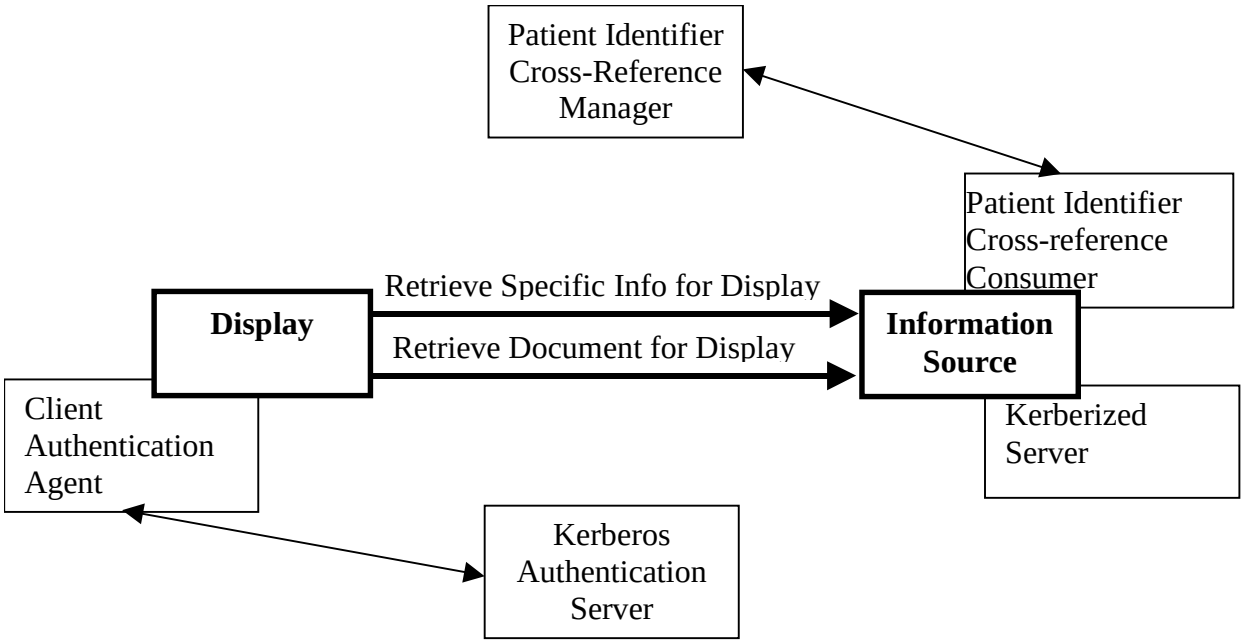

Figure E-1. Combined use of actors implementing multiple Integration Profiles

E.2 XDS Integration with RID

The RID Retrieve Document for Display transaction [ITI-12] is compatible with the XDS Retrieve Document transaction [ITI-17]. Thus, an RID Information Source implementing the Retrieve Document for Display transaction can be used to implement the XDS Retrieve Document transaction. In this instance, the RID Information Source must be a secure node [see ATNA].

E. 3 XDS Integration with PIX

All Patient IDs managed in the XDS transactions (either in XAD-Pid Domain or in an EHR-CR Domain) shall include the related Patient Domain ID (OID of the Assigning Authority) associated with the patient ID. It is recommended that this unambiguous patient identification be used with Patient IDs within the Documents also.

Because XDS is Document content neutral, there is no verification by the XDS Repository that the Patient IDs included inside the documents are consistent with the patient IDs managed by the Registry in the document entry related to that document.

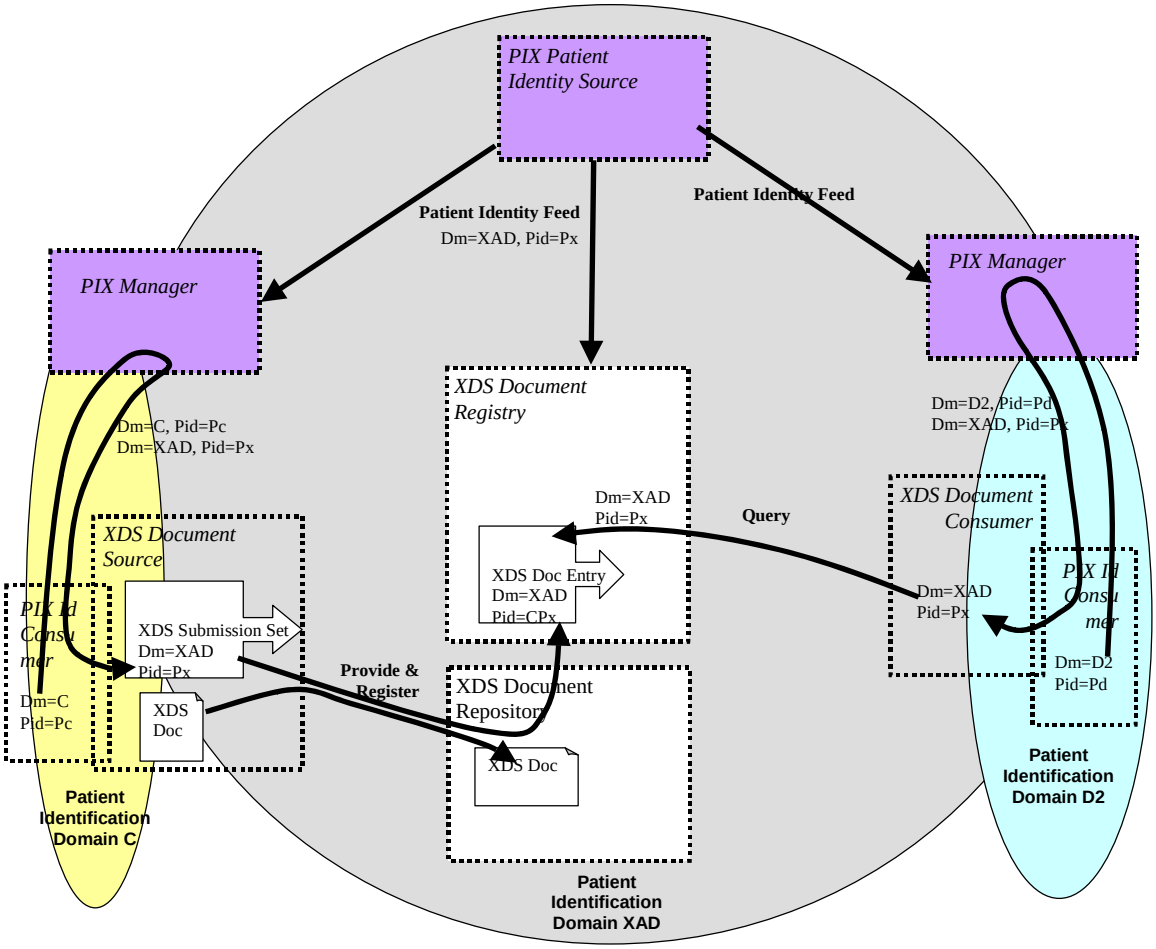

Figure E.3-1 XDS Affinity Domain with patient ID cross-referencing with IHE PIX Managers

Figure C.6-1 depicts an example of an XDS Affinity Domain with a Patient Identifier Domain (called XAD) and two EHR-CRs where the cross-referencing is performed by Patient Identifier Cross Referencing Managers internal to both the Document Source and the Document Consumer Domains (called C and D2 respectively).

A Document Source may choose to perform the cross-referencing of its own patient IDs in that of the XAD-Pid Domain by leveraging the IHE PIX Integration Profile (See Figure). The Patient ID Feed Transaction from the XAD Patient ID Source may be used to provide input to the Patient Identifier Cross-Referencing Manager used by the Document Source. The PIX manager may either be internal to the EHR-CRs or be shared across the XDS Affinity Domain.

E.4 XDS Integration with PWP

The XDS Document Source Actor in the XDS Integration Profile may choose to utilize the PWP Query Personnel White Pages [ITI-24] transaction to obtain information needed to fill the authorPerson and legalAuthenticatorName fields for the XDS Register Document Set [ITI-14] and Provide & Register Document Set [ITI-15] transactions.

The Personnel White Pages transaction defines, in ITI-TF 2:3.24.4.1.2.3.1, a “cn” attribute with “lang-x-ihe” that contains the information in the HL7 XCN (extended composite ID number and name for persons) format for personal information. These fields are optional in the PWP Integration Profile. A care delivery organization may choose to populate these fields in their Personnel White Pages Directory and utilize the ITI-24 transaction to support its XDS activities. This is not a required dependency, but is a possible reason to group a Document Source Actor with a Personnel White Pages Consumer Actor.

The PWP Integration Profile only provides the personnel information. Organizational information must be obtained via other means, e.g. extending the LDAP directory with organizational objects.

E.5 XDS Integration with PDQ

The Patient Demographics Query (PDQ) Integration Profile may be used in conjunction with the XDS Integration Profile to provide a lookup for XDS Affinity Domain Patient Identifiers to XDS Document Consumer and Document Source Actor. In this case a Patient Demographics Supplier Actor needs to be grouped with the XDS Patient Identifier Source Actor on one hand, and on the other hand a Patient Demographics Consumer Actor needs to be grouped with the Document Source/Consumer where one may want to query based on local patient traits and obtain a pick-list of candidate Patient Ids in the XAD Patient Identifier Domain. This offers a simpler solution that the use of the PIX Integration Profile.

E.6 XDM Integration with XDS, Content Integration Profiles, PIX, and DSG

The XDM Profile does not constrain the document types or purposes. Content Integration Profiles may impose such constraints.

The XDM Profile does not address the issue of patient reconciliation. The PIX and PDQ functionality might be available to a Portable Media Importer, but the XDM Profile does not require it. If there is no PIX or PDQ available to the Portable Media Importer, some other method for performing the necessary coercion of patient identifiers must be provided. This might be manual for Portable Media Importers that are intended for very small sites.

The Cross-enterprise Document Media Interchange (XDM) Integration Profile may be used in conjunction with the DSG Integration Profile to provide for the digital signature of the documents content and of the XDS metadata.

E. 7 XDM/XDR Distinction

Both XDR and XDM describe the exchange of a set of patients' documents. They are relevant in situations where XDS is not yet implemented or available at one of the participating organizations or where point-to-point (versus sharing through a registry) interaction is desired.

XDM is applicable in situations where the information receiver is an individual who will manually interpret or examine the data and associated documents. XDM allows for one exchange which contains documents relating to multiple patients and can be used in situations where no continuous networking capability is available on one or both of the participating healthcare providers.

XDR is applicable in situation where the information exchanged is going to an automated application or robust system capable of automated storage or process of documents relative to one patient. XDR requires continuous networking capability between the healthcare providers exchanging data.

Appendix F: Request to Standards Development Organizations

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Appendix G: Security Considerations

G.1 Cross Profile Considerations

IHE compliant systems usually process private healthcare information. This is subject to national privacy regulations, and possibly other state and contractual requirements. The IHE Infrastructure profiles do not fully define the security mechanisms necessary to protect this information. The Enterprise User Authentication profile provides one component of this solution.

IHE assumes that actors will be installed on nodes with the following characteristics:

- Each node has a security policy and procedure that applies to its operation. This is assumed to be part of the healthcare enterprise security policy.
- Any user (human, or application process) external to the node boundaries is submitted to an access control procedure in which the user/application will be authenticated.
- All required audit trail events are captured and recorded.

The profiles in this framework assume the following environment:

- Physical Security Environment
    - The equipment is assumed to be located in a physically protected and actively monitored area. This is normally the case with modality equipment because of other patient safety, privacy, and operational concerns. Similarly, the HIS systems and various archives are normally protected. Equipment like PACS workstations is sometimes placed in unprotected areas, but it is usually located where hospital staff monitors and limit access. It assumes that the threat of equipment modification is protected against by means of the physical security mechanisms.
    - The network equipment that connects the computers is also assumed to be physically protected against unauthorized connections and unauthorized modifications. In the treatment areas of most hospitals the network equipment is in ceilings, cableways, locked cabinets, and other protected areas. There is usually staff present to monitor that no unauthorized activity is taking place.
    - Local procedures and operations will be in place to ensure that the physical security assumptions are valid for other areas of the hospital, such as administrative offices, that may be at greater risk.
    - Remote locations, especially home offices, are not physically protected. Other means will be used to provide equivalent protection. This may include the use of technology such as VPN connections or HTTPS encryption. Use of encryption or VPN is not a complete replacement for physical security but may be part of an overall protection system.

- The home computer that is used for both personal and professional purposes is difficult to protect. It will be protected from inadvertent modification by malicious software or its use will be prohibited.

- Network Security Environment
  - In addition to the physical security of the network, there will be protection against network access by unsupervised systems. This is typically provided by mechanisms such as firewalls and VPNs.

The threat profile is assumed to be:

- Accidental and inadvertent misuse
- Individual abuse for personal gain, malice, revenge, or curiosity. The abusers are assumed to have only limited access to the underlying systems and software. They are not expert at the internal structure of the systems.
- Random untargeted abuse, such as from an Internet hacker.

The threat profile also assumes that the following threats are either not present or otherwise protected.

- Individual abuse by a system administrator, system developer, or other expert.
- Military or hostile government action
- Organized criminal attack

IHE addresses only those security requirements related to IT systems within the scope of IHE healthcare applications. It does not address security requirements for defending against network attacks, virus infection, etc.

IHE does not mandate the use of encryption because the performance impact of current encryption algorithms is excessive. Most hospital networks provide adequate security through physical and procedural mechanisms. The additional performance penalty for encryption is not justified for these networks. The profiles permit the use of encryption so that it can be used as part of an overall security plan.

G. 2 XDS Security Considerations

Security and privacy

Coordinating the security and privacy policies of all the care delivery organizations in an XDS Affinity Domain may be a challenge. An agreement is needed on security procedures, goals, auditing, record keeping, etc. This can result in changes to other enterprise policies, such as human resources procedures. XDS Affinity Domain members are delegating full access to their published data to the other members of the XDS Affinity Domain. This relationship requires a close ongoing partnership that ensures ongoing maintenance of policies, procedures, and activities. If laws change, relevant policies must be adjusted throughout the group. Corporate changes to group members affect the policies. Security events must be managed as a group. This must be managed as a long-term activity, not a one-time event.

Particular problem areas are likely to be:

- Authorized access and modification policies. The details of access policies are likely to have enterprise differences and conflicts that must be resolved. The XDS Affinity Domain relationships also introduce new policy requirements. For example, changes to employment (e.g. employee hiring and firing) must now include suitably rapid notifications to other XDS Affinity Domain members. Changes to privacy restrictions (e.g. divorces) now require full XDS Affinity Domain notifications, not merely enterprise notifications.
- Audit trail and access record keeping are often quite sensitive internal enterprise activities that must now be appropriately coordinated with the full XDS Affinity Domain.
- Changes to laws and regulations now affect not only the policies of the individual enterprises; they also must be reflected in the XDS Affinity Domain relationship contracts, policies, and procedures.
- Patient access and patient identity management. Patients usually have insecure computers. Patients often object to security procedures.
- Trans-border communication of Personal Health Information (PHI) often presents legal and regulatory issues.

ITI TF-2: Appendix J in volume II goes into more detail listing many of the threats, objectives, policies, and mitigations that need to be coordinated among XDS Affinity Domain members.

The XDS Integration Profile for two main reasons does not prescribe such Security and Privacy policies. First, it is clear that the broad range of possible solutions to these policies that will depend on the legal framework and the types of healthcare system, calls for XDS to be offer such flexibility. Decisions in this domain will have some impact on the implementations of XDS Actors, but it is expected that these will be minimal.

Appendix H: Intentionally Left Blank

Appendix I: Intentionally Left Blank

Appendix J: Content and Format of XDS Documents

The XDS Integration Profile purposely leaves a number of policies up to the XDS Affinity Domain to decide, including the structure and format of the content of XDS Documents to be shared, the mapping of content metadata into the XDS Document Registry, the coding of XDS Document metadata, the events that trigger an XDS Submission Request, and the policies concerning the use of XDS Folders to facilitate sharing.

It is important to recognize that until sufficient experience has been gained in cross-enterprise document sharing, it is not possible to establish common or even best practices in the use of the XDS Integration Profile. IHE has therefore chosen to abstain to make recommendations in these topics at this time.

IHE also recognizes that there will be a need for content-oriented integration profiles to be used in cooperation with this Integration Profile. It is expected that in the future the various IHE Domains (Patient Care Coordination, Cardiology, Laboratory, Radiology, IT Infrastructure, etc.) will produce IHE Integration Profiles refining the use of XDS within the domain. These various content-oriented integration profiles may rely on XDS, but would further constrain the forms of documents to be shared, or the uses of XDS features such as Folders and Submission Sets, et cetera.

Content Neutrality

XDS is content neutral. It neither prescribes nor prohibits the format, content, structure or representation of documents that can be retrieved from an XDS Document Repository. For the XDS Integration Profile to have immediate value to an XDS Affinity Domain, it must be able to adapt to the documents that are present and available from its members. Thus, prohibitions on content would only serve to limit the utility and adoption of the XDS Integration Profile. Similarly, XDS Affinity Domains must be able to adapt to emerging standards, which cannot be enumerated in any list of prescribed content formats.

IHE strongly recommends that XDS Affinity Domains adopt rules that require documents to comply with widely accepted standards where possible (*e.g.*, HL7 CDA, CEN ENV 13606, ASTM CCR, and DICOM Composite Object).

Document Headers and Metadata

Because XDS is content neutral, XDS cannot validate metadata contained within the body of an XDS document against the metadata supplied to the XDS Document Registry. XDS Affinity shall therefore select content where IHE has defined Integration Profiles, or until that point, the XDS Affinity Domains shall carefully define how the attributes in the XDS Document Registry are filled.

Metadata and the Patient Record

Although metadata in the document header may be duplicated in the XDS Document Registry, the XDS Document Registry metadata has a particular role in term of being part of the legal medical record stored. It is definitively not part of the clinical record as managed by the XDS Document Repositories where documents reside. Furthermore, XDS does not provide for transactions to “sign” or legally authenticate the content of an XDS Submission Set (See IHE Document Digital Signature Content Profile- DSG), although it offers the ability to track its author, if the XDS Affinity Domain so desires to enforce it. The contents of XDS Folders are tracked, through the Submission Sets that

contributed to placing document references in folders. However, the existence of document metadata in the registry and the potential medical acts involved in creating an XDS Submission Set or XDS Folder may make the contents of the XDS Document Registry part of the patient's legal medical record. It will be up to individual XDS Affinity Domains to decide how to address the issues involved with these clinical acts and to resolve them in accord with common sense, acceptable medical practices, and local regulations.

Appendix K : XDS Concept Details

K.1 XDS Document Concept

An XDS Document is the smallest unit of information that may be provided to a Document Repository Actor and be registered as an entry in the Document Registry Actor.

An XDS Document is a composition of clinical information that contains observations and services for the purpose of exchange with the following characteristics: Persistence, Stewardship, Potential for Authentication, and Wholeness. These characteristics are defined in the HL7 Clinical Document Architecture Release 1 specification.

An XDS Document may be human and/or application readable. In either cases, it shall comply with a published standard defining its structure, content and encoding. IHE intends to define content-oriented Integration Profiles relying on such content standards to be used in conjunction with XDS.

Furthermore:

1. When submitted for sharing, an XDS Document shall be provided to the Document Repository Actor as an octet stream with an associated MIME type.

2. When retrieved through the Retrieve Document transaction, an XDS Document shall be unchanged from the octet stream that was submitted (full fidelity repository).

   Note: An XDS Document may be a MIME multipart document (e.g. an HL7 CDA as its first part followed by attachments as files). The first part of the multi-part contains the primary part of the document, other parts are direct attachments to the primary part. The Document Repository handles this multi-part data set as an "opaque entity". The Document Repository does not need to analyze or process its multi-part structure nor the content of any parts in the context of the XDS Integration Profile.

   Note: An XDS Document may be retrieved using alternate methods using document specific retrieval methods. Such optional capabilities are not provided in the current specification of XDS, but are possibly candidates for addition as future options this Integration Profile.

3. An XDS Document shall be associated with metadata defined by the Document Source. This metadata information shall be placed by the XDS Registry Actor in an XDS Document Entry, and is used for query purposes by XDS Consumer Actors.

4. The XDS Integration Profile manages XDS Documents as a single unit of information, it does not provide mechanisms to access portions of an XDS Document. Only the Document Sources or Document Consumers have access to the internal information of the XDS Document.

5. An XDS Document is globally uniquely identified, so that no two XDS Documents with different content shall bear the same Unique Identifier. This identifier is unique across all XDS Affinity Domains, which allows potential merger of XDS Document Repositories from different domains, or exchange of XDS Documents between Clinical Affinitiy Domains, if so desired.

6. The XDS Document Registry Actor shall maintain a single document entry for each XDS Document stored in a Document Repository Actor. Duplicate copies of the same XDS Document (with the same unique identifier) may be stored and registered. Registration of an XDS Document with the same unique identifier but a different content is rejected.

7. This Integration Profile specifies the metadata required for each XDS document registered in the Document Registry. It is the responsibility of the Document Source to ensure that the XDS Document metadata reflects the actual content of the associated XDS Document. Neither the Document Repository nor the Document Registry checks this consistency.

8. The Document Source maintains the following responsibilities over the XDS Documents it has registered:

    a. It has rights to change the status of any of these Documents from "approved" to "deprecated" or to delete them outright.

    b. It has rights to submit an XDS Document with a "Parent Relationship" of replacement ("RPLC") for one of its previously submitted document[4].

XDS Affinity Domains should have policies and procedures to provide patient access to these operations where necessary. For example, in certain regions, patients may request the removal of documents from the EHR-LR. The Registry and Repositories implementations should be ready to support these local operations although there are no IHE transactions defined at this time.

K.2 Concept of an XDS Affinity Domain

An XDS Affinity Domain is made of a well-defined set of Document Repositories and Document Consumers that have agreed to share the clinical documents. An XDS Affinity Domain has a number of properties defined:

1. An XDS Affinity Domain does not deliver care. Only the EHR-CRs belonging to an XDS Affinity Domain as Document Sources and Consumers do.

2. An XDS Affinity Domain is managed by a single Document Registry Actor.

    Note: A distributed registry approach will be considered as a future and separate Integration Profile. For Document Source and Document Consumer Actors, the perception of a single Document Registry Actor hides the complexity of a distributed registry.

3. It includes any number of Document Repository Actors (a distributed configuration is the default, however, a centralized configuration with a grouped Registry/Repository is also supported).

4. It contains an explicit list of Document Consumer and Document Repository actors that participate in document sharing. The addition of a Document Repository or Document Consumer Actor is an administrative task that requires involvement of authorities maintaining the Registry and Repositories.

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[4] For example, in DICOM, where the document identity does not change even though its internal patient metadata may have been updated, the Document Source would submit an updated DICOM Document as a replacement for the existing one.

5. There is a chain of trust established between the users (healthcare staff) in each EHR-CR and the XDS Affinity Domain.
6. Document Repositories and Document Consumers may belong to more than one XDS Affinity Domain and share the same or different documents. This is an implementation strategy and will not be further described.
7. The XDS Affinity Domain supports a primary Patient Identification Domain that is used by the Document Source and Consumers to communicate with the Document Registry. When Document Sources and Consumers in the XDS Affinity Domain belong to different Patient Identifier Registration Domains, the Document Source and Consumers must cross-reference their own Patient Identifier Registration Domains to that of the Registry. They may use the IHE Patient Identifier Cross-referencing Integration Profile, the IHE Patient Demographics Query Integration Profile or other XDS Affinity Domain specific mechanisms for cross-referencing (See ITI TF-2 Appendix E Sections E.3 and E.5).
8. A Document Source may only contribute documents with Document Codes and Health Facility Codes that draw from a Vocabulary Value Set that is approved by the XDS Affinity Domain.

K.3 Other Principles of XDS

The XDS Integration Profile has been designed with the following limitations and principles:

1. A Document may contain references to other documents in its content which are not under the management of the XDS Document Registry. Such references may be available to the EHR-CR that registered the document that includes the reference. It is beyond the scope of XDS to provide access to such documents internal to the EHR-CR.
2. The XDS Repositories are not expected to perform any processing or translations on document content. Processing and translation are the responsibility of a Source EHR-CR or Consumer EHR-CR. The analysis, cross-document combination and presentation of document content are outside the scope of the XDS Integration Profile and its actors.
3. The custodianship for the clinical information contained in a registered document remains with the Source Actor of the EHR-CR. The EHR-LR offers only a “shared space” under the responsibility of each contributing EHR-CR. Through XDS, replacement or deletion of documents in the EHR-LR may only be initiated by the corresponding EHR-CR Source.
4. When an XDS Document that has already been registered in the XDS Registry of an XDS Affinity Domain is resubmitted as if it was a new XDS Document with the same Document Unique identifier, this “duplicate submission” is detected by the Repository and/or Registry based on the fact that the XDS Document Unique Identifier already exists in a Document Entry. The submission request to which that resubmitted Document belongs shall be rejected in the case where the identifiers match but the actual content differs (detected by use of a hash key computed by the Document Repository at the time of submission).

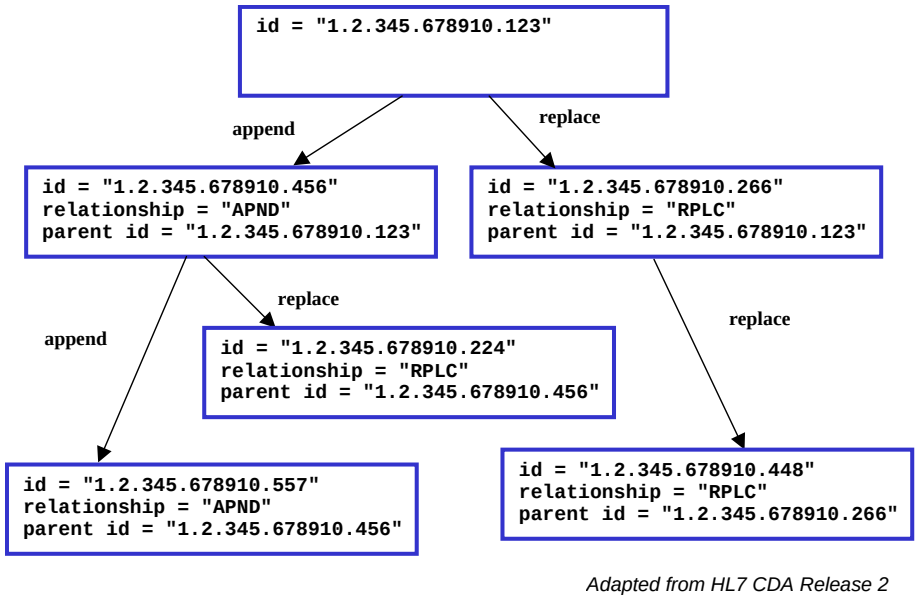
K.4 Document Identification

In order to reduce the number of unique identifiers associated with an XDS Document, the globally unique Document Id assigned by the document source and the unique XDS Document Id used by

the Repository are the same. It is strongly recommended to limit the use of the Document Entry UUID created per ebRS in order to reference the document entry for referencing internally to the encoding of the IHE transactions operations, and to encourage the use of the globally unique Document Id for all external operations (e.g. links maintained in data bases internal to the Document source Actor, links within documents, etc.).

The XDS Document Entry includes two separate attributes: an XDSDocument.uniqueId and XDSDocument.URI, a Universal Resource Identifier. The URI is a "self contained" web method that allows any Document Consumer to perform a Retrieve Document transaction (See ITI TF-2: Section 3.17). The Document Unique ID is a location independent identifier. As the result of XDS Document migration from one XDS Document Repository to another one within an XDS Affinity Domain, the URI would be changed, but not the Document unique ID.

K.5 Example of Document Relationship



Adapted from HL7 CDA Release 2

Figure K.5-1 Example of Document Relationships

These relationships are illustrated in the above figure. Typical scenarios are a simple replacement (e.g. XDSDocument.id "1.2.345.678910.266" replacing XDSDocument.id "1.2.345.678910.123") and a simple addendum (e.g. XDSDocument.id "1.2.345.678910.456" appends XDSDocument.id "1.2.345.678910.123"). More complex scenarios that might be anticipated include:

1. Replacement of an addendum (e.g. XDSDocument.id "1.2.345.678910.224" replaces XDSDocument.id "1.2.345.678910.456", which itself is an addendum to XDSDocument.id "1.2.345.678910.123") - expected behavior would be to render the replacement as the addendum (e.g. render XDSDocument.id "1.2.345.678910.224" as the addendum to XDSDocument.id "1.2.345.678910.123");
2. Addendum to a replaced document (e.g. XDSDocument.id "1.2.345.678910.456" appends XDSDocument.id "1.2.345.678910.123", which has been replaced by XDSDocument.id

"1.2.345.678910.266") - expected behavior would be to render the addendum along with the replacement (e.g. render XDSDocument.id "1.2.345.678910.456" as an addendum to XDSDocument.id "1.2.345.678910.266").

K.6 Off-Line transaction mode

Document Source Actors are allowed to be off-line part of the time, as in the case of a doctor's office system connected only by a dial-up line acting as a Document Source.

The Document Registry and Document Repositories should be designed to be on-line all the time (see note for exception).

Note: The Document Repository may be off-line also in the degenerate case of point-to-point e-mail transmission, where the XDS Affinity Domain is made only of two systems; on one hand a document source and on the other a document repository grouped with the Document registry and Document Consumer (See ITI TF-1: 10;5 Strategy 3).

Information sent to off-line systems will be supported through Internet e-mail protocols. E-mail protocols provide mechanisms for sending acknowledgments:

- Delivery receipts from the end-user, and
- Delivery failure notices from intermediate store-and-forward SMTP servers.

When using e-mail protocols, the asynchronous nature of the acknowledgments, which are delivered by e-mail messages, requires that the Send and Acknowledge components of the transaction be separated into distinct messages.

Body of the e-mail message should contain a simple notice (in English/ASCII), fixed subject line, address should be used for automated processing. An attachment formatted in the local language should contain instructions. Transaction should be included in a separate attachment.

Appendix L: XDS Affinity Domain Definition Checklist

The concept of an XDS Affinity Domain is defined in ITI TF-1:10 and Appendix K. ITI TF Appendix L originally provided an informative checklist for the key policies that need to be addressed in order to deploy an EHR-LR document sharing environment for an XDS Affinity Domain. However, it was recognized that this checklist was incomplete as it did not deal with many necessary XDS Affinity Domain deployment issues. In order to address these shortcomings, a new "Template for XDS Affinity Domain Deployment Planning" White Paper has been created:

http://www.ihe.net/Technical_Framework/index.cfm#IT

It takes the form of a template rather than a checklist because it acts more as an outline for all the issues that should be considered, rather than a checklist to be used to verify the correctness of a particular implementation. This new template can be used when defining policies for either an individual XDS Affinity Domain, or multiple XDS Affinity Domains within a particular nation or region.

Here is a summary of the topics defined in the new "Template for XDS Affinity Domain Deployment Planning":

- Organizational Rules
    - Structure, Roles, Transparency, Legal Considerations and Enforcement
- Operational Rules
    - Service Level Agreements, Daily Governance, Configuration Management, Data Retention, Archive, and Backup, and Disaster Recovery
- Membership Rules
    - Acceptance, Types of Membership, Membership Policies
- Connectivity to the XDS Affinity Domain from External Systems
    - System Architecture
- Global Architecture, Affinity Domain Actors, Transaction Support
    - Terminology and Content
- Refinement of Metadata and Content Attribute Use
    - Patient Privacy and Consent
- Access and Use, Patient consent, and Override Guidelines
    - Technical Security
- Authorization, Role Management, User/Role Authentication, Node Authentication, Certificates Management, Information Access Security, Information Integrity, Updates, and Maintenance Policies, Secure Audit Trails, Consistent Time, Audit Checks, and Risk analysis

Appendix M: Cross-Enterprise Document Sharing and IHE Roadmap

The IHE Cross-Enterprise Document Sharing Integration Profile is part of a family of IHE Integration Profiles grouped in a number of domain-specific Technical Frameworks Patient Care Coordination, Cardiology, Laboratory, Radiology, IT Infrastructure, etc.). XDS is a central foundation for Cross-Enterprise interoperability that may be combined with a number of the existing IHE Integration Profiles (See ITI TF-1:Appendix E). However a number of new IHE Integration Profiles need to be developed, pending the availability of the relevant base standards.

M.1 Document Content Integration Profiles for XDS

It is expected that the various IHE Domains (Cardiology, Laboratory, Radiology, IT Infrastructure, etc.) will produce new IHE Integration Profiles addressing the content of the documents that need to be shared. These various "content-oriented" Integration Profiles will rely on the XDS Integration Profile for managing the registration, discovery and access processes in a common manner.

Such an effort is underway with the IHE Patient Care Coordination Domain for medical summaries used in referrals and discharge summaries. See www.ihe.net.

M.2 Cross-Enterprise Dynamic Information Sharing

The management of dynamic information (non-document-oriented) such as allergy lists, medication lists, problem lists, etc is not addressed by XDS. However, a means to access this information in a structured form and to manage updates to such dynamic clinical information is a candidate for a specific Integration Profile.

M.3 Collaborative Workflow Process Management

There is a wide array of shared care delivery collaborative processes such as the placing and tracking of orders (e.g. drug prescriptions, radiology orders, etc.) for which XDS provides only a partial solution (the creation of the patient record with the resulting persistent artifacts). XDS offers a critical infrastructure for ePrescribing and eReferral in that it can ensure that the various providers share access to orders, prescriptions, dispensations, and results. The means to interoperate on the command/control part of these collaborative workflow processes is a candidate for specific Integration Profiles in the future.

M.4 Security and Privacy Management

The operation of any XDS Affinity Domain will require that a proper security model be put in place. It is expected that a range of security models should be possible. Although the XDS Integration Profile is not intended to include nor require any specific security model, it is expected that XDS implementers will group XDS Actors with actors from the IHE Audit Trail and Node Authentication and will need an Access Control capability that operates in such a cross-enterprise environment. Specific IHE Integration Profiles complementary to XDS are available (e.g. Cross-Enterprise User Authentication, Document Digital Signature, etc).

M.5 Federation of XDS Affinity Domains

XDS is an effective means to establish XDS Affinity Domains that include care delivery organizations at any level, local, regional or national. However, the establishment of independent but consistently XDS Affinity Domains will call for their federation, as patients expect their records to follow them as they move from region to region, or country to country. IHE foresees a need for transferring information from one XDS Affinity Domain to another, or to allow access from one XDS Affinity Domain to documents managed in other XDS Affinity Domains. XDS has been designed with this extension in mind. An XDS Domains Federation Integration Profile that complements XDS may be anticipated in the future.

Appendix N: Intentionally Left Blank

Appendix O: Intentionally Left Blank

Appendix P: Privacy Accessss Policies (Informative)

This Appendix provides information about when consent could be automated and consequently when BPPC could be used. Privacy consent can be summarized as: "I agree on my personal data being disclosed to some one under specific conditions".

Conditions are based on various factor(s) for example:

- type of person the data is disclosed to;
- type of data disclosed;
- type of access (normal access, emergency access...);
- security level in which the disclosure takes place (weak authentication vs. strong authentication);
- type of purpose for which the data is disclosed;
- timeframe (period of validity of the consent, window of disclosure...);

BPPC could be used when conditions can be described with a limited number of factors and when the factors can be defined and be easily interpreted by a Document Consumer implementing the Basic Patient Privacy Enforcement Option.

The XDS Affinity Domain Privacy Consent Policies could result in various actions, for example:

- limitation of the display of the existence of specific documents to the users of a Document Consumer
- limitation of the access to specific documents by the users of a Document Consumer
- display of a warning note (either concerning this access or to inform that further disclosure is not allowed, limited to some defined population, needed further consent...);
- collection of new consent (oral consent, patient authentication, electronically signed consent, paper consent...);

P.1 Consents in a sensitivity labeled and role based access control environment

One possible implementation may have a collection of policies and sensitivity markers that would form an access control matrix. An example simple access control matrix is shown in the table below.

Sensitivity / Functional Role	Billing Information	Administrative Information	Dietary Restrictions	General Clinical Information	Sensitive Clinical Information	Research Information	Mediated by Direct Care Provider
Administrative Staff	X	X					
Dietary Staff		X	X				
General Care Provider		X	X	X			
Direct Care Provider		X	X	X	X		X
Emergency Care Provider		X	X	X	X		X
Researcher						X	
Patient or Legal Representative	X	X	X	X	X		

Table P-1 Sample Access Control Policies

Each slice through the matrix vertically (by sensitivity marker), results in a single Patient Privacy Consent Policy. This vocabulary must then be configured in the XDS Affinity Domain. Thus configuring each application in the XDS Affinity Domain to recognize for each Patient Privacy Consent Policy identified, what types of accesses are allowed. Using the example above, the privacy consent policies might look like.

Privacy Consent Policy	Description
Billing Information	May be accessed by administrative staff and the patient or their legal representative.
Administrative Information	May be accessed by administrative or dietary staff or general, direct or emergency care providers, the patient or their legal representative.
Dietary Restrictions	May be accessed by dietary staff, general, direct or emergency care providers, the patient or their legal representative.
General Clinical Information	May be accessed by general, direct or emergency care providers, the patient or their legal representative.
Sensitive Information	May be accessed by direct or emergency care providers, the patient or their legal representative.

Research Information	May be accessed by researchers.
Mediated by Direct Care Provider	May be accessed by direct or emergency care providers.

Table P-2 Privacy Consent Policies When Expressed by Document Sensitivity

Other divisions of the access control matrix are possible, so long as a Privacy Consent Policy covers each cell granting access in the matrix.

The following list of references is provided as good references to understand the terms and concepts presented here. These references are not required by this profile.

- ISO/TS 21298 "Health informatics – Functional and structural roles".
- ISO/TS 22600 "Health Informatics – Privilege Management and Access Controls".
- CEN prEN 13606-4 "Health informatics — Electronic health record communication — Part 4: Security requirements and distribution rules"

P.3 Possible checklist for implementations

General (before anything else)

- Granularity of confidentiality implementation:
  - Granularity of document: all documents, document type, each document.
  - Granularity of user: all users, user type, each type.
- Depth of confidentiality implementation:
  - Is the existence (metadata) about a document that can't be read by the user shown in a list of available documents for this patient?
  - Is the user informed there are / might be not shown documents and how much?
  - Is there the possibility to manage different depth of confidentiality depending on users or document type?
- How to identify users, documents and policy?
- Does confidentiality management spread through further use (once the document is downloaded by a user)

While implementing

- Definition of default codes depending on site / hardware, document type, author, patient...
- Implementing options:
  - possibility of a list to choose from and how the list is constituted (out of all the possible value, out of the value acknowledged by patient...)
  - possibility to change default codes prior to publication
  - possibility to use different format depending on the confidentiality policy (only non-downloadable image, pdf, word...)
- Later modification of policy (possible directly when requesting a document or have to be validated before)

Prior to publication

- What elements should be checked before publication:
  - existence of a policy
  - existence of the policy used

- existence of a consent for that policy
- What additional information should be given (general consent policy, patient's specific consent policy...?)

Prior to allowing access to a document

- What elements should be checked before publication:
  - accessing user role
  - existence of the policy used vs. accessing user
- Specific accesses and impact on confidentiality policy:
  - emergency (specific policy, short cut of confidentiality policy...)
  - break glass
- What additional information should be given (general consent policy, patient' specific consent policy...)

P.4 Potential obligations

Possible things that the BPPC policies might include are not fully known at this time. The following is a list that has been discovered through use by researchers, health information exchanges, and vendors. The following are some thoughts of things that might be orchestrated by BPPC Policies.

General

1. Is the existence (metadata) about a document that can't be read by the user shown in a list of available documents for this patient
2. Map local role codes into some Affinity Domain defined role codes

Prior to implementation

3. the specific Document Source is configured with one site specific “normal” code to publish all of that Document Source documents against. For example an automatic blood-pressure device being used by one specific patient.
4. prompt user for the code to apply to the document (drop-down-list)
5. document-type based codes

Prior to publication

6. validate that the code to be published against has been acknowledged
7. support for a XDS Affinity Domain confidentialityCode that indicates that the patient has acknowledged the Patient Privacy Consent Policy that forbids the publication and/or use of documents in the XDS Affinity Domain (aka Opt-Out).

Prior to allowing access to a document

8. should documents with unrecognized codes be shown?
9. prompt the user with some site defined text "do you really want to do this?"
10. allow the user to review the base consent policy
11. allow the user to review the patient's specific Privacy Consent Acknowledgement Documents

12. allow the user to override a consent block (break-glass)
13. require that a new consent be acquired from the patient before using the documents in the XDS Affinity Domain
14. support for a XDS Affinity Domain confidentialityCode that indicates that the patient has acknowledged the Patient Privacy Consent Policy that forbids the publication and/or use of documents in the XDS Affinity Domain (aka Opt-Out).
15. validate that the code on the document has been acknowledged
16. confidentialityCode that would indicate that the Document can only be viewed, it cannot be incorporated or copied.
17. use of this document shall result in an ATNA emergency access audit event

P.5 Dynamic Use Models

It has also been suggested that documents should simply be published with the expected codes, and that only on use of a document that ALL current Patient Privacy Consent Acknowledgements are evaluated against with the code on the document. In this way revocation is more dynamic.

GLOSSARY

Actor: An entity within a use case diagram that can perform an action within a use case diagram. Possible actions are creation or consumption of a message

ADT: Admit, Discharge & Transfer.

Care Delivery Organization: A Care Delivery Organization refers to a broad variety of healthcare facilities: private practice, nursing home, ambulatory clinic, acute care in-patient facility, hospitals etc.

CCOW: ANSI certified technology neutral specification for the Health Level Seven Context Management Architecture (CMA). This architecture enables multiple applications to be automatically coordinated and synchronized in clinically meaningful ways at the point of use. The architecture specified in this document establishes the basis for bringing interoperability among healthcare applications to point-of-use devices, such as a personal computer that serves as a clinical desktop

Context Management Registry: An HTTP technology specific service defined by the HL7 Context Management "CCOW" Standard to locate an instance of a context manager servicing a specific desktop.

Context Session: A collection of participant applications that are sharing context on one or more subjects.

CDA: Clinical Document Architecture (specified by HL7).

CT: Consistent Time Integration Profile.

XDS Affinity Domain: A group of healthcare enterprises that have agreed to work together using a common set of policies and which share a common infrastructure of repositories and a registry.

Directory: A book containing the names and residences of the inhabitants of any place, or of classes of them; an address book; as, a business directory.

EHR-CR: An EHR-CR or Care-delivery Record abstracts the patient information managed by the IT system or set of systems of a Care Delivery Organization, which may support a broad variety of healthcare facilities: private practice, nursing home, ambulatory clinic, acute care in-patient facility, etc.

EHR-LR: The documents shared by the EHR-CR and tracked by the Registry form a Longitudinal Record for the patients that received care among the EHR-CRs of the XDS Affinity Domain. This is known as the EHR-LR.

eMPI: Enterprise Master Patient Index.

Encounter : An interaction between a patient and care provider(s) for the purpose of providing healthcare-related service(s). Healthcare services include health assessment.

Examples: outpatient visit to multiple departments, home health support (including physical therapy), inpatient hospital stay, emergency room visit, field visit (e.g., traffic accident), office visit, occupational therapy, telephone call.

EUA: Enterprise User Authentication Integration Profile.

Expected Actions: Actions which should occur as the result of a trigger event.

Globally Unique Identifier (GUID): An identifier of an entity, such as persistent document, that has been generated by an algorithm guaranteeing its global uniqueness.

HIMSS: Healthcare Information and Management Systems Society.

HIS: Hospital Information System.

IETF: Internet Engineering Task Force

IHE: Integrating the Healthcare Enterprise.

inetOrgPerson: The inetOrgPerson [RFC 2798] object class is a general purpose object class that holds attributes about people. The attributes it holds were chosen to accommodate information requirements found in typical Internet and Intranet directory service deployments. The inetOrgPerson object class is designed to be used within directory services based on the LDAP v3 [RFC 2251] and the X.500 family of protocols, and it should be useful in other contexts as well.

Interaction Diagram: A diagram that depicts data flow and sequencing of events.

IT: Information Technology.

JPEG: – Joint Photographic Experts Group.

KDC: Key Distribution Center (the Kerberos server that issues Ticket Granting Tickets and service tickets. See RFC1510).

LDAP: Lightweight Directory Access Protocol is designed to provide access to directories supporting the X.500 models, while not incurring the resource requirements of the X.500 Directory Access Protocol (DAP). This protocol is specifically targeted at management applications and browser applications that provide read/write interactive access to directories. When used with a directory supporting the X.500 protocols, it is intended to be a complement to the X.500 DAP.

Local Authentication: In the ATNA profile the term “local authentication” means that the user identification, authentication, and authorization method is chosen by the local system administration and does not necessarily comply with any IHE profile. It may be a local username password system, a secure token system, or any other system that is considered acceptable by the local security administration.

Movement: An event describing a change of the situation of the patient in the context of the encounter. This concept encompasses changes such as transfers of patient location, change of patient class, new attending doctor, new consulting doctor, new encounter starting, encounter closing, etc. The concept of Movement is a superset of the concept of “Transfer”.

MPI: Master Patient Index.

MRN: Medicare Record Number.

NEMA: National Electrical Manufacturers Association.

NTP: Network Time Protocol. This is the standard Internet protocol for synchronizing computer clocks. The web site http://www.ntp.org provides extensive background documentation at the introductory and expert level on how to synchronize computers.

OID: Object Identifier. (See also 'Globally Unique Identifier’).

PACS: Picture Archive and Communication System.

Patient: (When used in the context of ATNA) RFC-3881 defines the means of identifying the person who is a patient. The patient information in audit event records corresponds to the information available to identify a patient at the time the audit record was generated, and does not reflect later updates (e.g. patient reconciliation).

PatientID: (When used in the context of ATNA) A free text that holds the system-internal patient identifier being unique within that system domain. The patient identifier domain is that assigned to the system that generated the audit event record. The patient information in audit event records corresponds to the information available to identify a patient at the time the audit record was generated, and does not reflect later updates (e.g. patient reconciliation).

Patient Identifier Cross-reference Domain: Consists of a set of Patient Identifier Domains known and managed by a Patient Identifier Cross-reference Manager Actor. The Patient Identifier Cross-reference Manager Actor is responsible for providing lists of "alias" identifiers from different Patient Identifier Domains.

Patient Identifier Domain: A single system or a set of interconnected systems that all share a common identification scheme for patients. Such a scheme includes: (1) a single identifier-issuing authority, (2) an assignment process of an identifier to a patient, (3) a permanent record of issued patient identifiers with associated traits, and (4) a maintenance process over time. The goal of Patient Identification is to reduce errors.

Patient Mapping Agent: The CCOW defined component that provides for the mapping of patient identifiers across disparate patient identity domains.

Patient Privacy Consent Acknowledgement Document: A document that follows the BPPC Content Profile and captures the act of the patient acknowledging a specific XDS Affinity Domain defined Privacy Consent Policy.

Patient Privacy Consent Policy: A Patient Privacy Consent Policy further explains appropriate use of the XDS Affinity Domain in a way that provides choices to the patient. The BPPC profile places no requirements on the content of these policies nor the method used to develop these policies (See Appendix P for some guidance on developing these policies). A Privacy Consent Policy will identify who has access to information, and what information is governed by the policy (e.g., under what conditions will a document be marked as containing that type of information).

Patient Privacy Consent Policy Identifier: An Affinity Domain assigned identifier (OID) that uniquely identifies the Affinity Domain: Patient Privacy Consent Policy. There is one unique identifier (OID) for each Privacy Consent Policy within the Affinity Domain.

Patient Subject: The PSA defined subject that supports sharing the currently selected patient identifier amongst disparate applications running on the desktop.

PDF: Portable Document Format.

Personnel White Pages: Information on human workforce members within the authority of the PWP directory. This information has broad use among many clinical and non-clinical applications across the healthcare enterprise. The information can be used to enhance the clinical workflow (contact information), enhance the user interface (user friendly names and titles), and ensure identity.

PIX: Patient Identifier Cross-referencing Integration Profile.

PMA: Patient Mapping Agent component as defined by CCOW.

Principal: An end user, an application, a machine, or any other type of entity that may act as a requester in a transaction. A principal is typically represented in a transaction with a digital identity and the principal may have multiple valid digital identities to use with different transactions

Process Flow Diagram: A graphical illustration of the flow of processes and interactions among the actors involved in a particular example.

PSA: Patient-Synchronized Applications Integration Profile.

RID: Retrieve Information for Display Integration Profile.

RIS: Radiology Information System.

Role: The actions of an actor in a use case.

RSNA: Radiological Society of North America.

Scope: A brief description of the transaction.

Secure Domain: A network, hardware systems, secure nodes, and physical environment for which a single set of security policies is defined and enforced for access to its addressable objects.

Secure Node: A network-addressable system that conforms to a secure domain's access policies and management. A secure node often supports IHE actors.

SNTP: Simple Network Time Protocol. This is a reduced accuracy version of NTP. The protocol fields are the same, but the data values and algorithms used are greatly reduced accuracy so that it can be implemented on limited capacity systems.

Submission Set: A set of XDS documents registered together to a Document Repository concerning information related to one care event of a single patient, provided by an EHR system.

SUID: The Study Instance UID from a DICOM SOP instance, or collection of SOP instances.

TGT: Ticket Granting Ticket. The initial credentials that verify that the user has been authenticated. It is used to avoid repeated user authentication events and as a token to request access to services.

Trigger Event: An event such as the reception of a message or completion of a process, which causes another action to occur.

UID: Unique Identifier (See also Globally Unique Identifier).

Universal ID: Unique identifier over time within the UID type. Each UID must belong to one of specifically enumerated species. Universal ID must follow syntactic rules of its scheme.

Use Case: A graphical depiction of the actors and operation of a system.

Username: A sequence of characters, different from a password, that is used as identification and is required when logging on to a multi-user computer system, LAN, bulletin board system, or online service. Also called user ID, or uid.

User Assertion: A set of claims about an authenticated principal (user, application, system...) that is issued by an identity provider

User Subject: The PSA defined subject that supports sharing the user identity of the currently logged in to the applications on the desktop.

UTC: Universal Coordinated Time. This is the replacement for GMT. It defines a reference time base that is internationally recognized and supported.

Wet Signature: Ink on paper signature.

X-Assertion Provider: This is a SAML Identity Provider (IDP) or WS-Trust Security Token Service (STS), and is not further specified by IHE.

XDS Affinity Domain Policy: XDS Affinity Domain Policy that clearly defines the appropriate uses of the XDS Affinity Domain. Within this policy is a defined set of acceptable use Privacy Consent Policies that are published and understood.

XDS Document: An XDS Document is the smallest unit of information that may be provided to a Document Repository and registered in a Document Registry. An XDS Document may contain simple text, formatted text (e.g. HL7 CDA Release 1), images (e.g. DICOM) or structured and vocabulary coded clinical information (e.g. CDA Release 2, CCR), or may be made up of a mixture of the above types of content.

XDS Folder: An XDS Folder allows document sources to group the documents they submit with other related documents. What constitutes a Folder and the vocabulary associated with the specific Folders used by an EHR-CR is decided by an agreement between the care delivery organization members of an XDS Affinity Domain.

XUA: Cross-Enterprise User Assertion Integration Profile

