



Privacy Impact Assessment for the VA IT System called:

EHRM Forward Deployed Solution Set (FDSS) Non-Digital Imaging and Communications in Medicine (Non-DICOM) Imaging Extraction

VA Corporate Office (VACO)

Electronic Health Record Modernization Integration Office (EHRM-IO)

eMASS ID # 1134

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System Contacts:

System Contacts

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Abstract

The abstract provides the simplest explanation for “what does the system do?”.

The Electronic Health Record Modernization (EHRM) Forward Deployed Solution Set (FDSS) Non-Digital Imaging and Communications in Medicine (Non-DICOM) Imaging Extraction supports the overall extraction and migration of scanned images in Veterans Health Information Systems and Technology Architecture (VistA) Imaging from Department of Veterans Affairs (VA) Medical Centers (VAMC’s). Non-DICOM images extracted by the Non DICOM solution will be stored on Oracle Health Portable Encrypted Devices (CPED) to physically transport secure image data back to the Federal Enclave at Oracle Health Government Services (OHGS) data centers in Kansas City, MO (KC3), where images from CPEDs will be extracted to SAN Storage then into Ingestion server(s) for processing.

Overview

The overview is the most important section of the PIA. A thorough and clear overview gives the reader the appropriate context to understand the responses in the PIA. The overview should contain the following elements:

1 General Description

- A. *What is the IT system name and the name of the program office that owns the IT system?*

The IT system name is EHRM Forward Deployed Solution Set (FDSS) Non-DICOM Imaging Extraction and it is owned by the EHRM Integration Office (EHRM-IO).

- B. *What is the business purpose of the program, IT system, or technology and how it relates to the program office and agency mission?*

EHRM FDSS Non-DICOM Imaging Extraction supports the overall extraction and migration of scanned images in VistA Imaging from VA Medical Centers (VAMC’s).

- C. *Who is the owner or control of the IT system or project?*

The system is owned/controlled by the VA EHRM-IO and operated by Oracle Health, the Prime Contractor of EHRM-IO.

2. Information Collection and Sharing

- D. *What is the expected number of individuals whose information is stored in the system and a brief description of the typical client or affected individual?*

This Enterprise transactional system will be gradually deployed in all VAMC’s and CBOC’s. However, authorized end-users of Millennium (the EHR Core system) won’t have direct access to the system. Only a handful authorized EHRM-IO personnels,

including contractor staff, can access the system to perform administrative and maintenance jobs.

E. What is a general description of the information in the IT system and the purpose for collecting this information?

EHRM FDSS Non-DICOM Imaging Extraction supports the supports the overall migration and extraction of scanned images in VistA Imaging from VAMCs to the Federal Enclave at OHGS data centers (KC3).

F. What information sharing conducted by the IT system? A general description of the modules and subsystems, where relevant, and their functions.

Vista Imaging data will be transported by means of CPEDs, from VAMC's to the Federal Enclave at KC3.

G. Is the system operated in more than one site, and if so, a description of how use of the system and PII is maintained consistently in all sites and if the same controls are used across sites?

The system has been gradually deployed in VAMC's and CBOC's across the country with the same configuration and safeguarded by the same set of security and privacy controls, in accordance with VA Directive & Handbook 6500 series.

3. Legal Authority and SORN

H. What is the citation of the legal authority to operate the IT system?

The legal authority to collect data pursuant to the Privacy Act of 1974 is stated in VA SORN 24VA10A7, Patient Medical Records-VA, published in FR 85, 62406, on October 2, 2020. The authority to operate the system is stated in Title 38 U.S.C. § 501(b) and 304.

I. If the system is in the process of being modified and a SORN exists, will the SORN require amendment or revision and approval? If the system is using cloud technology, does the SORN for the system cover cloud usage or storage?

No SORN amendment or revision is expected.

4. System Changes

J. Will the completion of this PIA will result in circumstances that require changes to business processes?

No change to existing business processes is expected as result of this PIA completion.

K. Will the completion of this PIA could potentially result in technology changes?

The completion of this PIA will not result in any technology change of the underlined system.

Section 1. Characterization of the Information

The following questions are intended to define the scope of the information requested and collected as well as the reasons for its collection as part of the program, IT system, or technology being developed.

1.1 What information is collected, used, disseminated, created, or maintained in the system?

Identify and list all Sensitive Personal Information (SPI) that is collected and stored in the system, including Individually Identifiable Information (III), Individually Identifiable Health Information (IIHI), Protected Health Information (PHI), and Privacy- Protected Information. For additional information on these information types and definitions, please see VA Directives and Handbooks in the 6500 series (<https://vaww.va.gov/vapubs/>). If the system creates information (for example, a score, analysis, or report), list the information the system is responsible for creating.

If a requesting system receives information from another system, such as a response to a background check, describe what information is returned to the requesting system.

This question is related to privacy control AP-1, Authority To Collect, and AP-2, Purpose Specification.

Please check any information listed below that your system collects, uses, disseminates, creates, or maintains. If additional SPI is collected, used, disseminated, created, or maintained, please list those in the text box below:

- | | | |
|--|--|--|
| <input checked="" type="checkbox"/> Name | <input checked="" type="checkbox"/> Health Insurance Beneficiary Numbers | <input type="checkbox"/> Integrated Control Number (ICN) |
| <input checked="" type="checkbox"/> Social Security Number | Account numbers | <input type="checkbox"/> Military History/Service Connection |
| <input checked="" type="checkbox"/> Date of Birth | <input type="checkbox"/> Certificate/License numbers ¹ | <input type="checkbox"/> Next of Kin |
| <input checked="" type="checkbox"/> Mother's Maiden Name | <input type="checkbox"/> Vehicle License Plate Number | <input checked="" type="checkbox"/> Other Data Elements (list below) |
| <input checked="" type="checkbox"/> Personal Mailing Address | <input type="checkbox"/> Internet Protocol (IP) Address Numbers | |
| <input checked="" type="checkbox"/> Personal Phone Number(s) | <input checked="" type="checkbox"/> Medications | |
| <input checked="" type="checkbox"/> Personal Fax Number | <input checked="" type="checkbox"/> Medical Records | |
| <input checked="" type="checkbox"/> Personal Email Address | <input checked="" type="checkbox"/> Race/Ethnicity | |
| <input checked="" type="checkbox"/> Emergency Contact Information (Name, Phone Number, etc. of a different individual) | <input type="checkbox"/> Tax Identification Number | |
| <input type="checkbox"/> Financial Information | <input checked="" type="checkbox"/> Medical Record Number | |
| | <input checked="" type="checkbox"/> Gender | |

Other PII/PHI data elements: Electronic Data Interchange Personal Identifier (EDIPI) is used as the prime identifier and medical record number (MRN), Radiology number (RAD), Consult number (CON), Study number, Facility name, reason for image, Type of study, Scanned documents, Motion video and other non-textual data files, Non-DICOM clinical images, Benefits, X-ray technician name

PII Mapping of Components (Servers/Database)

EHRM FDSS Non-DICOM Imaging Extraction consists of one (1) key component (servers/databases/instances/applications/software/application programming interfaces (API). Each

¹ *Specify type of Certificate or License Number (e.g., Occupational, Education, Medical)

component has been analyzed to determine if any elements of that component collect PII. The type of PII collected by the system and the reasons for the collection of the PII are in the table below.

Note: Due to the PIA being a public facing document, please do not include server names in the table.

Internal Components Table

Component Name (Database, Instances, Application, Software, Application Program Interface (API) etc.) that contains PII/PHI	Does this system collect PII? (Yes/No)	Does this system store PII? (Yes/No)	Type of PII (SSN, DOB, etc.)	Reason for Collection/ Storage of PII	Safeguards
Non-DICOM Imaging Extraction application and database servers	Yes	No	Name, EDIPI, SSN, Date of birth, Mother's maiden name, Mailing address, Phone number(s), Fax number, Email address, Emergency Contact Information, Health Insurance Beneficiary Numbers/Account Numbers, Race/Ethnicity, MRN, Gender, Medication, Medical records, RAD, CON, Study number, Facility name, reason for image, Type of study, Scanned documents, Motion video and other non-textual data files, Non-DICOM clinical images, Benefits, X-ray technician name	Central archiving of medical imaging records	Transport layer security (TLS) and encrypted devices

1.2 What are the sources of the information in the system?

These questions are related to privacy controls DI-1, Data Quality, and IP-1, Consent.

1.2a List the individual, entity, or entities providing the specific information identified above. For example, is the information collected directly from the individual as part of an application for a benefit, or is it collected from other sources such as commercial data aggregators?

The source of information collected and processed by this system is VistA imaging objects gathered at each VAMC.

1.2b Describe why information from sources other than the individual is required? For example, if a program's system is using data from a commercial aggregator of information or data taken from

public Web sites, state the fact that this is where the information is coming from and then in question indicate why the system is using this source of data.

Information is not collected directly from individuals. They are VistA imaging non-DICOM objects collected from VA Medical Centers.

1.2c Does the system create information (for example, a score, analysis, or report), list the system as a source of information?

The system does not create new information.

1.3 How is the information collected?

These questions are related to privacy controls DI-1, Data Quality, and IP-1, Consent.

1.3a This question is directed at the means of collection from the sources listed in question 1.2. Information may be collected directly from an individual, received via electronic transmission from another system, or created by the system itself. Specifically, is information collected through technologies or other technologies used in the storage or transmission of information in identifiable form?

Information is not collected directly from individuals but from the local VAMC VistA Imaging system.

1.3b If the information is collected on a form and is subject to the Paperwork Reduction Act, what is the form's OMB control number and the agency form number?

Information is not collected on a form and is not subject to the Paperwork Reduction Act.

1.4 How will the information be checked for accuracy? How often will it be checked?

These questions are related to privacy controls DI-1, Data Quality, and DI-2, Data Integrity and Integrity Board.

1.4a Discuss whether and how often information stored in the system is checked for accuracy. Is information in the system checked against any other source of information (within or outside your organization) before the information is used to make decisions about an individual? For example, is there a computer matching agreement in place with another government agency? For systems that receive data from internal data sources or VA IT systems, describe the system checks to ensure that data corruption has not occurred during transmission.

The application server reads images from a file, checks the conformance of the image and creates a test report which describes in detail all detected violations of the applicable imaging standard.

1.4b Does the system check for accuracy by accessing a commercial aggregator of information, describe this process and the levels of accuracy required by the contract?

Data integrity, along with data confidentiality, is satisfied by means of deploying FIPS 140-2 encryption in the CPED.

1.5 What specific legal authorities, arrangements, and agreements defined the collection of information?

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List the full legal authority for operating the system, specifically the authority to collect the information listed in question 1.1. Provide the authorities in a manner understandable to any potential reader, i.e., do not simply provide a legal citation; use statute names or regulations in addition to citations. Legal authorities include Federal laws, regulations, statutes, and Executive Orders. This question is related to privacy control AP-1, Authority to Collect

The authority for the system to collect, use, and disseminate information about individuals that is maintained in systems of records by federal agencies, in accordance with the code of fair information practices established by the Privacy Act of 1974, as amended, Title 38 U.S.C. § 501(b) and 304, and VA System of Record Notice (SORN) 24VA10A7, Patient Medical Records-VA, published in FR 85, 62406, on October 2, 2020 (<https://www.govinfo.gov/content/pkg/FR-2020-10-02/pdf/2020-21426.pdf>). A biennial review of the SORN was conducted by the VHA Privacy Office in 2022 without any change recommended.

1.6 PRIVACY IMPACT ASSESSMENT: Characterization of the information

Consider the specific data elements collected and discuss the potential privacy risks and what steps, if any are currently being taken to mitigate those identified risks. (Work with your System ISSO to complete this section)

Consider the following Fair Information Practice Principles (FIPPs) when assessing the risk to individual privacy:

Principle of Purpose Specification: Explain how the collection ties with the purpose of the underlying mission of the organization and its enabling authority.

Principle of Minimization: Is the information directly relevant and necessary to accomplish the specific purposes of the program?

Principle of Individual Participation: Does the program, to the extent possible and practical, collect information directly from the individual?

Principle of Data Quality and Integrity: Are there policies and procedures for VA to ensure that personally identifiable information is accurate, complete, and current?

This question is related to privacy control AR-1, Governance and Privacy Program, and AR-2, Privacy Impact and Risk Assessment.

Follow the format below when entering your risk assessment:

Privacy Risk: The handling of CPED's transportation from VAMC to KC3 data center may be compromised, such as device gets lost.

Mitigation: VA EHRM-IO follows Committee for National Security System (CNSS) Instruction 4001, Controlled Cryptographic Items (CCI), section V, provisions 16 – Preparation for Shipment; and 17 – Transportation of Unkeyed CCI's in handling and shipping of CPED.

Section 2. Uses of the Information

The following questions are intended to clearly delineate the use of information and the accuracy of the data being used.

2.1 Describe how the information in the system that will be used in support of the program's business purpose.

Identify and list each use (both internal and external to VA) of the information collected or maintained. This question is related to privacy control AP-2, Purpose Specification.

PII/PHI Data Element	Internal Use	External Use
Name	Used to identify patient and matching record during migration.	Not used
Social security number (SSN)	Used to identify patient and matching record during migration.	
Electronic data interchange personal identifier (EDIPI)	Used as the prime identifier/ medical record number and is used for patient identity and record matching during migration	Not used
Date of Birth	Used to identify age and confirm patient identity.	Not used
Mother's maiden name	Patient record matching in migration	Not used
Mailing address	Patient record matching in migration	Not used
Phone number(s)	Patient record matching in migration	Not used
Fax number	Patient record matching in migration	Not used
Email address	Patient record matching in migration	Not used
Emergency contact information	Patient record matching in migration	Not used
Health insurance beneficiary numbers/Account numbers	Patient record matching in migration	Not used
Race/Ethnicity	Patient record matching in migration	Not used
Medical record number (MRN)	Patient record matching in migration	Not used
Gender identity	Patient record matching in migration	Not used
Medication	Patient record matching in migration	Not used
Medical records	Patient record matching in migration	Not used
Radiology number (RAD)	Patient record matching in migration	Not used
Consult number (CON)	Patient record matching in migration	Not used
Study number	Patient record matching in migration	Not used
Facility name	Patient record matching in migration	Not used
Non-DICOM image objects including attributes such as date taken or from what study	The attributes are used to identify and differentiate imaging objects	Not used

PII/PHI Data Element	Internal Use	External Use
Reason for image	Patient record matching in migration	Not used
Type of study	Patient record matching in migration	Not used
Scanned documents	Patient record matching in migration	Not used
Motion video and other non-textual data files	Patient record matching in migration	Not used
X-ray technician name	Patient record matching in migration	Not used

2.2 What types of tools are used to analyze data and what type of data may be produced?

These questions are related to privacy controls DI-1, Data Quality, DI-2, Data Integrity and Integrity Board, and SE-1, Inventory of Personally Identifiable Information.

2.2a Many systems sift through large amounts of information in response to a user inquiry or programmed functions. Systems may help identify areas that were previously not obvious and need additional research by agents, analysts, or other employees. Some systems perform complex analytical tasks resulting in, among other types of data, matching, relational analysis, scoring, reporting, or pattern analysis. Describe any type of analysis the system conducts and the data that is created from the analysis?

This is a transactional system without the capability to analyze data.

2.2b If the system creates or makes available new or previously unutilized information about an individual, explain what will be done with the newly derived information. Will it be placed in the individual's existing record? Will a new record be created? Will any action be taken against or for the individual identified because of the newly derived data? If a new record is created, will the newly created information be accessible to Government employees who make determinations about the individual? If so, explain fully under which circumstances and by whom that information will be used.

The system does not create new information.

2.3 How is the information in the system secured?

These questions are related to security and privacy controls SC-9, Transmission Confidentiality, and SC-28, Protection of Information at Rest.

2.3a What measures are in place to protect data in transit and at rest?

FIPS 140-2 approved cryptography is used for the CPED to protect data in transit and at rest.

2.3b If the system is collecting, processing, or retaining Social Security Numbers, are there additional protections in place to protect SSNs?

The security and privacy controls implemented for the system are determined sufficient to safeguard SSN as well as other data elements identified during the system and data security categorization process.

2.3c How is PII/PHI safeguarded in accordance with OMB Memorandum M-06-15?

The system security risk level is categorized at Moderate, in accordance with Federal Information Processing Standards (FIPS) Publication 199, and meets requirements set forth by OMB Memorandum M-06-15, Safeguarding Personally Identifiable Information. A set of more than 300 security and privacy controls have been selected and implemented commensurate to the system risk level, to include technical, operational, and administrative controls covering access controls, identification & authentication, personnel security, physical security, auditing and monitoring, incident response.

2.4 PRIVACY IMPACT ASSESSMENT: Use of the information.

Describe any types of controls that may be in place to ensure that information is handled in accordance with the uses described above. Example: Describe if training for users of the project covers how to appropriately use information. Describe the disciplinary programs or system controls (i.e. denial of access) that are in place if an individual is inappropriately using the information.

Consider the following FIPPs below to assist in providing a response:

Principle of Transparency: *Is the PIA and SORN, if applicable, clear about the uses of the information?*

Principle of Use Limitation: *Is the use of information contained in the system relevant to the mission of the project?*

This question is related to privacy control AR-4, Privacy Monitoring and Auditing, AR-5, Privacy Awareness and Training, and SE-2, Privacy Incident response.

2.4a How is access to the PII determined?

End-users of the EHR core system do not have direct access to this transactional system.

2.4b Are criteria, procedures, controls, and responsibilities regarding access documented?

User provisioning, identification and authentication processes are documented in the account management standard operating procedure, which covers criteria, procedures, roles and responsibilities, and applicable security and privacy controls in accordance with applicable Federal and VA policies, procedures, and standards.

2.4c Does access require manager approval?

Authorized EHRM-IO personnel with admin access to the system do need direct supervisor/manager approval.

2.4d Is access to the PII being monitored, tracked, or recorded?

Network, system, media transport auditing, monitoring controls are in place, in accordance with applicable Federal and VA information security & privacy policies, procedures, and standards.

2.4e Who is responsible for assuring safeguards for the PII?

The System Owner is ultimately responsible for assuring safeguards for the PII collected and processed by the system.

Section 3. Retention of Information

The following questions are intended to outline how long information will be retained after the initial collection.

3.1 What information is retained?

*Identify and list all information collected from question 1.1 that is **retained** by the system. This question is related to privacy controls DM-1, Minimization of Personally Identifiable Information, and DM-2, Data Retention and Disposal*

No PII is retained by the local servers. The CPEDs retain PII/PHI for the duration of transportation up to the point they arrive at their destination, which is the KC3 data center where VistA imaging objects will be loaded to SAN storage and extracted.

3.2 How long is information retained?

*In some cases, VA may choose to retain files in active status and archive them after a certain period of time. State active file retention periods, as well as archived records, in number of years, for the information and record types. For example, financial data held within your system may have a different retention period than medical records or education records held within your system, please be sure to list each of these retention periods. **The VA records officer should be consulted early in the development process to ensure that appropriate retention and destruction schedules are implemented.** If the system is using cloud technology, will it be following the NARA approved retention length and schedule? This question is related to privacy control DM-2, Data Retention and Disposal.*

VistA imaging objects are retained in the CPED's then being destroyed in accordance with disposition instructions set forth in the VA Record Control Schedule 10-1, item number 6000.2.(2) – Interim Electronic Source Information.

3.3 Has the retention schedule been approved by the VA records office and the National Archives and Records Administration (NARA)?

An approved records schedule must be obtained for any IT system that allows the retrieval of a record via a personal identifier. The VA records officer will assist in providing a proposed schedule. The schedule must be formally offered to NARA for official approval. Once NARA approves the proposed schedule, the VA records officer will notify the system owner. Please work with the system Privacy Officer and VA Records Officer to answer these questions. This question is related to privacy control DM-2, Data Retention and Disposal.

3.3a Are all records stored within the system of record indicated on an approved disposition authority?

Yes, records stored within the system are indicated on NARA-approved VA Record Control Schedule RCS 10-1 at <https://www.va.gov/vhapublications/RCS10/rcs10-1.pdf> .

3.3b Please indicate each records retention schedule, series, and disposition authority?

VA RCS10-1 Item number 6000.2.(2), N1-15-02-3, item 2

(<https://www.va.gov/vhapublications/RCS10/rcs10-1.pdf>) governs the retention of system-related information.

3.4 What are the procedures for the elimination or transfer of SPI?

Explain how records are destroyed, eliminated or transferred to NARA at the end of their mandatory retention period. Please give the details of the process. For example, are paper records shredded on site, or by a shredding company and accompanied by a certificate of destruction, etc.? This question is related to privacy control DM-2, Data Retention and Disposal.

In accordance with the disposition instructions set forth in the VA Record Control Schedule 10-1, item number 6000.2.(2) N1-15-02 -3, item 2–

(<https://www.va.gov/vhapublications/RCS10/rcs10-1.pdf>)

Interim Electronic Source Information, destruction of interim version of information is not to occur until it has been determined that the migrated information represents an exact duplicate of the previous version of the migrated information.

3.5 Does the system, where feasible, use techniques to minimize the risk to privacy by using PII for research, testing, or training?

Organizations often use PII for testing new applications or information systems prior to deployment. Organizations also use PII for research purposes and for training. These uses of PII increase the risks associated with the unauthorized disclosure or misuse of the information. Please explain what controls have been implemented to protect PII used for testing, training and research. This question is related to privacy control DM-3, Minimization of PII Used in Testing, Training and Research.

The system does not use PII for research, testing, or training.

3.6 PRIVACY IMPACT ASSESSMENT: Retention of information

Discuss the risks associated with the length of time data is retained and what steps, if any, are currently being taken to mitigate those identified risks. (Work with your System ISSO to complete all Privacy Risk questions inside the document this section).

While we understand that establishing retention periods for records is a formal process, there are policy considerations behind how long a project keeps information. The longer a project retains information, the longer it needs to secure the information and assure its accuracy and integrity. The proposed schedule should match the requirements of the Privacy Act to keep the minimum amount of PII for the minimum amount of time, while meeting the Federal Records Act. The schedule should align with the stated purpose and mission of the system.

Consider the following FIPPs below to assist in providing a response:

Principle of Minimization: *Does the project retain only the information necessary for its purpose? Is the PII retained only for as long as necessary and relevant to fulfill the specified purposes?*

Principle of Data Quality and Integrity: Has the PIA described policies and procedures for how PII that is no longer relevant and necessary is purged?

This question is related to privacy controls DM-1, Minimization of Personally Identifiable Information, and DM-2, Data Retention and Disposal.

Privacy Risk: A risk may arise when imaging objects are deleted from the CPED too early or not completely loaded onto the SAN storage.

Mitigation: The risk of incomplete data loading can be mitigated by means of validation and/or record check sum to ensure all objects are accounted for.

Section 4. Internal Sharing/Receiving/Transmitting and Disclosure

The following questions are intended to define the scope of information sharing/receiving/transmitting within VA.

4.1 With which internal organizations is information shared/received/transmitted? What information is shared/received/transmitted, and for what purpose? How is the information transmitted?

Identify and list the names of any program offices, contractor-supported IT systems, and any other organization or IT system within VA with which information is shared.

State the purpose for the internal sharing. If you have specific authority to share the information, provide a citation to the authority.

For each interface with a system outside your program office, state what specific data elements (PII/PHI) are shared with the specific program office, contractor-supported IT system, and any other organization or IT system within VA.

Describe how the information is transmitted. For example, is the information transmitted electronically, by paper, or by some other means? Is the information shared in bulk, on a case-by-case basis, or does the sharing partner have direct access to the information?

This question is related to privacy controls AP-2, Purpose Specification, AR-3, Privacy Requirements for Contractors and Service Providers, AR-8, Accounting of Disclosures, TR-1, Privacy Notice, and UL-1, Internal Use.

<i>List the Program Office or IT System information is shared/received with</i>	<i>List the purpose of the information being shared /received with the specified program office or IT system</i>	<i>List the specific PII/PHI data elements that are processed (shared/received/transmitted) with the Program Office or IT system</i>	<i>Describe the method of transmittal</i>
OI&T/VistA Imaging	Healthcare Treatment, continuity of care	Name, EDIPI, SSN, Date of birth, Mother’s maiden name, Mailing address, Phone number(s), Fax number, Email address, Emergency Contact Information, Health Insurance Beneficiary Numbers/Account Numbers, Race/Ethnicity, MRN, Gender, Medication, Medical records, RAD, CON, Study number, Facility name, reason for image, Type of study, Scanned documents, Motion video and other non-textual data files, Non-DICOM clinical images, Benefits, X-ray technician name	System-to-system electronic transmission using TLS

4.2 PRIVACY IMPACT ASSESSMENT: Internal sharing and disclosure

Discuss the privacy risks associated with the sharing of information within the Department and what steps, if any, are currently being taken to mitigate those identified risks. (Work with your System ISSO to complete all Privacy Risk questions inside the document this section).

This question is related to privacy control UL-1, Internal Use.

Privacy Risk: The extraction of VistA imaging objects from VistA Imaging Exchange to CPED may be compromised.

Mitigation: To mitigate the risk of internal sharing and disclosure and to ensure data integrity as well as data confidentiality, EHRM-IO follows Committee for National Security System (CNSS) Instruction 4001, Controlled Cryptographic Items (CCI), section V, provisions 16 – Preparation for Shipment; and 17 – Transportation of Unkeyed CCI’s in handling and shipping of CPED.

Section 5. External Sharing/Receiving and Disclosure

The following questions are intended to define the content, scope, and authority for information sharing external to VA, which includes Federal, State, and local governments, and the private sector.

5.1 With which external organizations (outside VA) is information shared/received? What information is shared/received, and for what purpose? How is the information transmitted and what measures are taken to ensure it is secure?

Is the sharing of information outside the agency compatible with the original collection? If so, is it covered by an appropriate routine use in a SORN? If not, please describe under what legal mechanism the IT system is allowed to share the information in identifiable form or personally identifiable information outside of VA.

Identify and list the names of any Federal, State, or local government agency or private sector organization with which information is shared.

For each interface with a system outside VA, state what specific data elements (PII/PHI) are shared with each specific partner.

What legal mechanisms, authoritative agreements, documentation, or policies are in place detailing the extent of the sharing and the duties of each party? For example, is the sharing of data compatible with your SORN? Then list the SORN and the applicable routine use from the SORN. Is there a Memorandum of Understanding (MOU), Computer Matching Agreement (CMA), or law that mandates the sharing of this information?

Describe how the information is transmitted to entities external to VA and what security measures have been taken to protect it during transmission.

This question is related to privacy control UL-2, Information Sharing with Third Parties

Data Shared with External Organizations

<i>List External Program Office or IT System information is shared/received with</i>	<i>List the purpose of information being shared / received / transmitted with the specified program office or IT system</i>	<i>List the specific PII/PHI data elements that are processed (shared/received/transmitted) with the Program or IT system</i>	<i>List the legal authority, binding agreement, SORN routine use, etc. that permit external sharing (can be more than one)</i>	<i>List the method of transmission and the measures in place to secure data</i>
Defense Health Agency (DHA) – Federal EHR System	Treatment, continuity of care	Name, EDIPI, SSN, Date of birth, Mother’s maiden name, Mailing address, Phone number(s), Fax number, Email address, Emergency Contact Information, Health Insurance Beneficiary Numbers/Account Numbers, Race/Ethnicity, MRN, Gender, Medication, Medical records, RAD, CON, Study number, Facility name, reason for image, Type of study, Scanned	MOU between DoD and VA for Sharing of Personal Information, June 18, 2024	Cerner portable encrypted device(s) (CPED’s)

		documents, Motion video and other non-textual data files, Non-DICOM clinical images, Benefits, X-ray technician name		
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5.2 PRIVACY IMPACT ASSESSMENT: External sharing and disclosure

Discuss the privacy risks associated with the sharing of information outside the Department and what steps, if any, are currently being taken to mitigate those identified risks.

If no External Sharing listed on the table above, (State there is no external sharing in both the risk and mitigation fields).

Discuss whether access controls have been implemented and whether audit logs are regularly reviewed to ensure appropriate sharing outside of the Department. For example, is there a Memorandum of Understanding (MOU), contract, or agreement in place with outside agencies or foreign governments.

Discuss how the sharing of information outside of the Department is compatible with the stated purpose and use of the original collection.

This question is related to privacy control AR-2, Privacy Impact and Risk Assessment, AR-3, Privacy Requirements for Contractors and Service Providers, and AR-4, Privacy Monitoring and Auditing

Privacy Risk: VA patient data is now collected and retained in a shared database as part of the Federal EHR may expose to certain privacy/security risks such as unauthorized access or being used for purposes other than the stated purpose and use of the original collection.

Mitigation: Beside the 2024 MOU signed between the then-Secretaries of DoD and VA, the two agencies have entered into several inter-agency MOA, MOU/ISA, in line with the RMF and applicable OMB Memoranda, CNSSI, DoD and VA policies and procedures to ensure data safeguarding and information privacy controls are implemented as having designed to prevent and/or detect violation or compromise situations, maintaining an acceptable risk level for the operating systems, both in Prod and Pre-Prod environments.

Section 6. Notice

The following questions are directed at providing notice to the individual of the scope of information collected, the right to consent to uses of the information, and the right to decline to provide information.

6.1 Was notice provided to the individual before collection of the information? If yes, please provide a copy of the notice as an Appendix-A 6.1 on the last page of the document. Also provide notice given to individuals by the source system (A notice may include a posted privacy policy, a Privacy Act notice on forms, or a system of records notice published in the Federal Register.) If notice was not provided, why not?

These questions are related to privacy control TR-1, Privacy Notice, and TR-2, System of Records Notices and Privacy Act Statements, and TR-3, Dissemination of Privacy Program Information.

6.1a This question is directed at the notice provided before collection of the information. This refers to whether the person is aware that his or her information is going to be collected. A notice may include a posted privacy policy, a Privacy Act statement on forms, or a SORN published in the Federal Register, Notice of Privacy Practice provided to individuals for VHA systems. If notice was provided in the Federal Register, provide the citation.

Same privacy control sets applicable to the source or parent system will be used for this “child” system. With reference to the “Notice” requirements, beside the SORN publication in the Federal Register in October 2020 as having mentioned in 1.5, the current publication of the VHA Notice of Privacy Practices (NOPP) can be found in the VHA webpage, <http://www.va.gov/health/>, under the “Resources” section. A copy of the NOPP is provided to the Veteran upon enrollment and a revised/latest NOPP mailed to eligible veterans every three years by the VHA. A copy of the NOPP must be provided to non-Veteran/humanitarian patients in person when they present for services.

6.1b If notice was not provided, explain why. If it was provided, attach a copy of the current notice.

A notice specific to the FDSS Non-DICOM system is not provided as the initial collection of data occurs under the VistA legacy and EHR Core systems and therefore this requirement has been met by those systems.

6.1c Describe how the notice provided for the collection of information is adequate to inform those affected by the system that their information has been collected and is being used appropriately. Provide information on any notice provided on forms or on Web sites associated with the collection.

A notice specific to this system is not provided as the collection of data occurs under the EHR Core system and therefore this requirement has been met by the VistA legacy and EHR Core system. Accordingly, the Notice of Privacy Practice (NOPP) is a document which explains the collection and use of protected information to individuals applying for VHA benefits. The NOPP (Appendix A) is provided when the Veteran enrolls or when updates are made to the NOPP, copies are mailed to all VHA beneficiaries (every 3 years). Employees and contractors are required to review, sign and abide by the National Rules of Behavior on an annual basis, that outlines the requirements and expectations for appropriate use of Veteran PHI/PII maintained in VA systems. In addition to NOPP distributions are the SORN publications in the Federal Register in October 2020 as mentioned in 1.5 above.

6.2 Do individuals have the opportunity and right to decline to provide information? If so, is a penalty or denial of service attached?

This question is directed at whether the person from or about whom information is collected can decline to provide the information and if so, whether a penalty or denial of service is attached. This question is related to privacy control IP-1, Consent, IP-2, Individual Access, and IP-3, Redress.

Yes, as outlined in the NOPP for the EHR Core system. Specifically, individuals do have an opportunity to decline to provide information at any time. However, to apply for enrollment in the VA health care system, all Veterans are required to fill out VA Form 10-10EZR. The

information provided on this form will be used by VA to determine eligibility for medical benefits. The applicant is not required to disclose their financial information; however, VA is not currently enrolling new applicants who decline to provide their financial information unless they have other qualifying eligibility factors. If a financial assessment is not used to determine the applicant's eligibility for cost-free medication, travel assistance or waiver of the travel deductible, and the applicant chooses not to disclose personal financial information, the applicant will not be eligible for these benefits. More details and instruction for VA Form 10-10EZr can be found through the Resources section of the VHA webpage at [va.gov/health/](https://www.va.gov/health/) or at this link <https://www.va.gov/find-forms/about-form-10-10ezr/>.

6.3 Do individuals have the right to consent to particular uses of the information? If so, how does the individual exercise the right?

This question is directed at whether an individual may provide consent for specific uses, or the consent is given to cover all uses (current or potential) of his or her information. If specific consent is required, how would the individual consent to each use? This question is related to privacy control IP-1, Consent.

Yes, as outlined in the NOPP for the VistA legacy and EHR Core system. Specifically, Right to Request Restriction: Veterans/patients do have the right to request that VHA not use or disclose all or part of their health information to carry out treatment, payment or health care operations, or that VHA not use or disclose all or part of their health information with individuals such as their relatives or friends involved in their care, including use or disclosure for a particular purpose or to a particular person. Reference the NOPP on how to submit a request for restriction. VHA, however, as a "Covered Entity" under the law, is not required to agree to such restriction, except in the case of a disclosure restricted under 45 CFR § 164.522(a)(1) (vi). This provision applies only if the disclosure of the Veteran's or patient's health information is to a health plan for the purpose of payment or health care operations and the Veteran's health information pertains solely to a health care service or visit which is paid out of pocket in full by the Veteran/patient. However, VHA is not legally able to accept an out-of-pocket payment from a Veteran for the full cost of a health care service or visit. The Administration can only accept payment from a Veteran for co-payments. Therefore, this provision does not apply to VHA and VHA is not required or able to agree to a restriction on the disclosure of a Veteran's/patient's health information to a health plan for the purpose of receiving payment for health care services provided by VHA. Additionally, VHA is not able to restrict access to the patient health information by DoD providers with whom the patient has a treatment relationship. Lastly, Individuals have the right to consent to the use of their information. Individuals are directed to use the 10-5345 Release of Information (ROI) form describing what information is to be sent out and to whom it is being sent to. Patients have the right to opt-out of VA facility directories. Individuals can request further limitations on other disclosures. A veteran, guardian or court appointed Power of Attorney can submit a request to the facility Privacy Officer to obtain information. **6.4**

PRIVACY IMPACT ASSESSMENT: Notice

Describe the potential risks associated with potentially insufficient notice and what steps, if any, are currently being taken to mitigate those identified risks. (Work with your System ISSO to complete all Privacy Risk questions inside the document this section).

Consider the following FIPPs below to assist in providing a response:

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Principle of Transparency: *Has sufficient notice been provided to the individual?*

Principle of Use Limitation: *Is the information used only for the purpose for which notice was provided either directly to the individual or through a public notice? What procedures are in place to ensure that information is used only for the purpose articulated in the notice?*

This question is related to privacy control TR-1, Privacy Notice, AR-2, Privacy Impact and Risk Assessment, and UL-1, Internal Use.

Privacy Risk: An individual may not receive notice that their information is being collected, maintained, processed, or disseminated by the VA prior to providing the information.

Mitigation: This risk is mitigated by the common practice of providing the NOPP when Veterans apply for benefits and every three years thereafter to include any changes made to the notice. Additionally, NOPPs are provided to non-Veteran beneficiaries at each episode of care and periodic monitoring is performed to check that the signed NOPP acknowledgment form has been scanned into the beneficiaries' electronic health record. Additional mitigation is provided by making the System of Record Notices (SORNs) and PIA available for review online, as discussed in question 6.1 and the Overview section of this PIA. Additional mitigation is provided by making the System of Record Notices (SORNs) 24VA10A2 – Medical Patient Records, VA- (https://www.oprm.va.gov/privacy/systems_of_records.aspx) and NOPP <http://www.va.gov/health/> available for review online.

Section 7. Access, Redress, and Correction

The following questions are directed at an individual's ability to ensure the accuracy of the information collected about him or her.

7.1 What are the procedures that allow individuals to gain access to their information?

These questions are related to privacy control IP-2, Individual Access, and AR-8, Accounting of Disclosures.

7.1a Cite any procedures or regulations your program has in place that allow access to information. These procedures, at a minimum, should include the agency's FOIA/Privacy Act practices, but may also include additional access provisions. For example, if your program has a customer satisfaction unit, that information, along with phone and email contact information, should be listed in this section in addition to the agency's procedures. See 5 CFR 294 and the VA FOIA Web page at <http://www.foia.va.gov/> to obtain information about FOIA points of contact and information about agency FOIA processes.

As having stated in the VHA NOPP, Veterans/patients have the right to review and obtain a copy of their health information by means of completing VA Form 10-5345a – Individuals' Request for a Copy of their Own Health Information, to the facility Privacy Officer of the VHA facility that provided or paid for their care. Form 10-5345a can be obtained from the facility webpage or the VA online repository at the link <https://www.va.gov/find-forms/about-form-10-5345a>. Additionally, Veterans/patients can gain access to their health

record by enrolling in the VA patient portal, myHealthVet, at <https://www.myhealth.va.gov/index.html>

7.1b If the system is exempt from the access provisions of the Privacy Act, please explain the basis for the exemption or cite the source where this explanation may be found, for example, a Final Rule published in the Code of Federal Regulations (CFR)?

This system is not exempt from the access provisions of the Privacy Act.

7.1c If the system is not a Privacy Act system, please explain what procedures and regulations are in place that covers an individual gaining access to his or her information?

This is a Privacy Act system.

7.2 What are the procedures for correcting inaccurate or erroneous information?

Describe the procedures and provide contact information for the appropriate person to whom such issues should be addressed? If the correction procedures are the same as those given in question 7.1, state as much. This question is related to privacy control IP-3, Redress, and IP-4, Complaint Management.

Right to Request Amendment of Health Information: Veterans/patients have the right to request an amendment (correction) of their health information in Federal EHR records if they believe it is incomplete, inaccurate, untimely, or unrelated to their care. A request in writing must be submitted to the facility Privacy Officer, specifying the information to be corrected, including a reason to support the request for amendment. A decision to approve or deny is made by the practitioner who entered the data and relayed to the Veteran in writing by the facility Privacy Officer. Appeal rights are provided if a request is denied. The goal is to complete any evaluation and determination within 30 days. A request for amendment of information contained in a system of records must be delivered to the System Manager, or designee, for the concerned system of records, and the facility Privacy Officer, or designee, to be date stamped; and is filed appropriately. In reviewing requests to amend or correct records, the System Manager must be guided by the criteria set forth in VA regulation 38 CFR 1.579. That is, VA must maintain in its records only such information about an individual that is accurate, complete, timely, relevant, and necessary. Lastly, individuals have the right to review and change their contact or demographic information at time of appointment or upon arrival to the VA facility and/or submit a change of address request form to the facility business office for processing.

7.3 How are individuals notified of the procedures for correcting their information?

How are individuals made aware of the procedures for correcting his or her information? This may be through notice at collection or other similar means. This question is meant to address the risk that even if procedures exist to correct information, if an individual is not made fully aware of the existence of those procedures, then the benefits of the procedures are significantly weakened. This question is related to privacy control IP-3, Redress, and IP-4, Complaint Management.

The NOPP, outlining the procedure for Veterans/patients request amendment (correction) of their health information, is provided to the Veteran/patient at the time their information being collected during enrollment and every three years thereafter. If they enroll in the patient portal, a digital version of the NOPP is also available for their awareness. Veterans/patients are expected to review and understand the said procedures as well as the NOPP in its completeness, so that they can properly exercise their rights. Particularly, the procedures also address the situation when a request for amendment is denied - Veterans/patients will be notified of such decision in writing and given information about their right to appeal the decision. In response, the Veterans/patients may do any of the following: file an appeal, file a “Statement of Disagreement” which will be included in their health record, or ask that their initial request for amendment accompany all future disclosures of the disputed health information. Reference the VHA NOPP, which can be found in the Resources section of the VHA webpage (<https://www.va.gov/health/>). Publications of the SORNs referenced in 1.5 are also a means of notification. Lastly, individuals are provided written notice of the amendment process in the written amendment acknowledgement and response letters.

7.4 If no formal redress is provided, what alternatives are available to the individual?

*Redress is the process by which an individual gains access to his or her records and seeks corrections or amendments to those records. Redress may be provided through the Privacy Act and Freedom of Information Act (FOIA), and also by other processes specific to a program, system, or group of systems. **Example: Some projects allow users to directly access and correct/update their information online. This helps ensure data accuracy.***

This question is related to privacy control IP-3, Redress, and IP-4, Complaint Management.

The processes outlined in 7.2 and 7.3 are considered formal redress process. To ensure data accuracy and maintain quality of care, patients are encouraged to actively review and verify information included in their health records.

7.5 PRIVACY IMPACT ASSESSMENT: Access, redress, and correction

*Discuss what risks there currently are related to the Department’s access, redress, and correction policies and procedures for this system and what, if any, steps have been taken to mitigate those risks. **For example, if a project does not allow individual access, the risk of inaccurate data needs to be discussed in light of the purpose of the project. For example, providing access to ongoing law enforcement activities could negatively impact the program’s effectiveness because the individuals involved might change their behavior.** (Work with your System ISSO to complete all Privacy Risk questions inside the document this section).*

Consider the following FIPPs below to assist in providing a response:

Principle of Individual Participation: Is the individual provided with the ability to find out whether a project maintains a record relating to him?

Principle of Individual Participation: If access and/or correction is denied, then is the individual provided notice as to why the denial was made and how to challenge such a denial?

Principle of Individual Participation: Is there a mechanism by which an individual is able to prevent information about him obtained for one purpose from being used for other purposes without his knowledge?

This question is related to privacy control IP-3, Redress.

Privacy Risk: Individuals whose records contain incorrect or out-of-date information may be exposed to the risk of not receiving prescription medications, notification of appointments, or test results timely. Certain incorrect information in a patient medical record could result in improper diagnosis and treatments.

Mitigation: Various accuracy checks are designed and implemented in different workflows of the DHMSM EHR Core system. VHA built-in procedure requires staff verify information in patient medical records and correct information identified as incorrect during each patient's medical appointments. Staff are informed of the importance of maintaining compliance with VA Request of Information policies and procedures and the importance of remaining alert to information correction requests.

Individual patients have the right to request an amendment (correction) to their health information in VHA records if they believe it is incomplete, inaccurate, untimely, or unrelated to your care. The individuals must submit request in writing, specify the information that they want corrected, and provide a reason to support their request for amendment. All amendment requests should be submitted to the facility Privacy Officer at the VHA health care facility that maintains the patient's information or health records. Reference "Right to Request Amendment of Health Information" under VHA Notice of Privacy Practices (NOPP) (<https://www.va.gov/health/>)

Section 8. Technical Access and Security

The following questions are intended to describe technical safeguards and security measures.

8.1 What procedures are in place to determine which users may access the system, and are they documented?

These questions are related to privacy control AR-7, Privacy-Enhanced System Design and Development.

8.1a Describe the process by which an individual receives access to the system?

The User Role Assignment Standard Operating Procedure (URA SOP), version 1.5, dated December 15, 2022, developed and managed by the National User Role Access Coordinator (URAC) Lead, under the EHRM Office of Functional Champion (OFC) Deployment Manager, outlines the objectives, scope, methodology, timing and duration, tools and resources, roles and responsibilities, and procedure, to complete the conversion of user roles, including training, from the legacy EHR system (VistA) to the new one (Millennium EHR). While the Computerized Patient Record System (CPRS) in VistA, by design, has permission for each user that can be added, removed, and otherwise customized depending on the user's needs, the new EHR/Millennium uses several "roles" pre-defined by the vendor and set at the national level. Each user of the new system is assigned one or several role(s) that define their access right (authorization). The 'User Role Assignment' (URA) process is essentially to optimize the conversion of a user's legacy permission(s) to the available

role(s) (equivalent to access rights) in Millennium. Once the role(s) for each user have been assigned, the local URAC(s) will follow the procedures documented in the EHRM Access Office Access Management Guide, to complete new user provisioning in Millennium. Concurrently, the local URAC(s) will monitor and ensure the user complete assigned training courses before the site go-live date.

8.1b Identify users from other agencies who may have access to the system and under what roles these individuals have access to the system. Who establishes the criteria for what PII can be shared?

Only authorized VA users can access this EHRM system.

8.1c Describe the different roles in general terms that have been created to provide access to the system? For example, certain users may have "read-only" access while others may be permitted to make certain amendments or changes to the information.

Only VA authorized personnel users can access the system to perform administrative and maintenance jobs.

8.2 Will VA contractors have access to the system and the PII? If yes, what involvement will contractors have with the design and maintenance of the system? Has a contractor confidentiality agreement, Business Associate Agreement (BAA), or a Non-Disclosure Agreement (NDA) been developed for contractors who work on the system?

If so, how frequently are contracts reviewed and by whom? Describe the necessity of the access provided to contractors to the system and whether clearance is required. If Privacy Roles and Responsibilities have been established to restrict certain users to different access levels, please describe the roles and associated access levels. Explain the need for VA contractors to have access to the PII. This question is related to privacy control AR-3, Privacy Requirements for Contractors, and Service Providers.

Yes. Authorized contractor staff of VA EHRM-IO can access the system for administrative and maintenance purposes. All contractor personnel must comply with VA cybersecurity and data safeguarding requirements, including the Contractor Confidentiality Agreement, the Non-Disclosure Agreement, and the Subcontractor Business Associate Agreement revision signed in July 2023 between Oracle Health Government Services and the VA EHRM-IO.

8.3 Describe what privacy training is provided to users either generally or specifically relevant to the program or system?

VA offers privacy and security training. Each program or system may offer training specific to the program or system that touches on information handling procedures and sensitivity of information. Please describe how individuals who have access to PII are trained to handle it appropriately. This question is related to privacy control AR-5, Privacy Awareness and Training.

All eligible and authorized VA users of the system must read and acknowledge the VA National Rules of Behavior (ROB) or VA Contractor's ROB pertaining to everyday behavior expected of Organizational Users, prior to gaining access to any VA/Federal information

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system or sensitive information. The rules are included as part of the annual VA Privacy and Information Security Awareness and Rules of Behavior (WBT) course, ID# 10176, which all VA network authorized users must complete via the VA's Talent Management System (TMS). After the user's initial acceptance of the Rules, the user must re-affirm their acceptance annually as part of the renew/refreshing privacy and security awareness training. Acceptance is obtained via electronic acknowledgment and is tracked through the TMS system. The questions are directed at providing notice to the individual of the scope of information collected, the right to consent to uses of the information, and the right to decline to provide information. System administrators are required to complete additional role-based training. Additionally, these users also need to complete course ID# 10203, HIPAA and Privacy training annually since they will have direct access to PHI in the Millennium system in particular, and the Federal EHR system in general. The curriculum of TMS courses identified and assigned to a user by the URA process is to address different purposes other than privacy awareness & training.

8.4 Has Authorization and Accreditation (A&A) been completed for the system?

Yes, A&A has been completed for the system.

8.4a *If Yes, provide:*

1. *The Security Plan Status:* Approved
2. *The System Security Plan Status Date:* 7 March 2023
3. *The Authorization Status:* Authorization to Operate
4. *The Authorization Date:* 27 April 2023
5. *The Authorization Termination Date:* 26 April 2025
6. *The Risk Review Completion Date:* 21 April 2023
7. *The FIPS 199 classification of the system (LOW/MODERATE/HIGH):* MODERATE

Please note that all systems containing SPI are categorized at a minimum level of "moderate" under Federal Information Processing Standards Publication 199.

8.4b *If No or In Process, provide your Initial Operating Capability (IOC) date.*

N/A

Section 9 – Technology Usage

The following questions are used to identify the technologies being used by the IT system or project.

9.1 Does the system use cloud technology? If so, what cloud model is being utilized?

If so, Does the system have a FedRAMP provisional or agency authorization? If the system does use cloud technology, but does not have FedRAMP authorization, explain how the Cloud Service Provider (CSP) solution was assessed and what FedRAMP documents and processes were used for the assessment in order to comply with VA Handbook 6517. Types of cloud models include: Software as a Service (SaaS), Infrastructure as a Service (IaaS), Platform as a Service (PaaS), Commercial off the Shelf (COTS), Desktop as a Service (DaaS), Mobile Backend as a Service (MBaaS), Information Technology Management as a Service (ITMAaS). This question is related to privacy control UL-1, Information Sharing with Third Parties.

No, the system does not use cloud technology.

9.2 Does the contract with the Cloud Service Provider, Contractors and VA customers establish who has ownership rights over data including PII? (Provide contract number and supporting information about PII/PHI from the contract). (Refer to question 3.3.2 of the PTA) This question is related to privacy control AR-3, Privacy Requirements for Contractors, and Service Providers.

No, the system does not use cloud technology.

9.3 Will the CSP collect any ancillary data and if so, who has ownership over the ancillary data?

Per NIST 800-144, cloud providers hold significant details about the accounts of cloud consumers that could be compromised and used in subsequent attacks. Ancillary data also involves information the cloud provider collects or produces about customer-related activity in the cloud. It includes data collected to meter and charge for consumption of resources, logs and audit trails, and other such metadata that is generated and accumulated within the cloud environment.

This question is related to privacy control DI-1, Data Quality.

No, the system does not use cloud technology.

9.4 NIST 800-144 states, “Organizations are ultimately accountable for the security and privacy of data held by a cloud provider on their behalf.” Is this principle described in contracts with customers? Why or why not?

What are the roles and responsibilities involved between the organization and cloud provider, particularly with respect to managing risks and ensuring organizational requirements are met? This question is related to privacy control AR-3, Privacy Requirements for Contractors and Service Providers.

N/A – The system does not use cloud technology/cloud service provider.

9.5 If the system is utilizing Robotics Process Automation (RPA), please describe the role of the bots.

Robotic Process Automation is the use of software scripts to perform tasks as an automated process that executes in parallel with or in place of human input. For example, will the automation move or touch PII/PHI information. RPA may also be referred to as “Bots” or Artificial Intelligence (AI).

The system does not utilize RPA.

Section 10. References

Summary of Privacy Controls by Family

Summary of Privacy Controls by Family

ID	Privacy Controls
AP	Authority and Purpose
AP-1	Authority to Collect
AP-2	Purpose Specification
AR	Accountability, Audit, and Risk Management
AR-1	Governance and Privacy Program
AR-2	Privacy Impact and Risk Assessment
AR-3	Privacy Requirements for Contractors and Service Providers
AR-4	Privacy Monitoring and Auditing
AR-5	Privacy Awareness and Training
AR-7	Privacy-Enhanced System Design and Development
AR-8	Accounting of Disclosures
DI	Data Quality and Integrity
DI-1	Data Quality
DI-2	Data Integrity and Data Integrity Board
DM	Data Minimization and Retention
DM-1	Minimization of Personally Identifiable Information
DM-2	Data Retention and Disposal
DM-3	Minimization of PII Used in Testing, Training, and Research
IP	Individual Participation and Redress
IP-1	Consent
IP-2	Individual Access
IP-3	Redress
IP-4	Complaint Management
SE	Security
SE-1	Inventory of Personally Identifiable Information
SE-2	Privacy Incident Response
TR	Transparency
TR-1	Privacy Notice
TR-2	System of Records Notices and Privacy Act Statements
TR-3	Dissemination of Privacy Program Information
UL	Use Limitation
UL-1	Internal Use
UL-2	Information Sharing with Third Parties

Signature of Responsible Officials

The individuals below attest that the information provided in this Privacy Impact Assessment is true and accurate.

Privacy Officer, Angela Pluff

Information Systems Security Officer, Jeramy Drake

Information Systems Owner, Michael Hartzell

APPENDIX A-6.1

Please provide a link to the notice or verbiage referred to in Section 6 (a notice may include a posted privacy policy; a Privacy Act notice on forms; screen shot of a website collection privacy notice).

In the <http://www.va.gov/health/> webpage, the current PDF copy of the “VA Privacy Practices” is listed in the “Resources” section on the right.

SORN 24VA10A7, Patient Medical Records-VA: <https://www.govinfo.gov/content/pkg/FR-2020-10-02/pdf/2020-21426.pdf>

HELPFUL LINKS:

Record Control Schedules:

<https://www.va.gov/vhapublications/rcs10/rcs10-1.pdf>

National Archives (Federal Records Management):

<https://www.archives.gov/records-mgmt/grs>

VA Publications:

<https://www.va.gov/vapubs/>

VA Privacy Service Privacy Hub:

<https://dvagov.sharepoint.com/sites/OITPrivacyHub>

VA Privacy Service:

<https://department.va.gov/privacy/>