Privacy Impact Assessment for the VA IT System called:

Managed Service – Laboratory System Reengineering PathNet (LSRP) Assessing Clinical Services

Veterans Health Administration

Date PIA submitted for review:

05/31/2023

System Contacts:

<table>
<thead>
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<tbody>
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</tbody>
</table>
Abstract
The abstract provides the simplest explanation for “what does the system do?” and will be published online to accompany the PIA link.

The purpose of this project is to replace the legacy Laboratory Information Management System (LSRP) with a Commercial Off-The-Shelf (COTS) LSRP. The selected COTS product and managed service, Cerner Millennium PathNet Remote Hosting Option (RHO) that resides at the Cerner Technology Centers in Kansas City, will allow the VA to meet future requirements of Electronic Medical Record, HealtheVet and interoperability between the Department of Defense (DoD) and Public Health Services (PHS) as per public law 107-287.

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. The IT system name and the name of the program office that owns the IT system.

Managed Service – LSRP Assessing

Clinical Services

B. The business purpose of the program, IT system, or technology and how it relates to the program office and agency mission.

The LSRP automates laboratory workflows by interfacing with other VA Health Information System clinical and revenue-cycle workflows to process lab test orders, manage specimens, support laboratory quality checks, and support the delivery of test results to the appropriate clinical systems and providers to enable effective and efficient care delivery. The LSRP also supports laboratory test order processing workflows for 3rd party laboratory testing services. The COTS LSRP and RHO manages service enables the VA to meet future requirements of Electronic Medical Record, HealtheVet and interoperability between the Department of Defense (DoD) and Public Health Services (PHS) as per public law 107-287

C. Indicate the ownership or control of the IT system or project.

VA Controlled / non-VA Owned and Operated
2. Information Collection and Sharing

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

The system is expected to hold the laboratory data of approximately 40,000 Veterans. The LSRP system processes the PII and PHI of all VA patients who receive laboratory services from the VA.

E. A general description of the information in the IT system and the purpose for collecting this information.

The data processed by the LSRP includes VA Veteran or primary subject’s personal contact information (name, address, telephone, etc.); personal identifiers (Social Security Number (SSN), financial account number); family relation; service information; medical record information. Records can be retrieved using full name, SSN, and financial account number. Records can also be retrieved via searches on Medical Record Number, birth date, gender, and unique identifiers assigned by the LSRP (accession numbers). The purpose of the collection is for processing, retainment and sharing.

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

All of the PII and PHI processed by the LSRP is received from VistA as well as Quest Diagnostics and LabCorp Reference Lab systems via HL7 interfaces.

G. Whether the system is 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 is operational in only one site.

3. Legal Authority and SORN

H. A citation of the legal authority to operate the IT system.

The legal authority to operate the system falls under Title 38, United States Code, Section 501.

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?

The system is not in the process of being modified.

D. System Changes

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

The information selected below must match the information provided in question 2.1 as well as the data elements columns in 4.1 and 5.1.
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:

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

- Family Relation (spouse, children, parents, grandparents)
- Service Information
- Medical Information
- Laboratory test orders and results

PII Mapping of Components (Servers/Database)

LSRP consists of 1 key components (servers/databases). Each component has been analyzed to determine if any elements of that component collect PII. The type of PII collected by LSRP 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 the server names in the table. The first table of 3.9 in the PTA should be used to answer this question.

Internal Database Connections

<table>
<thead>
<tr>
<th>Database Name of the information system collecting/storing PII</th>
<th>Does this system collect PII? (Yes/No)</th>
<th>Does this system store PII? (Yes/No)</th>
<th>Type of PII (SSN, DOB, etc.)</th>
<th>Reason for Collection/Storage of PII</th>
<th>Safeguards</th>
</tr>
</thead>
<tbody>
<tr>
<td>Millennium</td>
<td>Yes</td>
<td>Yes</td>
<td>• Name</td>
<td>Perform laboratory test for the</td>
<td>800-53 rev 4 High Baseline,</td>
</tr>
</tbody>
</table>
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?

All of the PII processed by the LSRP is received from VistA as well as Quest Diagnostics and LabCorp Reference Lab systems via HL7 interfaces. The LSRP does not collect data directly from a patient.

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.

All of the PII and PHI processed by the LSRP is received from VistA as well as Quest Diagnostics and LabCorp Reference Lab systems via HL7 interfaces.

1.2c If the system creates information (for example, a score, analysis, or report), list the system as a source of information.

The system does not create any additional 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?

All of the PII and PHI processed by the LSRP is received from VistA as well as Quest Diagnostics and LabCorp Reference Lab systems via HL7 interfaces.

1.3b If the information is collected on a form and is subject to the Paperwork Reduction Act, give the form’s OMB control number and the agency form number.
The information is not collected on a form

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 LSRP provides the capability to support workflows that include manual order review and quality checks. The LSRP relies on the underlying integrity and accuracy of the data provided by source systems.

1.4b If the system checks for accuracy by accessing a commercial aggregator of information, describe this process and the levels of accuracy required by the contract.

The system does not access a commercial aggregator.

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

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.

Title 38, United States Code, Sections 501(b) and 304. 24VA10A7 - Patient Medical Records – VA

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:**
Both Personally Identifiable Information (PII) and a variety of other Sensitive Personal Information (SPI), such as Protected Health Information (PHI) is collected. Due to the highly sensitive nature of this data, there is a risk that, if the data were accessed by an unauthorized individual or otherwise breached, serious personal, professional, or financial harm may result for the individuals affected.

**Mitigation:**
A variety of security measures designed to ensure that the information is not inappropriately disclosed or released is employed. These measures include access control; awareness and training; audit and accountability; certification, accreditation, and security assessments; configuration management; contingency planning; identification and authentication; incident response; maintenance; media protection; physical and environmental protection; planning; personnel security; risk assessment; systems and services acquisition; system and communications protection; and system and information integrity. Our facility employs all security controls in the respective high impact security control baseline unless specific exceptions have been allowed based on the tailoring guidance provided in National Institute of Standards and Technology (NIST) Special Publication 800-37.
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 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.

The information collected will be used to provide high quality laboratory services in support of the VA’s mission of utilizing high quality, effective, and efficient Information Technology services to provide benefits and services to the Veterans of the United States.

- Name – Used to identify the patient during appointments and in other communication.
- Social Security Number – Used as a patient identifier.
- Date of Birth – Used to identify age and confirm patient identity.
- Mailing Address – Used as a patient identifier.
- Phone Number – Used as a patient identifier.
- Health Insurance – Used to link records between LSRP and VISTA systems.

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.

No data analysis of this kind is conducted.

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.

Results from laboratory services are appended to the patient’s current record in VistA.

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?

AES-256 encryption is in place

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

AES-256 encryption is in place

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

Industry standard encryption is in place - AES-256.

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?

LSRP provides the capability to limit access to patient information to authorized users of the LSRP through role-based access control. The LSRP provides security and user event logging including the actions taken by users on the data.

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

Yes
2.4c Does access require manager approval?

Access does require manager approval

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

Access to PII is being monitored, tracked and recorded

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

The Information System Owner (ISO) is responsible.

Coordinates with the Information System Security Officer (ISSO) and Privacy Officer (PO) to complete a Privacy Threshold Analysis (PTA) at least annually, and a Privacy Impact Assessment (PIA) at least every three years

Monitors and ensures compliance with Privacy Controls

Delegates appropriate Privacy duties

Requires maintenance of evidence and documentation as eMASS artifacts

Approves/authorizes access to VA LSRP PII/PHI

Is responsible for VA LSRP Privacy compliance

Performs Privacy review and approval duties as assigned or tasked

Collaborates with data owners, POs, ISSOs and the Privacy Service to obtain/confirm VA LSRP System of Records Notices (SORN)

Updates all SORNs as dictated by law or VA policy

Requires all personnel maintain confidentiality of PII/PHI

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*

Veteran or primary subject's personal contact information (name, address, telephone, etc.); family relation; service information; medical information. Records can be retrieved using full name, SSN,
and financial account number. Records can also be retrieved via searches on accession number, Medical Record Number, birth date, and gender.

### 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.

Clinical information is retained in accordance with VA Records Control Schedule 10-1. Demographic information is updated as applications for care are submitted and retained in accordance with VA Records Control Schedule 10-1. Retention period for the PHI is 75 years after date of last episode of patient care.

### 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. 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

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

VA Records Control Schedule 6000.2(b) (https://www.va.gov/vhapublications/RCS10/rcs10-1.pdf)

### 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.
Prior to termination or completion of this contract, Cerner LSRP and their affiliated business partners must receive written approval from the VHA before any VA/VHA provided information is destroyed. Any data destruction done on behalf of the VA/VHA must be done in accordance with National Archives and Records Administration (NARA) approved records schedules found in VHA RCS 10-1. Destroyed in accordance with records control schedule making the data unidentifiable. Electronic data and files of any type, including Protected Health Information (PHI), Sensitive Personal Information (SPI), and more are destroyed in accordance with the Department of Veterans’ Affairs Handbook 6500.1, Electronic Media Sanitization (November 3, 2008), http://www.va.gov/vapubs/viewPublication.asp?Pub_ID=416&FType=2. When required, this data is deleted from their file location and then permanently deleted from the deleted items or Recycle bin. Magnetic media is wiped and sent out for destruction per VA Handbook 6500.1. Digital media is shredded or sent out for destruction per VA Handbook 6500.1. Huntington VAMC additionally follows Field Security Service (FSS) Bulletin #176 dated April 9, 2014 for MediaSanitization Program, https://www.vendorportal.ecms.va.gov/FBODocumentServer/DocumentServer.aspx?DocumentId=4214767&FileName=36C25518Q0329-011.pdf as well as FSS Standard Operating Procedures (SOP) MP-6 Electronic Media Sanitization.

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.

Follow the format below:

**Privacy Risk:**

The risk associated with storing this data increases the longer the data is stored.

**Mitigation:**

To mitigate the risk posed by information retention, Cerner LSRP adheres to the VA RCS schedules for each category or data it maintains. When the retention date is reached for a record, the medical center and Cerner LSRP will carefully dispose of the data by the determined method as described in RCS 10-1

**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?  
**NOTE:** Question 3.9 (second table) on Privacy Threshold Analysis should be used to answer this question.

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.
Data Shared with Internal Organizations

<table>
<thead>
<tr>
<th>List the Program Office or IT System information is shared/received with</th>
<th>List the purpose of the information being shared/received with the specified program office or IT system</th>
<th>List the specific PII/PHI data elements that are processed (shared/received/transmitted) with the Program Office or IT system</th>
<th>Describe the method of transmittal</th>
</tr>
</thead>
</table>
| Veterans Health Administration | Laboratory Services | • Name  
• Address  
• Email  
• Telephone number  
• Social security number  
• Date of birth | Electronically pulled from VistA through computerized patient record system (CPRS) to VALSRP program |

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.

Follow the format below:

**Privacy Risk:**

Policies & procedures, Data protection deficiency, Insider Threat, Unauthorized Access

**Mitigation:**

Access Control, policies and guidelines on sensitive data, regular training, implement technical controls such as firewalls, intrusion detection systems, encryption

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.

NOTE: Question 3.10 on Privacy Threshold Analysis should be used to answer this question.

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

<table>
<thead>
<tr>
<th>List External Program Office or IT System information is shared/received with</th>
<th>List the purpose of information being shared / received / transmitted with the specified program office or IT system</th>
<th>List the specific PII/PHI data elements that are processed (shared/received/transmitted) with the Program or IT system</th>
<th>List the legal authority, binding agreement, SORN routine use, etc. that permit external sharing (can be more than one)</th>
<th>List the method of transmission and the measures in place to secure data</th>
</tr>
</thead>
</table>
| Quest Diagnostics | Lab orders may be transmitted to and results received from Quest if facility is used | • Name  
• Address  
• Email  
• Telephone number  
• Social security number  
• Date of birth | National MOU/ISA | Site to Site (S2S), IPSEC Tunnel, Secure FTP |
| LabCorp Reference Lab | Lab orders may be transmitted to and results received from LabCorp if facility is used | • Name  
• Address  
• Email  
• Telephone number | National MOU/ISA | Site to Site (S2S), IPSEC Tunnel, Secure FTP |
| Cerner (LSRP)                  | To automate laboratory workflows by interfacing with other VA Health Information System clinical and revenue-cycle workflows. | • Name  
• Address  
• Email  
• Telephone number  
• Social security number  
• Date of birth | National MOU/ISA | Site to Site (S2S), IPSEC Tunnel, Secure FTP |

### 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.

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

Follow the format below:

**Privacy Risk:**
Unauthorized access, data protection, data breaches, compliance

**Mitigation:**
Encrypted data transfer, authentication and access control, monitor & audit, MOUS/ISA

### 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, TR-2, and TR-3, System of Records Notices and Privacy Act Statements, and 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.

Notice:
The Department of Veterans Affairs provides additional notice of this system by publishing 2 System of Record Notices (SORNs):


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

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.

Cerner LSRP mitigates this risk by ensuring that it provides individuals notice of information collection and notice of the system’s existence through the System of Record Notice https://www.govinfo.gov/content/pkg/PAI-2019-VA/xml/PAI-2019-VA.xml#24VA10P2.

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.

LSRP does not collect information directly from the patient.
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.

LSRP does not collect information directly from the patient.

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:

**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.

Follow the format below:

**Privacy Risk:**
There is a risk that veterans and other members of the public will not know that the Cerner LSRP exists or that it collects, maintains, and/or disseminates Personally Identifiable Information (PII) and other Sensitive Personal Information (SPI) about them.

**Mitigation:**
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.

Darryl Webb, FOIA Officer
202-738-2974
vacofoiaservice@va.gov

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).

The system is not exempt

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.

Individuals must follow established procedures to gain access to their information under the guidelines of the Privacy Act, Freedom of Information Act (FOIA), and Health Insurance Portability and Accountability Act (HIPAA). When requesting access to one’s own records, patients are asked to complete VA Form I 0-5345a: Individuals’ Request for a Copy of their Own Health Information, which can be obtained from the medical center or online at http://www.va.gov/vafonns/medical/pdf/vha-10-5345a-fill.pdf. Additionally, veterans and their dependents can gain access to their Electronic Health Record (EHR) by enrolling in the MyHealthVet program, VA's online personal health record. More information about MyHealthVet is available at https://www.myhealth.va.gov/index.html.

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,
When a Veteran has concerns that something written in his/her medical records is inaccurate, incomplete or needs to be removed entirely, they have the right to request an amendment of the record. The Veteran must submit their request in writing, specify the information that they want corrected, and provide a reason to support the request for amendment. All amendment requests must be submitted to the facility Privacy Officer. The Privacy Officer refers the request and related record to the health care provider who authored the information in order for the provider to determine if the record needs to be amended as requested. If the author is not available, the documentation is referred to that provider’s supervisor. If the amendment is granted, the facility Privacy Officer will work with the responsible record custodian to amend their records. If the amendment is not granted, the Privacy Officer will notify the Veteran of the decision and include appeal rights.

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 facility staff and providers are educated to refer the Veteran to the Privacy Officer for requests to correct their records. At the time of the request the individual is sent an acknowledgement letter and they are sent a letter at the completion of processing regarding the outcome of the requested correction.

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.

Redress is provided through the Privacy Act for the individual to view and request correction to the inaccurate or erroneous information. If the request is denied, the individual may appeal the decision by writing to the Office of General Counsel (024); Department of Veterans Affairs; 810 Vermont Avenue, N.W.; Washington, D.C. 20420. The Privacy Act and HIPAA permit the individual to also complete a Statement of Disagreement to the information that was denied correction. The facility would be able to include a rebuttal to the Statement of Disagreement. The Statement of Disagreement, rebuttal, and denial letter would be attached to the information.
that was requested to be corrected and would be released with the information at any time the information was authorized for release.

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.

Follow the format below:

Privacy Risk:
If a system does not allow individual access, the risk of inaccurate information may occur. Additionally, patients should have access to the information so they can keep abreast of labs which are outside of normal and plan their healthcare accordingly. If a healthcare provider depends on inaccurate information, the patient could be given the wrong treatment.

Mitigation:
The Privacy Risk is low-moderate, as the information is processed through the Privacy Act, HIPAA, and FOIA. FOIA protects specific records with exemptions. When information is processed under FOIA, the exemptions and an explanation of the exemption are included in the response to the request. The individual has a right to access their individual information under the Privacy Act, when that information is part of a Privacy Act System of Records. An individual’s identity is confirmed in requesting access, redress, and correction of information through legal authority (POA, Guardian, Next of Kin), photo identification and/or wet signature, which protect the information from being used without the individual’s knowledge. Appeal rights are given to an individual upon denial of a correction.

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.

Types of accounts are defined as being “named accounts” and “support accounts”. “Named accounts” have rights and privileges based on a pre-determined profile for an individual’s role. “Support accounts” are temporary and provide short-term access for troubleshooting or project-oriented work on a client’s processing environment.

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?

No users identified

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.

These accounts, generated by Cerner Technology Services (CTS) Enterprise Security Identity Access Management, grant limited access appropriate to the type of troubleshooting and/or project work required as outlined in the Performance Work Statement.

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.

Only VA-approved Cerner contractor personnel will be able to access the data. Approved personnel will go through the VA onboarding process, receive a clearance, and complete required training before access is granted.

Training Management System (TMS)

- VA Privacy and Information Security Awareness and Rules of Behavior (WBT)
- Privacy and HIPAA Training
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 users of the LSRP system are assigned privacy and security awareness training on an annual basis. The training is conducted through the VA TMS system that tracks and monitors students training records. Training is setup to test students understanding of VA policy and procedures regarding use and handling of PHI and PII information.

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

8.4a If Yes, provide:

1. The Security Plan Status: Approved
2. The System Security Plan Status Date: 05-Jun-2023
3. The Authorization Status: Authorization to Operate (ATO)
4. The Authorization Date: 16-Mar-2023
5. The Authorization Termination Date: 09/12/2023
6. The Risk Review Completion Date: 03/15/2023
7. The FIPS 199 classification of the system (LOW/MODERATE/HIGH): HIGH

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.

This document is part of the ATO submission package; however a current 180-day ATO-C exists for the system Dated 06 January 2021.

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.
Note: For systems utilizing the VA Enterprise Cloud (VAEC), no further responses are required after 9.1. (Refer to question 3.3.1 of the PTA)

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.

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.

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.

The system does not use cloud technology

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 use Robotics Process Automation (RPA)
## Section 10. References

### Summary of Privacy Controls by Family

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Signature of Responsible Officials

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

Diana L Bowen
166536
Digitally signed by Diana L Bowen
Date: 2023.06.28 12:13:02 -04'00'

Privacy Officer, Diana Bowen

Craig J. Heitz
564885
Digitally signed by Craig J. Heitz
Date: 2023.06.28 13:09:28 -05'00'

Information Systems Security Officer, Craig Heitz

Christopher Brown 101386
Digitally signed by Christopher Brown 101386
Date: 2023.06.29 09:04:50 -05'00'

Information Systems Owner, Christopher Brown
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).

HELPFUL LINKS:

Record Control Schedules:

General Records Schedule 1.1: Financial Management and Reporting Records (FSC):

National Archives (Federal Records Management):
https://www.archives.gov/records-mgmt/grs

VHA Publications:
https://www.va.gov/vhapublications/publications.cfm?Pub=2

VA Privacy Service Privacy Hub:
https://dvagov.sharepoint.com/sites/OITPrivacyHub

Notice of Privacy Practice (NOPP):
VHA Notice of Privacy Practices
VHA Handbook 1605.04: Notice of Privacy Practices