



Privacy Impact Assessment for the VA IT System called:

COVID-19 Patient Manager VA OIT Office of the CTO

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<< 3/19/2021 >>

System Contacts:

System Contacts

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Abstract

The abstract provides the simplest explanation for “what does the system do?” and will be published online to accompany the PIA link.

Covid-19 Patient Manager (CPM) is a standards-based (utilizing Substitutable Medical Applications and Reusable Technologies (SMART) on Fast Healthcare Interoperability Resources (FHIR)) web application that will be accessible from within the electronic health record and will provide guideline-based recommendations on care for patients affected by Covid-19. This application requires the development of technical infrastructure to interact with electronic health record data, launch from within the electronic health record, authenticate users, maintain appropriate security, and other capabilities. This technical infrastructure will constitute a clinical decision support platform that can be used to quickly launch additional applications within the electronic health record and assist physicians with rapidly responding to future healthcare emergencies.

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:

- *The IT system name and the name of the program office that owns the IT system.*
- *The business purpose of the program, IT system, or technology and how it relates to the program office and agency mission.*
- *The expected number of individuals whose information is stored in the system and a brief description of the typical client or affected individual.*
- *If your system is a regional GSS, VistA, or LAN, include a list of the hospitals/medical centers, or other regional offices that fall under your system. Additionally, what region is the system under?*
- *A general description of the information in the IT system.*
- *Any information sharing conducted by the IT system. A general description of the modules and subsystems, where relevant, and their functions.*
- *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.*
- *A citation of the legal authority to operate the IT system.*
- *Whether the completion of this PIA will result in circumstances that require changes to business processes*
- *Whether the completion of this PIA could potentially result in technology changes*
- *If the system is in the process of being modified and a SORN exists, will the SORN require amendment or revision and approval?*

- *Does the system use cloud technology? 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.*
- *Does a contract with Cloud Service Provider, Contractors and VA customers establish who has ownership rights over data including PII?*
- *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 is the magnitude of harm if privacy related data is disclosed, intentionally or unintentionally? Would the reputation of the CSP or its customers (VA) be affected?*

1. Covid-19 Patient Manager (CPM) is owned by VA OIT, Office of the Chief Technology Officer, and is being developed in collaboration with VA Clinical Informatics and Data Management Office (CIDMO).
2. The business purpose is as follows: CPM addresses Customer Service and Transforming Business Operations. This will help clinicians to administer a high standard of care by adhering to new clinical guidelines. It will also help clinicians to work more efficiently by assisting with managing and acting upon the high volumes of data clinicians must work with, including rapidly evolving inpatient data, outpatient symptoms monitoring, and other patient-generated data. This clinical decision support application will provide recommendations at the point of care, overcoming current shortfalls to integrate them into the electronic health record (the centerpiece of clinician workflow).
3. Individuals’ information will not be stored by the CPM system.
4. The CPM system is not a regional GSS, VistA, or LAN.
5. A general description of the information accessed by the system is as follows: The CPM system will access patients’ electronic health record data (patient name, date of birth, sex assigned at birth, problem list, vital signs, laboratory results, imaging study results, medications) in order to compute evidence-based recommendations about next steps in care. This data is not stored by the CPM system. The logic of these evidence-based recommendations is supplied and validated by institutions including the Agency for Healthcare Research and Quality (AHRQ) and the American College of Emergency Physicians (ACEP). The system does not write back to the electronic health record; it is read-only.
6. The CPM system does not share any data. All data is retrieved from VistA through the Lighthouse Health API or Veterans Data Integration and Federation (VDIF).
7. This system will be used in multiple VA Medical Centers. The same controls will be used across all sites. CPM is a standards-based (utilizing Substitutable Medical Applications and Reusable Technologies (SMART) on Fast Healthcare Interoperability Resources (FHIR)) web application, and it is through use of these standards that consistency of the PII accessed by the system will be maintained.

8. The Legal Authority to operate the CPM system is: SOR 24VA10A7 Patient Medical Records; Title 38, USC 501(b) and 304
9. The completion of this PIA will not result in circumstances that require changes to business processes.
10. The completion of this PIA will not result in technology changes.
11. No SORN is necessary for the CPM system.
12. The CPM system uses cloud technology. The CPM system is deployed on AWS FedRAMP cloud in accordance with all applicable regulations. AWS Government Cloud is authorized under FedRAMP.
13. The CPM system does not store any data. Data ownership is not applicable.
14. The CPM system does not store any data. Organizational accountability is not applicable.
15. The CPM system does not store any data. Release of data is not applicable.

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://vawww.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 type="checkbox"/> Personal Mailing Address | <input type="checkbox"/> Personal Fax Number |
| <input type="checkbox"/> Social Security Number | <input type="checkbox"/> Personal Phone Number(s) | <input type="checkbox"/> Personal Email Address |
| <input checked="" type="checkbox"/> Date of Birth | | <input type="checkbox"/> Emergency Contact Information (Name, Phone) |
| <input type="checkbox"/> Mother's Maiden Name | | |

Number, etc. of a different individual)

Financial Account Information

Health Insurance Beneficiary Numbers

Account numbers

Certificate/License numbers

Vehicle License Plate Number

Internet Protocol (IP) Address Numbers

Current Medications

Previous Medical Records

Race/Ethnicity

Tax Identification Number

Medical Record Number

Other Unique Identifying Number (list below)

<<Add Additional Information Collected But Not Listed Above Here (For Example, A Personal Phone Number That Is Used As A Business Number)>>

We are processing the following information: Patient identifiers (name (first and last), date of birth, Internal Control Number (ICN)), problem list, diagnoses, vital signs, diagnostic tests, laboratory results, medications, procedure history.

PII Mapping of Components

CPM consists of 1 key components. Each component has been analyzed to determine if any elements of that component collect PII. The type of PII collected by VistA and the reasons for the collection of the PII are in the table below.

PII Mapped to Components

PII Mapped to Components

Components of the information system (servers) collecting/storing PII	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
VistA	CPM system does not store PII. CPM system accesses PII from VistA through the Lighthouse Health API or through VDIF.	CPM system does not store PII.	Patient identifiers (name (first and last), date of birth, ICN), problem list, diagnoses, vital signs, diagnostic tests, laboratory results, medications,	PII is accessed by the CPM system so that clinical recommendations can be made to VA physicians.	The CPM system does not store data. All OIT Server security controls are in place, as are VAEC security controls.

			procedure history		

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

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?

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 1.3 indicate why the system is using this source of data.

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

This question is related to privacy controls DI-1, Data Quality, and IP-1, Consent.

The information is collected from the electronic health record (VistA). This information is accessed from the electronic health record because the individual patient may not know the information. The system uses this information to create a Covid-19 severity score and recommendations on next steps for care, which can be copy/pasted by a healthcare provide from the CPM application into an electronic health record note if the provider would like to do that.

1.3 How is the information collected?

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 technology used in the storage or transmission of information in identifiable form?

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.

This question is related to privacy controls DI-1, Data Quality, and IP-1, Consent.

Information is collected electronically through an API with the VA Medical Center’s VistA.

1.4 What is the purpose of the information being collected, used, disseminated, created, or maintained?

Include a statement of why the particular SPI is collected, maintained, used, or disseminated in the system is necessary to the program's or agency's mission. Merely stating the general purpose of the system without explaining why this particular type of information should be collected and stored is not an adequate response to this question.

If the system collects, uses, disseminates, or maintains publicly available or commercial data, include a discussion of why commercial data is relevant and necessary to the system's purpose. This question is related to privacy control AP-2, Purpose Specification.

The CPM system requires this data to address Customer Service and Transforming Business Operations. This system and the data it accesses will help clinicians to administer a high standard of care by adhering to new clinical guidelines regarding Covid-19 care. It will also help clinicians to work more efficiently by assisting with managing and acting upon the high volumes of data clinicians must work with, including rapidly evolving inpatient data, outpatient symptoms monitoring, and other patient-generated data. This clinical decision support application will provide recommendations at the point of care, overcoming current shortfalls to integrate them into the electronic health record.

The system does not collect, use, disseminate, or maintain publicly available or commercial data.

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

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.

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. This question is related to privacy controls DI-1, Data Quality, and DI-2, Data Integrity and Integrity Board.

Information is checked for accuracy within VistA. Clinical practitioners review and sign all treatment information and Business Office/Health Information Management Service reviews data obtained and assists with corrections.

1.6 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

SOR 24VA10A - Patient Medical Records Title 38, United States Code, Section 501(b) and 304.
Link: https://www.oprm.va.gov/docs/Current_SORN_List_02_02_2021.pdf

1.7 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.

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 CPM system accesses and displays PII and other sensitive PHI. IF this information were breached or accidentally released to inappropriate parties or the public, it could result in personal, and/or emotional harm to the individuals whose information were inappropriately accessed.

Mitigation: The CPM system is careful to display only the information necessary to accomplish the VA mission of assisting physicians with making clinical decisions about caring for individual patients with Covid-19. The clinician-facing application requires authentication and

authorization to ensure that the user accessing the application is appropriately authorized to access the data displayed by the application. Workstations at VA Medical Centers are designed to maximize physical security of the information being displayed on a given screen. The system logs are securely maintained, and access to the audit logs is limited to personnel with security-related roles and auditors.

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 CPM system does not store or maintain any data. The CPM system retrieves and displays data from VistA via Lighthouse Health API or VDIF FHIR API. This data is used to compute recommendations for next steps in care for patients with Covid-19. These recommendations and data are displayed to authenticated physicians so that they may make decisions about next steps in patient care.

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

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.

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.

This question is related to privacy controls DI-1, Data Quality, DI-2, Data Integrity and Integrity Board, and SE-1, Inventory of Personally Identifiable Information

The CPM system generates recommendations about a patient's care based on their clinical data; however, the recommendations are not stored or written back into the electronic health record.

2.3 PRIVACY IMPACT ASSESSMENT: Use of the information. How is access to the PII determined? Are criteria, procedures, controls, and responsibilities regarding access documented? Does access require manager approval? Is access to the PII being monitored, tracked, or recorded? Who is responsible for assuring safeguards for the PII?

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.

Add answer here:

The CPM system utilizes SSOi for authorization of users. Access to the CPM system is restricted to authorized VA Medical Center personnel who have access to the underlying VistA data.

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

The CPM system does not retain data.

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.
This question is related to privacy control DM-2, Data Retention and Disposal.*

N/A – the CPM system does not retain data.

3.3 Has the retention schedule been approved by the VA records office and the National Archives and Records Administration (NARA)? If so please indicate the name of the records retention schedule.

*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.*

N/A – the CPM system does not retain data.

3.4 What are the procedures for the elimination of SPI?

*Explain how records are destroyed or eliminated at the end of the 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*

N/A – the CPM system does not retain data.

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. Have policies and procedures been developed to minimize the use of PII for testing, training, and research? This question is related to privacy control DM-3, Minimization of PII Used in Testing, Training and Research

N/A – the CPM system does not retain data.

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.

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: N/A – the CPM system does not retain data.

Mitigation: N/A – the CPM system does not retain data.

Section 4. Internal Sharing/Receiving/Transmitting and Disclosure

The following questions are intended to define the scope of information sharing/receiving/transmitting within VA. NOTE: Question 5 on Privacy Threshold Analysis should be used to answer this question.

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.

Data Shared with Internal Organizations

<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 data element types such as PII/PHI that are shared/received with the Program Office or IT system</i>	<i>Describe the method of transmittal</i>
VistA via Lighthouse Health API or Veterans Data Integration and Federation (VDIF) API	To provide physicians with clinical data relevant to making decision about patients with Covid-19.	Patient identifiers (name (first and last), date of birth, ICN), problem list, diagnoses, vital signs, diagnostic tests, laboratory results, medications, procedure history	Secure Representational State Transfer (REST) Service between CPM and Lighthouse Health API or VDIF.

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.

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

Follow the format below:

Privacy Risk: The privacy risk associated with displaying data within the Department of Veterans Affairs is that the data may be disclosed to individuals who do not require access or have a need to know. Inappropriate/unauthorized disclosure heightens the threat of the information being misused.

Mitigation: The CPM system is careful to display only the information necessary to accomplish the VA mission of assisting physicians with making clinical decisions about caring for individual patients with Covid-19. The clinician-facing application requires authentication and authorization to ensure that the user accessing the application is appropriately authorized to access the data displayed by the application. Workstations at VA Medical Centers are designed to maximize physical security of the information being displayed on a given screen. The system logs are securely maintained, and access to the audit logs is limited to personnel with security-related roles and auditors.

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: This question is #7 in the Privacy Threshold Analysis.

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 data element types such as PII/PHI that are shared/received 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>
N/A	N/A	N/A	N/A	N/A

If specific measures have been taken to meet the requirements of OMB Memoranda M-06-15 and M-06-16, note them here.

CPM does not share/receive data from external systems.

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: CPM does not share/receive data from external systems.

Mitigation: CPM does not share/receive data from external systems.

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 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?

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. If notice was provided in the Federal Register, provide the citation.

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

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. This question is 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.

N/A – CPM does not collect or retain data. The system does not collect data from clinicians or providers either. It presents read-only information to them, which they can use in their decision-making. They do not write anything back to the electronic health record from the CPM system.

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

N/A – CPM does not collect or retain data. The system does not collect data from clinicians or providers either. It presents read-only information to them, which they can use in their decision-making. They do not write anything back to the electronic health record from the CPM system.

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

N/A – CPM does not collect or retain data. The system does not collect data from clinicians or providers either. It presents read-only information to them, which they can use in their decision-making. They do not write anything back to the electronic health record from the CPM system.

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.

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:

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?

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.

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).

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 question is related to privacy control IP-2, Individual Access, and AR-8, Accounting of Disclosures.

N/A – CPM does not collect or retain data.

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.

N/A – CPM does not collect or retain data.

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.

N/A – CPM does not collect or retain data.

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.

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

Example: Some projects allow users to directly access and correct/update their information online. This helps ensure data accuracy.

N/A – CPM does not collect or retain data.

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.

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:

Mitigation:

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?

Describe the process by which an individual receives access to the system.

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?

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.

This question is related to privacy control AR-7, Privacy-Enhanced System Design and Development.

Veterans Health Administration (VHA) has established policies and procedures for the identification and authorization of VistA/CPRS users. CPM follows these previously established mechanisms.

Access is restricted to VA employees who must complete both the Privacy and HIPAA Focused and Information Security training. Specified access is granted based on the employee's functional category. Role based training is required for individuals with significant information security responsibilities to include but not limited to Information Security Officer (ISO), local Chief Information Officer (CIO), System Administrators, Network Administrators, Database Managers, Users of VA Information Systems or VA Sensitive Information. Users submit access requests based on need to know and job duties. Supervisor, ISO and OI&T approval must be obtained prior to access granted. These requests are submitted for VA employees, contractors and all outside agency requests and are processed through the appropriate approval processes. Once access is granted, individuals can log into the system(s) through dual authentication, i.e., a PIV card with a complex password combination. Once inside the system, individuals are authorized to access information on a need to know basis.

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 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.

VA contractors that have access to the computer system are only delegated keys and menu functions needed to complete their duty task. They are required to complete annual Privacy,

Security, and Rules of Behavior training. Contractors having access to PHI/PII are required to have a Business Associate Agreement (BAA) (nationally with the Veterans Health Administration (VHA) or locally with facility). Contracts are reviewed on an annual basis by the Contracting Officer Representative (COR). The Privacy Officer and Information Security Officer monitor that the annual Privacy, Security, and Rules of Behavior (ROB) training is completed by contractors and business associates. Any local BAAs are monitored by Privacy Officer to ensure compliance with the Health Insurance Portability and Accountability Act (HIPAA). VA contractors under contract to perform system development and test system activities shall use redacted test patient data. No PII/PHI data is used in development or test systems. All contractors accessing VA systems or data are required to sign an NDA prior to beginning their work.

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.

Personnel that will be accessing information systems must read and acknowledge their receipt and acceptance of the VA National ROB or VA Contractor's ROB prior to gaining access to any VA information system or sensitive information. The rules are included as part of the security awareness training which all personnel must complete via the VA's Talent Management System (TMS). After the user's initial acceptance of the ROB, the user must re-affirm their acceptance annually as part of the security awareness training. Acceptance is obtained via electronic acknowledgment and is tracked through the TMS system. System administrators are required to complete additional role-based training. Users with access to PHI are required to complete HIPAA privacy training annually.

- PRIVACY AND HIPPA TRAINING
- VA PRIVACY & VA INFORMATION SECURITY

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

If Yes, provide:

1. *The date the Authority to Operate (ATO) was granted,*
2. *Whether it was a full ATO or ATO with Conditions,*
3. *The amount of time the ATO was granted for, and*
4. *The FIPS 199 classification of the system (LOW/MODERATE/HIGH).*

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

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

ATO was granted on 4/23/21, and ATO was granted for 180 days, until 10/20/21. Categorization for the system is moderate.

Section 9. 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

ID	Privacy Controls
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, Christian Loftus

Information Systems Security Officer, Richard Alomar-Loubriel

System Owner, Shane Elliott

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).