Privacy Impact Assessment for the VA IT System called:

Inspire Sleep Physician Programmer
Pulmonary (Sleep)
VHA James H. Quillen VA Medical Center

Date PIA submitted for review:
June 13, 2022

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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.

The Inspire Physician Programmer is a tablet computer and a telemetry cable. The tablet computer has specific software that will allow the Sleep Provider to activate and make adjustments to the Implantable Pulse Generator (IPG) that was surgically placed into the patient at a Community Care facility. It is the only component in the Inspire system that contains software. The Physician Programmer is a fancy remote that can change the settings to the implant that is inside the patient’s body without having to touch the patient.

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.
- Indicate the ownership or control of the IT system or project.
- The expected number of individuals whose information is stored in the system and a brief description of the typical client or affected individual.
- A general description of the information in the IT system and the purpose for collecting this information.
- 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? If the system is using cloud technology, does the SORN for the system cover cloud usage or storage?

Inspire Physician Programmer owned by Medicine Service and will be used solely by James H. Quillen VA Medical Center (JHQVAMC). The purpose of this system is to provide our veterans an alternative treatment to Obstructive Sleep Apnea. This system is a tablet computer and a telemetry cable. The tablet computer has specific software that will allow the Sleep Provider to make adjustments to the Implantable Pulse Generator (IPG) that was surgically placed into the patient at a Community Care facility. It is the only component in the Inspire system that contains software. The Physician Programmer is a fancy remote that can change the settings to the implant that is inside the patient’s body without
having to touch the patient. The telemetry head communicates with the IPG through the skin via short-range radiofrequency (RF) telemetry; telemetry communication allows the physician to adjust the settings and programs the IPG without touching the person. The telemetry head is powered by a wall outlet connection and wirelessly communicates (via wireless Bluetooth link) with the physician programmer tablet. The physician programmer has the capability to monitor respiratory waveforms, program stimulation modes, adjust stimulation parameter values, and store waveforms and settings. The Physician Programmer is basically only operational when connected to a patient’s device. When it is connected to the patient’s device, it pulls automatically the serial number of the IPG. The Sleep provider enters the patient’s first name, last name, and last 4 of Social Security number only to be able to identify the specific patient for future clinic visits to make adjustments to the IPG. Each Patient is unique may require multiple visits in order to achieve the adequate response to Inspire treatment.

No information sharing will be conducted with the system. This system only receives the serial number from the IPG and the Sleep Provider enters the First Name, Last Name, and the last four of the Social Security number. The completion of this PIA will result in no changes in either the business processes or technology changes.

The following VA System of Record Notice (SORN) applies to the Inspire Physician Programmer:


The legal authorities to operate the Inspire Physician Programmer are Title 5, United States Code, Section 501. Title 38 United States Codes, sections 3686(b), 3323(a); 10 U.S.C.16131, and 38 CFR 21.74256(b).

**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/](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.*

Version Date: October 1, 2021
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 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
- Gender
- Integration Control Number (ICN)
- Military History/Service Connection
- Next of Kin
- Other Unique Identifying Information (list below)

Serial Number of the Implantable Pulse Generator (IPG) (Automatically pulled into the Physician Programmer)

PII Mapping of Components

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

PII Mapped to Components

Note: Due to the PIA being a public facing document, please do not include the server names in the table.
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 gathered from the patient during their appointment and the Implantable Pulse Generator serial number is automatically pulled into the Inspire Physician Programmer.

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.

The information is entered by the Sleep Provider and the Implantable Pulse Generator serial number is automatically pulled into the Inspire Physician Programmer.
1.4 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.

Patient’s information can be checked for accuracy by verifying with their Electronic Health Record.

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

James H. Quillen VA Medical Center (JHQVAMC) Inspire Physician Programmer is maintained under the Veterans Benefits Act, Chapter 73: Veterans Health Administration-Organization and Functions, Title 38, United States Code§7301; per VA information systems and regulatory bodies. Title 38, United States Code, sections 3686(b), 3323(a); Title 10 United States Code 16131, and Title 38 section CFR 21.74256(b).

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.

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:** Collects both personally identifiable information (PII) and a variety of other sensitive personal information (SPI) such as protected health information (PHI). There is a risk that an unauthorized person could get information and that it would result in serious personal, professional, or financial harm.

**Mitigation:** JHQVAMC makes sure by a variety of security measures created to ensure the information is not inadvertently released or disclosed. These measures include access control, awareness/training, audit and accountability, certification, accreditation, security, risk assessment, configuration management, and media protection. Our facility ensures all security controls are in place.

**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.

Name: Use as an Identifier
Last Social Security Number: Used as an Identifier
Serial Number of the Inspire Implantable Pulse Generator (IPG): Used as an Identifier.

This information will be used to keep track of the individual patients due to they may have multiple visits to get the settings of the IPG working efficiently.

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

Reports on how the treatment is working or not working for the patient that has the Inspire Implantable Pulse Generator (IPG) device surgically implanted.

2.3 How is the information in the system secured?
   2.3a What measures are in place to protect data in transit and at rest?

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

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

This question is related to security and privacy controls SC-9, Transmission Confidentiality, and SC-28, Protection of Information at Rest

There are controls in place to assure information is handled in accordance with the uses described above include annual mandatory online information security and HIPAA & Privacy focused training; face-to-face training for all incoming employees conducted by the Information Security Officer and Privacy Officer; regular audits of individuals accessing sensitive information; and formal rounds during which personal examination of all areas within the facility to ensure information is being appropriately used and controlled.

VA systems are intended to be used by authorized VA network users for viewing and retrieving information only except as otherwise explicitly authorized for official business and limited personal use under VA policy.

Information from this system resides on and transmits through computer systems and networks funded by the VA. All access or use constitutes understanding and acceptance that there is no reasonable expectation of privacy in the use of Government networks or systems. All access or use of this system constitutes user understanding and acceptance of these terms and constitutes
unconditional consent to review and action including (but not limited to) monitoring recording copying auditing inspecting investigating restricting access blocking tracking disclosing to authorized personnel or any other authorized actions by all authorized VA and law enforcement personnel.

2.4 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:

Only Sleep providers will have access to the Inspire Physician Programmer. The Sleep provider will ensure that the Physician Programmer is kept in a secure location when not in use.

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 information that will be retained on the Physician Programmer will be what the Sleep provider will enter the Patient’s name, last 4 of Social Security number, settings that have been
set for each patient and the serial number of the Implantable Pulse Generator (IPG) which is automatically pulled from it that was surgically placed previously.

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. If the system is using cloud technology, will it be following the NARA approved retention length and schedule?

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.

The data should be retained on the Inspire Physician Programmer for 6 years after the provider has left the facility and routine records will be disposed of when the agency determines they are no longer needed for administrative or other operational purposes.

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.

NARA retention schedule are Schedules GRS 20, item 1c and GRS 24 item 6a.

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

All Hard drives will be destroyed following VA and ISO Guidelines when the system is no longer in use.
Electronic data and files of any type, including Protected Health Information (PHI), Sensitive Personal Information (SPI), Human Resources records, and more are destroyed in accordance with the Department of Veterans’ Affairs Handbook 6500.1, Electronic Media Sanitization (November 3, 2008), https://www.vendorportal.ecms.va.gov/FBODocumentServer/DocumentServer.aspx?DocumentId=4214767&FileName=36C25518Q0329-011.pdf 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.

Additionally, James H. Quillen VA Medical Center follows Field Security Service (FSS) Bulletin #176 dated April 9, 2014 for Media Sanitization Program, SOPs - FSS - All Documents 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. 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.

There will not be any research, testing or training being done on the Inspire Physician Programmer.

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:** The risk to maintaining data within the Inspire Physician Programmer is the longer the time frame information is kept, the greater the risk that information possibly will be compromised or breached.

**Mitigation:** To mitigate the risk posed by information retention, the JHQVAMC adheres to the VA RC schedules for each category of data it maintains. When the retention date is reached for a record, the medical center will carefully dispose of the data by the determined method as described in question 3.4.

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**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>
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### 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:** N/A

**Mitigation:** N/A

### 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**

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<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>
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<tr>
<td>N/A</td>
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**5.2 PRIVACY IMPACT ASSESSMENT: External sharing and disclosure**

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

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:** N/A
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.

The JHQVAMC provides notice of information collection in several ways. The initial method of notification is in person during individual interviews or in writing via the Privacy Act statement on forms and applications completed by the individual. The Notice of Privacy Practice (NOPP) [http://vaww.va.gov/vhapublications/ViewPublication.asp?pub_ID=3147](http://vaww.va.gov/vhapublications/ViewPublication.asp?pub_ID=3147) is a document which explains the collection and use of protected information to individuals applying for VHA benefits. A signed statement acknowledging that they individual read and understood the NOPP is scanned into each applicant’s electronic file. When updates are made to the NOPP copies are mailed to all VHA beneficiaries.

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

This question is directed at whether the person from or about whom information is collected can decline to provide the information and if so, whether a penalty or denial of service is attached. This question is related to privacy control IP-1, Consent, IP-2, Individual Access, and IP-3, Redress.
Yes, individuals do have an opportunity to decline to provide information at any time. No, there is not a penalty or denial of service for declining to provide information.

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

Individuals have a right to deny the use of their health information and/or Individually Identifiable Health Information (IIHI) and for the purpose of research. Individuals can request further limitations on other disclosures. A veteran, guardian or court appointed Power of Attorney can submit a request to the facility Privacy Officer to obtain information. JHQVAMC can approve or deny these requests. However, if the request to provide information is accepted JHQVAMC must conform to the restrictions. Additionally, individuals have the right to consent to particular uses of information.

6.4 PRIVACY IMPACT ASSESSMENT: Notice

Describe the potential risks associated with potentially insufficient notice and what steps, if any, are currently being taken to mitigate those identified risks.

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 an individual may not know that their information is being collected or how it will be used.

Mitigation: JHQVAMC mitigates this risk by ensuring that it provides individuals notice of information collection and notice of the system’s existence through the methods discussed in question 6.1.
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.

When requesting access to one’s own records, patients are asked to complete VA Form 10-5345a: Individuals’ Request for a Copy of their Own Health Information, which can be obtained from the medical center or online at https://www.va.gov/find-forms/about-form-10-5345a . Additionally, veterans and their dependents can gain access to their Electronic Health Record (EHR) by enrolling in the myHealthevet program, VA’s online personal health record. More information about myHealthevet at https://www.myhealth.va.gov/index.html. In addition to the procedures discussed above, the SORNs listed in question 6.1 each address record access, redress, and correction. Links to all VA SORNs can be found at http://www.rms.oit.va.gov/sor_records.asp.

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.

VA Form Request to Amend Record will be filled out for correction of patient record.

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.

Notice provided to patient by business office and turned into the privacy officer. The SORNs listed in question 6.1 each discuss and notify members of the public of the procedures related to record access, redress, and correction. Links to all VA SORNs can be found at [http://vaww.oprm.va.gov/privacy/systems_of_records.aspx](http://vaww.oprm.va.gov/privacy/systems_of_records.aspx). Individuals may request correction of their information by contacting a Medical Support Assistant, the Chief of HIMS, the Patient Advocate and or the Release of Information Office (ROI). Individuals are provided verbal notice of amendment process by the PO and/or Health Information Management Systems (HIMS) Chief at time of request.

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.

The Privacy Officer provides appeal rights to the Office of General Counsel or VHA Privacy Office via the written response to the Veteran regarding the outcome of the amendment request.

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:** Risk to patient if incorrect data has been entered.

**Mitigation:** JHQVAMC ensures the patient’s record is correct by checking information prior to each appointment, and at each appointment. Individuals can gain access to their information based on the procedures above and correct any inaccurate or erroneous information.

**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.

Only Sleep providers will have access to the Inspire Physician Programmer.

**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.
No, there is no need for contractors to access the device.

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 are required Privacy and Information Security Awareness training to be completed on an annual basis. The Talent Management System offers the following applicable privacy courses: VA 10176: Privacy and Information Security Awareness and Rules of Behavior; Privacy and HIPPA Focused Training VA 10203; Annual Government Ethics 3812493A 3812493.

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

If Yes, provide:

1. The Security Plan Status,
2. The Security Plan Status Date,
3. The Authorization Status,
4. The Authorization Date,
5. The Authorization Termination Date,
6. The Risk Review Completion Date,
7. 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.

The Inspire sleep system has no connectivity. As a medical device, it will be a minor application and will fall under the MOU-MDIA ATO in Emass.

1. The Security Plan Status, Approved
2. The Security Plan Status Date, 22-Feb-2022
3. The Authorization Status, ATO
4. The Authorization Date, 22 April 2022
5. The Authorization Termination Date, 22 April 2023
6. The Risk Review Completion Date, 11 April 2022
7. The FIPS 199 classification of the system (LOW/MODERATE/HIGH), LOW
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).

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.

N/A

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)

This question is related to privacy control AR-3, Privacy Requirements for Contractors and Service Providers.

N/A

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.

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

What are the roles and responsibilities involved between the organization and cloud provider, particularly with respect to managing risks and ensuring organizational requirements are met?

This question is related to privacy control AR-3, Privacy Requirements for Contractors and Service Providers.

N/A

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).

Inspire Physician Programmer uses short-range radio frequency telemetry to communicate with the Implantable Pulse Generator (IPG) which automatically pulls the serial number, turns on the IPG and change the settings to the IPG.
### 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.

**DEBORAH WEBB**
Digitally signed by DEBORAH WEBB
Date: 2022.06.16 15:16:58 -04'00'

Privacy Officer, Deborah Webb

**CLYDE D. JOHNSON 3470296**
Digitally signed by CLYDE D. JOHNSON 3470296
Date: 2022.06.16 14:48:16 -04'00'

Information Systems Security Officer, Clyde Johnson

**ADIL WARSY**
Digitally signed by ADIL WARSY
Date: 2022.06.16 16:07:25 -04'00'

Information Systems Owner, Adil Warsy
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).

https://vaww.va.gov/vhapublications/ViewPublication.asp?pub_ID=3147