PRIVACY IMPACT ASSESSMENT (PIA)

PRESCRIBING AUTHORITY: DoD Instruction 5400.16, "DoD Privacy Impact Assessment (PIA) Guidance". Complete this form for Department of Defense (DoD) information systems or electronic collections of information (referred to as an "electronic collection" for the purpose of this form) that collect, maintain, use, and/or disseminate personally identifiable information (PII) about members of the public, Federal employees, contractors, or foreign nationals employed at U.S. military facilities internationally. In the case where no PII is collected, the PIA will serve as a conclusive determination that privacy requirements do not apply to system.

1. DOD INFORMATION SYSTEM/ELECTRONIC COLLECTION NAME:

KARL STORZ IOR

2. DOD COMPONENT NAME:

Defense Health Agency

3. PIA APPROVAL DATE:

07/24/2025

PMO - Integrated Clinical Systems (CAE)

SECTION 1: PII DESCRIPTION SUMMARY (FOR PUBLIC RELEASE)

a. The PII is: *(Check one. Note: Federal contractors, military family members, and foreign nationals are included in general public.)*

- ☐ From members of the general public
- ☐ From Federal employees
- ☒ from both members of the general public and Federal employees
- ☐ Not Collected *(if checked proceed to Section 4)*

b. The PII is in a: *(Check one.)*

- ☐ New DoD Information System
- ☐ New Electronic Collection
- ☒ Existing DoD Information System
- ☐ Existing Electronic Collection
- ☐ Significantly Modified DoD Information System

c. Describe the purpose of this DoD information system or electronic collection and describe the types of personal information about individuals collected in the system.

The KARL STORZ Integrated Operating Room (IOR) System allows video, still image or required data from an imaging device, camera, navigation system or computer display to be centrally controlled and displayed on monitors or display consoles within the OR Suite or dedicated remote device. The IOR is a suite of products:
1. The KARL STORZ AIDA BELLA WD 300 a dedicated appliance (consisting of hard- and software) intended for documentation of audio-visual and patient data during a diagnostic or therapeutic procedure. It allows for the capture and annotation of the surgical procedure. Audio-visual data recorded and distributed by the KARL STORZ AIDA BELLA WD 300 is not intended for diagnostic or therapeutic purposes. This system is physically located within the IOR and does not interface with the MTF network.
2. The NEO IP system enables integrated video signals to be centrally routed to the connected displays and video devices. NEO IP enables functions such as control of room cameras, video conferencing systems, lighting, telephone, etc., using a central touch panel. An android tablet (NEO IP Android) provides GUI interface for the Neo IP Controller (NEO IP Ubuntu). This system is physically located within the IOR and does not interface with the MTF network, but allows hospital owned and managed devices to receive video data via HDMI cable.
3. StreamConnect NEO is a Web based solution providing Live Streaming from room cameras located in the OR to any networked PC over a browser. This system interfaces with the MTF network and can become a member server of the site domain.
4. IOR hardware includes cameras, video decoders, video encoders, endoflator, and monitors. These devices/sub-components do not interface with the MTF network, and are physically located within the IOR. The KARL STORZ IOR system is indirectly connected to the GIG via the StreamConnect server.

Personally Identifiable Information (PII) collected includes: demographic and medical information. The categories of individuals with records in this system include: Military members of the Armed Forces, Military retirees, and their family members; DoD Civilian employees; Foreign Nationals; members of the US Coast Guard and Public Health Service; military academy cadets and midshipmen; and other individuals who receive medical treatment at DoD treatment facilities/activities.

d. Why is the PII collected and/or what is the intended use of the PII? *(e.g., verification, identification, authentication, data matching, mission-related use, administrative use)*

The PII collected is for mission-related purposes to support the delivery of health care services, match an individual with his/her medical diagnostic reports and to ensure accuracy when these reports are integrated in the medical records for that individual.

e. Do individuals have the opportunity to object to the collection of their PII? ☐ Yes ☒ No

(1) If "Yes," describe the method by which individuals can object to the collection of PII.

(2) If "No," state the reason why individuals cannot object to the collection of PII.

The PII is obtained from an existing DoD information system or electronic collection.

f. Do individuals have the opportunity to consent to the specific uses of their PII? ☐ Yes ☒ No

(1) If "Yes," describe the method by which individuals can give or withhold their consent.

(2) If "No," state the reason why individuals cannot give or withhold their consent.

The PII is obtained from an existing DoD information system or electronic collection.

g. When an individual is asked to provide PII, a Privacy Act Statement (PAS) and/or a Privacy Advisory must be provided. *(Check as appropriate and provide the actual wording.)*

☐ Privacy Act Statement ☐ Privacy Advisory ☒ Not Applicable

The PII is obtained from an existing DoD information system or electronic collection.

h. With whom will the PII be shared through data/system exchange, both within your DoD Component and outside your Component? *(Check all that apply)*

☒ Within the DoD Component	Specify.	DHA MTF health care providers
☒ Other DoD Components *(i.e. Army, Navy, Air Force)*	Specify.	Navy and Air Force MTFs
☒ Other Federal Agencies *(i.e. Veteran's Affairs, Energy, State)*	Specify.	US Coast Guard, Veterans Administration, Public Health Service, Center for Disease Control
☐ State and Local Agencies	Specify.	
☒ Contractor *(Name of contractor and describe the language in the contract that safeguards PII. Include whether FAR privacy clauses, i.e., 52.224-1, Privacy Act Notification, 52.224-2, Privacy Act, and FAR 39.105 are included in the contract.)*	Specify.	The Manufacturer servicing the device may have access to some data. There may also be contractor radiologists providing radiology support who will need direct access to patient studies. Contracts for Manufacturers and radiologists accessing this device include a standard Military Health System (MHS), Health Insurance Portability and Accountability Act (HIPAA), Business Associate Agreement, DoD/HIPAA guidelines, and MEDCOM Information Assurance (IA) guidelines.
☐ Other *(e.g., commercial providers, colleges).*	Specify.	

i. Source of the PII collected is: *(Check all that apply and list all information systems if applicable)*

☐ Individuals ☐ Databases

☒ Existing DoD Information Systems ☐ Commercial Systems

☐ Other Federal Information Systems

j. How will the information be collected? *(Check all that apply and list all Official Form Numbers if applicable)*

☐ E-mail ☐ Official Form (*Enter Form Number(s) in the box below)*

☐ In-Person Contact ☐ Paper

☐ Fax ☐ Telephone Interview

☒ Information Sharing - System to System ☐ Website/E-Form

☐ Other *(If Other, enter the information in the box below)*

k. Does this DoD Information system or electronic collection require a Privacy Act System of Records Notice (SORN)?

A Privacy Act SORN is required if the information system or electronic collection contains information about U.S. citizens or lawful permanent U.S. residents that is retrieved by name or other unique identifier. PIA and Privacy Act SORN information must be consistent.

☐ Yes ☒ No

If "Yes," enter SORN System Identifier

SORN Identifier, not the Federal Register (FR) Citation. Consult the DoD Component Privacy Office for additional information or http://dpcld.defense.gov/Privacy/SORNs/

or

If a SORN has not yet been published in the Federal Register, enter date of submission for approval to Defense Privacy, Civil Liberties, and Transparency Division (DPCLTD). Consult the DoD Component Privacy Office for this date.

If "No," explain why the SORN is not required in accordance with DoD Regulation 5400.11-R: Department of Defense Privacy Program.

l. What is the National Archives and Records Administration (NARA) approved, pending or general records schedule (GRS) disposition authority for the system or for the records maintained in the system?

(1) NARA Job Number or General Records Schedule Authority. GRS 5.2, item 020 (DAA-GRS-2022-0009-0002)

(2) If pending, provide the date the SF-115 was submitted to NARA.

(3) Retention Instructions.

FILE NUMBER: 103-14
FILE TITLE: Intermediary Records
DISPOSITION: Temporary. Cut off and destroy upon creation or update of the final record, or when no longer needed for business use, whichever is later.

m. What is the authority to collect information? A Federal law or Executive Order must authorize the collection and maintenance of a system of records. For PII not collected or maintained in a system of records, the collection or maintenance of the PII must be necessary to discharge the requirements of a statue or Executive Order.

(1) If this system has a Privacy Act SORN, the authorities in this PIA and the existing Privacy Act SORN should be similar.
(2) If a SORN does not apply, cite the authority for this DoD information system or electronic collection to collect, use, maintain and/or disseminate PII. (If multiple authorities are cited, provide all that apply).

(a) Cite the specific provisions of the statute and/or EO that authorizes the operation of the system and the collection of PII.

(b) If direct statutory authority or an Executive Order does not exist, indirect statutory authority may be cited if the authority requires the operation or administration of a program, the execution of which will require the collection and maintenance of a system of records.

(c) If direct or indirect authority does not exist, DoD Components can use their general statutory grants of authority ("internal housekeeping") as the primary authority. The requirement, directive, or instruction implementing the statute within the DoD Component must be identified.

10 USC 3013, Secretary of the Army; 10 USC 1071-1085, Medical and Dental Care; 50 USC Supplement IV, Appendix 454, as amended, Persons liable for Training and Service; 42 USC Chapter 117 Sections 11131-152, Reporting of Information; 10 USC 10997a and 1097b, Tricare Prime and Tricare Program; 10 USC 1079, Contracts for Medical Care for Spouses and Children; 10 USC 1079a, CHAMPUS; 10 USC 1086, Contracts for Health Benefits for Certain Members, Former Members, and Their Dependents; DoDI 6015.23, Delivery of Healthcare at Military Treatment Facilities (MTFs); DoDD 6040.37, Confidentiality of Medical Quality Assurance (QA) Records; DoD 6010.8-R, Civilian Health and Medical Program of the Uniformed Services (CHAMPUS)

n. Does this DoD information system or electronic collection have an active and approved Office of Management and Budget (OMB) Control Number?

Contact the Component Information Management Control Officer or DoD Clearance Officer for this information. This number indicates OMB approval to collect data from 10 or more members of the public in a 12-month period regardless of form or format.

☐ Yes ☒ No ☐ Pending

(1) If "Yes," list all applicable OMB Control Numbers, collection titles, and expiration dates.
(2) If "No," explain why OMB approval is not required in accordance with DoD Manual 8910.01, Volume 2, " DoD Information Collections Manual: Procedures for DoD Public Information Collections."
(3) If "Pending," provide the date for the 60 and/or 30 day notice and the Federal Register citation.

The information collected in this system is for the diagnosis and treatment of medical conditions and not considered a public information collection in accordance with DoDM 8910.01, Volume 2, Enclosure 3, paragraph 8b(5).