

Manufacturer Disclosure Statement for Medical Device Security -- MDS2

Hologic, Inc. Faxitron Specimen Imaging RD-04179 Rev 001 30-Apr-2021

Question ID	Question	See note	IEC TR 80001-2-2:2012	NIST SP 800-53 Rev. 4	ISO 27002:2013
DOC-1	Manufacturer Name	Hologic, Inc.			
DOC-2	Device Description	Specimen Radiography System			
DOC-3	Device Model	Faxitron Path+; PathVisionXL			
DOC-4	Document ID	RD-04179 Rev 001			
DOC-5	Manufacturer Contact Information	Steve Bolduc steven.bolduc@Hologic.com The Faxitron Faxitron Path+; PathVisionXL Systems are a specimen imaging device. The system is able to capture images and perform procedures with no network connectivity. However it is typically connected to a network to achieve query/retrieve, archiving, printing, interfacing with a RIS, etc.			
DOC-6	Intended use of device in network-connected environment:	4/30/2021			
DOC-7	Document Release Date	4/30/2021			
DOC-8	Coordinated Vulnerability Disclosure: Does the manufacturer have a vulnerability disclosure program for this device?	No			
DOC-9	ISAO: Is the manufacturer part of an Information Sharing and Analysis Organization?	No			
DOC-10	Diagram: Is a network or data flow diagram available that indicates connections to other system components or expected external resources?	Yes, available upon request.			
DOC-11	SaMD: Is the device Software as a Medical Device (i.e. software-only, no hardware)?	No			
DOC-11.1	Does the SaMD contain an operating system?	N/A			
DOC-11.2	Does the SaMD rely on an owner/operator provided operating system?	N/A			
DOC-11.3	Is the SaMD hosted by the manufacturer?	N/A			
DOC-11.4	Is the SaMD hosted by the customer?	N/A			
		Yes, No, N/A, or See Notes			
	MANAGEMENT OF PERSONALLY IDENTIFIABLE INFORMATION				
MPII-1	Can this device display, transmit, store, or modify personally identifiable information (e.g. electronic Protected Health Information (ePHI))?	Yes		AR-2	A.15.1.4
MPII-2	Does the device maintain personally identifiable information?	Yes		AR-2	A.15.1.4
MPII-2.1	Does the device maintain personally identifiable information temporarily in volatile memory (i.e., until cleared by power-off or reset)?	Yes		AR-2	A.15.1.4
MPII-2.2	Does the device store personally identifiable information persistently on internal media?	Yes			
MPII-2.3	Is personally identifiable information preserved in the device's non-volatile memory until explicitly erased?	Yes			
MPII-2.4	Does the device store personally identifiable information in a database?	Yes			
MPII-2.5	Does the device allow configuration to automatically delete local personally identifiable information after it is stored to a long term solution?	No		AR-2	A.15.1.4

MPII-2.6	Does the device import/export personally identifiable information with other systems (e.g., a wearable monitoring device might export personally identifiable information to a server)?	Yes	—	AR-2	A.15.1.4
MPII-2.7	Does the device maintain personally identifiable information when powered off, or during power service interruptions?	Yes	—	AR-2	A.15.1.4
MPII-2.8	Does the device allow the internal media to be removed by a service technician (e.g., for separate destruction or customer retention)?	Yes	—		
MPII-2.9	Does the device allow personally identifiable information records be stored in a separate location from the device's operating system (i.e. secondary internal drive, alternate drive partition, or remote storage location)?	Yes	—	AR-2	A.15.1.4
MPII-3	Does the device have mechanisms used for the transmitting, importing/exporting of personally identifiable information?	Yes	—	AR-2	A.15.1.4
MPII-3.1	Does the device display personally identifiable information (e.g., video display, etc.)?	Yes	—	AR-2	A.15.1.4
MPII-3.2	Does the device generate hardcopy reports or images containing personally identifiable information?	Yes	Note 2	AR-2	A.15.1.4
MPII-3.3	Does the device retrieve personally identifiable information from or record personally identifiable information to removable media (e.g., removable-HDD, USB memory, DVD-R/RW, CD-R/RW, tape, CF/SD card, memory stick, etc.)?	Yes	Note 4	AR-2	A.15.1.4
MPII-3.4	Does the device transmit/receive personally identifiable information via dedicated cable connection (e.g., RS-232, RS-423, USB, FireWire, etc.)?	Yes		AR-2	A.15.1.4
MPII-3.5	Does the device transmit/receive personally identifiable information via a wired network connection (e.g., RJ45, fiber optic, etc.)?	Yes	Note 6	AR-2	A.15.1.4
MPII-3.6	Does the device transmit/receive personally identifiable information via a wireless network connection (e.g., WiFi, Bluetooth, NFC, infrared, cellular, etc.)?	No		AR-2	A.15.1.4
MPII-3.7	Does the device transmit/receive personally identifiable information over an external network (e.g., Internet)?	No		AR-2	A.15.1.4
MPII-3.8	Does the device import personally identifiable information via scanning a document?	No			
MPII-3.9	Does the device transmit/receive personally identifiable information via a proprietary protocol?	No	—		
MPII-3.10	Does the device use any other mechanism to transmit, import or export personally identifiable information?	No	—	AR-2	A.15.1.4
Management of Private Data notes:				AR-2	A.15.1.4

AUTOMATIC LOGOFF (ALOF)

The device's ability to prevent access and misuse by unauthorized users if device is left idle for a period of time.

ALOF-1	Can the device be configured to force reauthorization of logged-in user(s) after a predetermined length of inactivity (e.g., auto-logoff, session lock, password protected screen saver)?	Yes	Note 7	Section 5.1, ALOF	AC-12	None
ALOF-2	Is the length of inactivity time before auto-logoff/screen lock user or administrator configurable?	Yes	Note 7	Section 5.1, ALOF	AC-11	A.11.2.8, A.11.2.9

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AUDIT CONTROLS (AUDT)

The ability to reliably audit activity on the device.

AUDT-1	Can the medical device create additional audit logs or reports beyond standard operating system logs?	No	---
AUDT-1.1	Does the audit log record a USER ID?	Yes	---
AUDT-1.2	Does other personally identifiable information exist in the audit trail?	No	---
AUDT-2	Are events recorded in an audit log? If yes, indicate which of the following events are recorded in the audit log:	Yes	---
AUDT-2.1	Successful login/logout attempts?	Yes	---
AUDT-2.2	Unsuccessful login/logout attempts?	Yes	---
AUDT-2.3	Modification of user privileges?	Yes	---
AUDT-2.4	Creation/modification/deletion of users?	Yes	---
AUDT-2.5	Presentation of clinical or PII data (e.g. display, print)?	No	---
AUDT-2.6	Creation/modification/deletion of data?	No	---
AUDT-2.7	Import/export of data from removable media (e.g. USB drive, external hard drive, DVD)?	No	---
AUDT-2.8	Receipt/transmission of data or commands over a network or point-to-point connection?	No	---
AUDT-2.8.1	Remote or on-site support?	No	---
AUDT-2.8.2	Application Programming Interface (API) and similar activity?	No	---
AUDT-2.9	Emergency access?	N/A	---
AUDT-2.10	Other events (e.g., software updates)?	No	---
AUDT-2.11	Is the audit capability documented in more detail?	No	---
AUDT-3	Can the owner/operator define or select which events are recorded in the audit log?	No	---
AUDT-4	Is a list of data attributes that are captured in the audit log for an event available?	Yes	---
AUDT-4.1	Does the audit log record date/time?	Yes	---
AUDT-4.1.1	Can date and time be synchronized by Network Time Protocol (NTP) or equivalent time source?	Yes	---
AUDT-5	Can audit log content be exported?	Yes	---
AUDT-5.1	Via physical media?	Yes	---
AUDT-5.2	Via IHE Audit Trail and Node Authentication (ATNA) profile to SIEM?	No	---
AUDT-5.3	Via Other communications (e.g., external service device, mobile applications)?	No	---
AUDT-5.4	Are audit logs encrypted in transit or on storage media?	No	---
AUDT-6	Can audit logs be monitored/reviewed by owner/operator?	Yes	---
AUDT-7	Are audit logs protected from modification?	No	---
AUDT-7.1	Are audit logs protected from access?	No	---
AUDT-8	Can audit logs be analyzed by the device?	No	---

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Section 5.2, AUDT	AU-1	A.5.1.1, A.5.1.2, A.6.1.1, A.12.1.1, A.18.1.1, A.18.2.2
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
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Section 5.2, AUDT	AU-2	None
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Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None
Section 5.2, AUDT	AU-2	None

AUTHORIZATION (AUTH)

The ability of the device to determine the authorization of users.

AUTH-1	Does the device prevent access to unauthorized users through user login requirements or other mechanism?	Yes	---
AUTH-1.1	Can the device be configured to use federated credentials management of users for authorization (e.g., LDAP, OAuth)?	Yes	Active Directory

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Section 5.3, AUTH	IA-2	A.9.2.1
Section 5.3, AUTH	IA-2	A.9.2.1

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AUTH-1.2	Can the customer push group policies to the device (e.g., Active Directory)?	See Notes	Note 8	Section 5.3, AUTH	IA-2	A.9.2.1
AUTH-1.3	Are any special groups, organizational units, or group policies required?	Yes	Note 9	Section 5.3, AUTH	IA-2	A.9.2.1
AUTH-2	Can users be assigned different privilege levels based on 'role' (e.g., user, administrator, and/or service, etc.)?	No	—	Section 5.3, AUTH	IA-2	A.9.2.1
AUTH-3	Can the device owner/operator grant themselves unrestricted administrative privileges (e.g., access operating system or application via local root or administrator account)?	Yes	—	Section 5.3, AUTH	IA-2	A.9.2.1
AUTH-4	Does the device authorize or control all API access requests?	No	—	Section 5.3, AUTH	IA-2	A.9.2.1
AUTH-5	Does the device run in a restricted access mode, or 'kiosk mode', by default?	No	—			

CYBER SECURITY PRODUCT UPGRADES (CSUP)

The ability of on-site service staff, remote service staff, or authorized customer staff to install/upgrade device's security patches.

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CSUP-1	Does the device contain any software or firmware which may require security updates during its operational life, either from the device manufacturer or from a third-party manufacturer of the software/firmware? If no, answer "N/A" to questions in this section.	Yes	Windows			
CSUP-2	Does the device contain an Operating System? If yes, complete 2.1-2.4.	Yes	—			
CSUP-2.1	Does the device documentation provide instructions for owner/operator installation of patches or software updates?	No				
CSUP-2.2	Does the device require vendor or vendor-authorized service to install patches or software updates?	Yes	—			
CSUP-2.3	Does the device have the capability to receive remote installation of patches or software updates?	Yes				
CSUP-2.4	Does the medical device manufacturer allow security updates from any third-party manufacturers (e.g., Microsoft) to be installed without approval from the manufacturer?	No				
CSUP-3	Does the device contain Drivers and Firmware? If yes, complete 3.1-3.4.	Yes	—			
CSUP-3.1	Does the device documentation provide instructions for owner/operator installation of patches or software updates?	No	—			
CSUP-3.2	Does the device require vendor or vendor-authorized service to install patches or software updates?	Yes	—			
CSUP-3.3	Does the device have the capability to receive remote installation of patches or software updates?	No	—			
CSUP-3.4	Does the medical device manufacturer allow security updates from any third-party manufacturers (e.g., Microsoft) to be installed without approval from the manufacturer?	No	—			
CSUP-4	Does the device contain Anti-Malware Software? If yes, complete 4.1-4.4.	Yes				
CSUP-4.1	Does the device documentation provide instructions for owner/operator installation of patches or software updates?	Yes	Note 10			
CSUP-4.2	Does the device require vendor or vendor-authorized service to install patches or software updates?	See Notes	Note 10			

CSUP-4.3	Does the device have the capability to receive remote installation of patches or software updates?	Yes	Note 10			
CSUP-4.4	Does the medical device manufacturer allow security updates from any third-party manufacturers (e.g., Microsoft) to be installed without approval from the manufacturer?	See Notes	Note 10			
CSUP-5	Does the device contain Non-Operating System commercial off-the-shelf components? If yes, complete 5.1-5.4.	Yes	—			
CSUP-5.1	Does the device documentation provide instructions for owner/operator installation of patches or software updates?	No	—			
CSUP-5.2	Does the device require vendor or vendor-authorized service to install patches or software updates?	Yes	—			
CSUP-5.3	Does the device have the capability to receive remote installation of patches or software updates?	No	—			
CSUP-5.4	Does the medical device manufacturer allow security updates from any third-party manufacturers (e.g., Microsoft) to be installed without approval from the manufacturer?	No	—			
CSUP-6	Does the device contain other software components (e.g., asset management software, license management)? If yes, please provide details or reference in notes and complete 6.1-6.4.	No	—			
CSUP-6.1	Does the device documentation provide instructions for owner/operator installation of patches or software updates?	No	—			
CSUP-6.2	Does the device require vendor or vendor-authorized service to install patches or software updates?	No	—			
CSUP-6.3	Does the device have the capability to receive remote installation of patches or software updates?	No	—			
CSUP-6.4	Does the medical device manufacturer allow security updates from any third-party manufacturers (e.g., Microsoft) to be installed without approval from the manufacturer?	No	—			
CSUP-7	Does the manufacturer notify the customer when updates are approved for installation?	Yes	Note 11			
CSUP-8	Does the device perform automatic installation of software updates?	No	—			
CSUP-9	Does the manufacturer have an approved list of third-party software that can be installed on the device?	Yes	Note 10			
CSUP-10	Can the owner/operator install manufacturer-approved third-party software on the device themselves?	Yes	Note 10			
CSUP-10.1	Does the system have mechanism in place to prevent installation of unapproved software?	No	—			
CSUP-11	Does the manufacturer have a process in place to assess device vulnerabilities and updates?	Yes	Note 12			
CSUP-11.1	Does the manufacturer provide customers with review and approval status of updates?	Yes	Note 11			
CSUP-11.2	Is there an update review cycle for the device?	Yes	Note 13			

HEALTH DATA DE-IDENTIFICATION (DIDT)

The ability of the device to directly remove information that allows identification of a person.

DIDT-1	Does the device provide an integral capability to de-identify personally identifiable information?	No	—
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Section 5.6, DIDT

None

ISO 27038

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DIDT-1.1	Does the device support de-identification profiles that comply with the DICOM standard for de-identification?	No	—	Section 5.6, DIDT	None	ISO 27038
DATA BACKUP AND DISASTER RECOVERY (DTBK)				IEC TR 80001-2-2:2012	NIST SP 800-53 Rev. 4	ISO 27002:2013
<i>The ability to recover after damage or destruction of device data, hardware, software, or site configuration information.</i>						
DTBK-1	Does the device maintain long term primary storage of personally identifiable information / patient information (e.g. PACS)?	No	—			
DTBK-2	Does the device have a “factory reset” function to restore the original device settings as provided by the manufacturer?	See Notes	Note 3	Section 5.7, DTBK	CP-9	A.12.3.1
DTBK-3	Does the device have an integral data backup capability to removable media?	Yes	Note 14	Section 5.7, DTBK	CP-9	A.12.3.1
DTBK-4	Does the device have an integral data backup capability to remote storage?	Yes	Note 14			
DTBK-5	Does the device have a backup capability for system configuration information, patch restoration, and software restoration?	Yes	Note 14			
DTBK-6	Does the device provide the capability to check the integrity and authenticity of a backup?	No	—	Section 5.7, DTBK	CP-9	A.12.3.1
EMERGENCY ACCESS (EMRG)				IEC TR 80001-2-2:2012	NIST SP 800-53 Rev. 4	ISO 27002:2013
<i>The ability of the device user to access personally identifiable information in case of a medical emergency situation that requires immediate access to stored personally identifiable information.</i>						
EMRG-1	Does the device incorporate an emergency access (i.e. “break-glass”) feature?	No	—	Section 5.8, EMRG	SI-17	None
HEALTH DATA INTEGRITY AND AUTHENTICITY (IGAU)				IEC TR 80001-2-2:2012	NIST SP 800-53 Rev. 4	ISO 27002:2013
<i>How the device ensures that the stored data on the device has not been altered or destroyed in a non-authorized manner and is from the originator.</i>						
IGAU-1	Does the device provide data integrity checking mechanisms of stored health data (e.g., hash or digital signature)?	No		Section 5.9, IGAU	SC-28	A.18.1.3
IGAU-2	Does the device provide error/failure protection and recovery mechanisms for stored health data (e.g., RAID-5)?	No	Note 15	Section 5.9, IGAU	SC-28	A.18.1.3
MALWARE DETECTION/PROTECTION (MLDP)				IEC TR 80001-2-2:2012	NIST SP 800-53 Rev. 4	ISO 27002:2013
<i>The ability of the device to effectively prevent, detect and remove malicious software (malware).</i>						
MLDP-1	Is the device capable of hosting executable software?	Yes	—	Section 5.10, MLDP		
MLDP-2	Does the device support the use of anti-malware software (or other anti-malware mechanism)?	Yes	Note 10	Section 5.10, MLDP	SI-3	A.12.2.1
MLDP-2.1	Does the device include anti-malware software by default?	Yes		Section 5.10, MLDP	CM-5	A.9.2.3, A.9.4.5, A.12.1.2, A.12.1.4, A.12.5.1
MLDP-2.2	Does the device have anti-malware software available as an option?	Yes		Section 5.10, MLDP	AU-6	A.12.4.1, A.16.1.2, A.16.1.4

MLDP-2.3	Does the device documentation allow the owner/operator to install or update anti-malware software?	Yes		Section 5.10, MLDP	CP-10	A.17.1.2
MLDP-2.4	Can the device owner/operator independently (re-)configure anti-malware settings?	Yes		Section 5.10, MLDP	AU-2	None
MLDP-2.5	Does notification of malware detection occur in the device user interface?	Yes				
MLDP-2.6	Can only manufacturer-authorized persons repair systems when malware has been detected?	Yes				
MLDP-2.7	Are malware notifications written to a log?	Yes				
MLDP-2.8	Are there any restrictions on anti-malware (e.g., purchase, installation, configuration, scheduling)?	No				
MLDP-3	If the answer to MLDP-2 is NO, and anti-malware cannot be installed on the device, are other compensating controls in place or available?	N/A	---	Section 5.10, MLDP	SI-2	A.12.6.1, A.14.2.2, A.14.2.3, A.16.1.3
MLDP-4	Does the device employ application whitelisting that restricts the software and services that are permitted to be run on the device?	No	---	Section 5.10, MLDP	SI-3	A.12.2.1
MLDP-5	Does the device employ a host-based intrusion detection/prevention system?	No	---	Section 5.10, MLDP	SI-4	None
MLDP-5.1	Can the host-based intrusion detection/prevention system be configured by the customer?	Yes	---	Section 5.10, MLDP	CM-7	A.12.5.1
MLDP-5.2	Can a host-based intrusion detection/prevention system be installed by the customer?	Yes	---	Section 5.10, MLDP		

NODE AUTHENTICATION (NAUT)

The ability of the device to authenticate communication partners/nodes.

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NAUT-1	Does the device provide/support any means of node authentication that assures both the sender and the recipient of data are known to each other and are authorized to receive transferred information (e.g. Web APIs, SMTP, SNMP)?	Yes		Section 5.11, NAUT	SC-23	None
NAUT-2	Are network access control mechanisms supported (E.g., does the device have an internal firewall, or use a network connection white list)?	Yes	Note 16	Section 5.11, NAUT	SC-7	A.13.1.1, A.13.1.3, A.13.2.1, A.14.1.3
NAUT-2.1	Is the firewall ruleset documented and available for review?	Yes	Available upon request			
NAUT-3	Does the device use certificate-based network connection authentication?	No				

CONNECTIVITY CAPABILITIES (CONN)

All network and removable media connections must be considered in determining appropriate security controls. This section lists connectivity capabilities that may be present on the device.

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CONN-1	Does the device have hardware connectivity capabilities?	Yes	---			
CONN-1.1	Does the device support wireless connections?	No	---			
CONN-1.1.1	Does the device support Wi-Fi?	No	---			
CONN-1.1.2	Does the device support Bluetooth?	No	---			
CONN-1.1.3	Does the device support other wireless network connectivity (e.g. LTE, Zigbee, proprietary)?	No	---			
CONN-1.1.4	Does the device support other wireless connections (e.g., custom RF controls, wireless detectors)?	No	---			
CONN-1.2	Does the device support physical connections?	Yes	---			
CONN-1.2.1	Does the device have available RJ45 Ethernet ports?	Yes	---			
CONN-1.2.2	Does the device have available USB ports?	Yes	---			

CONN-1.2.3	Does the device require, use, or support removable memory devices?	Yes	Note 5
CONN-1.2.4	Does the device support other physical connectivity?	No	
	Does the manufacturer provide a list of network ports and protocols that are used or may be used on the device?	Yes	Available upon request.
CONN-2	Can the device communicate with other systems within the customer environment?	Yes	—
CONN-3	Can the device communicate with other systems external to the customer environment (e.g., a service host)?	Yes	—
CONN-4	Does the device make or receive API calls?	No	—
CONN-5	Does the device require an internet connection for its intended use?	No	—
CONN-6	Does the device support Transport Layer Security (TLS)?	Yes	Note 17
CONN-7	Is TLS configurable?	Yes	Note 17
CONN-7.1	Does the device provide operator control functionality from a separate device (e.g., telemedicine)?	No	

PERSON AUTHENTICATION (PAUT)

The ability to configure the device to authenticate users.

PAUT-1	Does the device support and enforce unique IDs and passwords for all users and roles (including service accounts)?	Yes	Note 18
PAUT-1.1	Does the device enforce authentication of unique IDs and passwords for all users and roles (including service accounts)?	Yes	Note 18
PAUT-2	Is the device configurable to authenticate users through an external authentication service (e.g., MS Active Directory, NDS, LDAP, OAuth, etc.)?	Yes	Active Directory
PAUT-3	Is the device configurable to lock out a user after a certain number of unsuccessful logon attempts?	Yes	Note 19
PAUT-4	Are all default accounts (e.g., technician service accounts, administrator accounts) listed in the documentation?	No	
PAUT-5	Can all passwords be changed?	No	—
PAUT-6	Is the device configurable to enforce creation of user account passwords that meet established (organization specific) complexity rules?	Yes	Note 20
PAUT-7	Does the device support account passwords that expire periodically?	Yes	Note 21
PAUT-8	Does the device support multi-factor authentication?	No	—
PAUT-9	Does the device support single sign-on (SSO)?	Yes	Active Directory
PAUT-10	Can user accounts be disabled/locked on the device?	Yes	—
PAUT-11	Does the device support biometric controls?	No	
PAUT-12	Does the device support physical tokens (e.g. badge access)?	No	—
PAUT-13	Does the device support group authentication (e.g. hospital teams)?	Yes	
PAUT-14	Does the application or device store or manage authentication credentials?	Yes	Note 22
PAUT-14.1	Are credentials stored using a secure method?	Yes	Note 22

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Section 5.12, PAUT

IA-2

A.9.2.1

Section 5.12, PAUT

IA-2

A.9.2.1

Section 5.12, PAUT

IA-5

A.9.2.1

Section 5.12, PAUT

IA-2

A.9.2.1

Section 5.12, PAUT

SA-4(5)

A.14.1.1, A.14.2.7, A.14.2.9, A.15.1.2

Section 5.12, PAUT

IA-2

A.9.2.1

Section 5.12, PAUT

IA-2

A.9.2.1

Section 5.12, PAUT

IA-2

A.9.2.1

Section 5.12, PAUT

IA-2

A.9.2.1

Section 5.12, PAUT

IA-2

A.9.2.1

PHYSICAL LOCKS (PLOK)

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Physical locks can prevent unauthorized users with physical access to the device from compromising the integrity and confidentiality of personally identifiable information stored on the device or on removable media

PLOK-1	Is the device software only? If yes, answer "N/A" to remaining questions in this section.	No	---	Section 5.13, PLOK	PE- 3(4)	A.11.1.1, A.11.1.2, A.11.1.3
PLOK-2	Are all device components maintaining personally identifiable information (other than removable media) physically secure (i.e., cannot remove without tools)?	Yes	---	Section 5.13, PLOK	PE- 3(4)	A.11.1.1, A.11.1.2, A.11.1.3
PLOK-3	Are all device components maintaining personally identifiable information (other than removable media) physically secured behind an individually keyed locking device?	No	---	Section 5.13, PLOK	PE- 3(4)	A.11.1.1, A.11.1.2, A.11.1.3
PLOK-4	Does the device have an option for the customer to attach a physical lock to restrict access to removable media?	No	---	Section 5.13, PLOK	PE- 3(4)	A.11.1.1, A.11.1.2, A.11.1.3

ROADMAP FOR THIRD PARTY COMPONENTS IN DEVICE LIFE CYCLE (RDMP)

Manufacturer's plans for security support of third-party components within the device's life cycle.

RDMP-1	Was a secure software development process, such as ISO/IEC 27034 or IEC 62304, followed during product development?	Yes	---	Section 5.14, RDMP	CM-2	None
RDMP-2	Does the manufacturer evaluate third-party applications and software components included in the device for secure development practices?	Yes	---	Section 5.14, RDMP	CM-8	A.8.1.1, A.8.1.2
RDMP-3	Does the manufacturer maintain a web page or other source of information on software support dates and updates?	Yes	---	Section 5.14, RDMP	CM-8	A.8.1.1, A.8.1.2
RDMP-4	Does the manufacturer have a plan for managing third-party component end-of-life?	Yes	---	Section 5.14, RDMP	CM-8	A.8.1.1, A.8.1.2

SOFTWARE BILL OF MATERIALS (SBoM)

A Software Bill of Material (SBoM) lists all the software components that are incorporated into the device being described for the purpose of operational security planning by the healthcare delivery organization. This section supports controls in the RDMP section.

SBOM-1	Is the SBoM for this product available?	Yes	See SBoM sheet within this document.			
SBOM-2	Does the SBoM follow a standard or common method in describing software components?	No	---			
SBOM-2.1	Are the software components identified?	Yes	---			
SBOM-2.2	Are the developers/manufacturers of the software components identified?	Yes	---			
SBOM-2.3	Are the major version numbers of the software components identified?	Yes	---			
SBOM-2.4	Are any additional descriptive elements identified?	Yes	---			
SBOM-3	Does the device include a command or process method available to generate a list of software components installed on the device?	No	---			
SBOM-4	Is there an update process for the SBoM?	Yes	Note 23			

SYSTEM AND APPLICATION HARDENING (SAHD)

IEC TR 80001-2-2:2012	NIST SP 800-53 Rev. 4	ISO 27002:2013
IEC TR 80001-2-2:2012	NIST SP 800-53 Rev. 4	ISO 27002:2013
IEC TR 80001-2-2:2012	NIST SP 800-53 Rev. 4	ISO 27002:2013

The device's inherent resistance to cyber attacks and malware.

SAHD ID	Question	Response	Notes	Section	Control	References
SAHD-1	Is the device hardened in accordance with any industry standards?	No		Section 5.15, SAHD	CM-7 AC-17(2)/IA-3	A.12.5.1* A.6.2.1, A.6.2.2, A.13.1.1, A.13.2.1, A.14.1.2/None
SAHD-2	Has the device received any cybersecurity certifications?	No		Section 5.15, SAHD	SA-12(10)	A.14.2.7, A.15.1.1, A.15.1.2, A.15.1.3
SAHD-3	Does the device employ any mechanisms for software integrity checking?	No	—			
SAHD-3.1	Does the device employ any mechanism (e.g., release-specific hash key, checksums, digital signature, etc.) to ensure the installed software is manufacturer-authorized?	No				
SAHD-3.2	Does the device employ any mechanism (e.g., release-specific hash key, checksums, digital signature, etc.) to ensure the software updates are the manufacturer-authorized updates?	No		Section 5.15, SAHD	CM-8	A.8.1.1, A.8.1.2
SAHD-4	Can the owner/operator perform software integrity checks (i.e., verify that the system has not been modified or tampered with)?	No		Section 5.15, SAHD	AC-3	A.6.2.2, A.9.1.2, A.9.4.1, A.9.4.4, A.9.4.5, A.13.1.1, A.14.1.2, A.14.1.3, A.18.1.3
SAHD-5	Is the system configurable to allow the implementation of file-level, patient level, or other types of access controls?	Yes	Note 24	Section 5.15, SAHD	CM-7	A.12.5.1*
SAHD-5.1	Does the device provide role-based access controls?	Yes	Note 24	Section 5.15, SAHD	CM-7	A.12.5.1*
SAHD-6	Are any system or user accounts restricted or disabled by the manufacturer at system delivery?	No		Section 5.15, SAHD	CM-8	A.8.1.1, A.8.1.2
SAHD-6.1	Are any system or user accounts configurable by the end user after initial configuration?	Yes		Section 5.15, SAHD	CM-7	A.12.5.1*
SAHD-6.2	Does this include restricting certain system or user accounts, such as service technicians, to least privileged access?	No		Section 5.15, SAHD	CM-7	A.12.5.1*
SAHD-7	Are all shared resources (e.g., file shares) which are not required for the intended use of the device disabled?	No	—	Section 5.15, SAHD	CM-7	A.12.5.1*
SAHD-8	Are all communication ports and protocols that are not required for the intended use of the device disabled?	No	—	Section 5.15, SAHD	SA-18	None
SAHD-9	Are all services (e.g., telnet, file transfer protocol [FTP], internet information server [IIS], etc.), which are not required for the intended use of the device deleted/disabled?	No	—	Section 5.15, SAHD	CM-6	None
SAHD-10	Are all applications (COTS applications as well as OS-included applications, e.g., MS Internet Explorer, etc.) which are not required for the intended use of the device deleted/disabled?	No	—	Section 5.15, SAHD	SI-2	A.12.6.1, A.14.2.2, A.14.2.3, A.16.1.3
SAHD-11	Can the device prohibit boot from uncontrolled or removable media (i.e., a source other than an internal drive or memory component)?	No				
SAHD-12	Can unauthorized software or hardware be installed on the device without the use of physical tools?	See Notes	Note 26			
SAHD-13	Does the product documentation include information on operational network security scanning by users?	No	—			
SAHD-14	Can the device be hardened beyond the default provided state?	See Notes	Note 27			
SAHD-14.1	Are instructions available from vendor for increased hardening?	Yes	Available upon request/discussion.			
SHAD-15	Can the system prevent access to BIOS or other bootloaders during boot?	Yes	Note 25			
SAHD-16	Have additional hardening methods not included in 2.3.19 been used to harden the device?	No	—			

SECURITY GUIDANCE (SGUD)

Availability of security guidance for operator and administrator of the device and manufacturer sales and service.

IEC TR 80001-2-2:2012

NIST SP 800-53 Rev. 4

ISO 27002:2013

SGUD-1	Does the device include security documentation for the owner/operator?	Yes	Note 28
SGUD-2	Does the device have the capability, and provide instructions, for the permanent deletion of data from the device or media?	Yes	Note 29
SGUD-3	Are all access accounts documented?	Yes	Available upon request.
SGUD-3.1	Can the owner/operator manage password control for all accounts?	No	---
SGUD-4	Does the product include documentation on recommended compensating controls for the device?	No	---

Section 5.16, SGUD

AT-2/PL-2

A.7.2.2, A.12.2.1/A.14.1.1

Section 5.16, SGUD

MP-6

A.8.2.3, A.8.3.1, A.8.3.2, A.11.2.7

Section 5.16, SGUD

AC-6,IA-2

A.9.1.2, A.9.2.3, A.9.4.4, A.9.4.5/A.9.2.1

HEALTH DATA STORAGE CONFIDENTIALITY (STCF)

The ability of the device to ensure unauthorized access does not compromise the integrity and confidentiality of personally identifiable information stored on the device or removable media.

IEC TR 80001-2-2:2012

NIST SP 800-53 Rev. 4

ISO 27002:2013

STCF-1	Can the device encrypt data at rest?	No	---
STCF-1.1	Is all data encrypted or otherwise protected?	No	---
STCF-1.2	Is the data encryption capability configured by default?	No	---
STCF-1.3	Are instructions available to the customer to configure encryption?	No	---
STCF-2	Can the encryption keys be changed or configured?	No	---
STCF-3	Is the data stored in a database located on the device?	Yes	---
STCF-4	Is the data stored in a database external to the device?	No	---

Section 5.17, STCF

SC-28

A.8.2.3

Section 5.17, STCF

SC-28

A.8.2.3

TRANSMISSION CONFIDENTIALITY (TXCF)

The ability of the device to ensure the confidentiality of transmitted personally identifiable information.

IEC TR 80001-2-2:2012

NIST SP 800-53 Rev. 4

ISO 27002:2013

TXCF-1	transmitted only via a point-to-point dedicated cable?	Yes	---
TXCF-2	Is personally identifiable information encrypted prior to transmission via a network or removable media?	No	---
TXCF-2.1	If data is not encrypted by default, can the customer configure encryption options?	No	---
TXCF-3	Is personally identifiable information transmission restricted to a fixed list of network destinations?	No	---
TXCF-4	Are connections limited to authenticated systems?	No	---
TXCF-5	Are secure transmission methods supported/implemented (DICOM, HL7, IEEE 11073)?	No	---

Section 5.18, TXCF

CM-7

A.12.5.1

Section 5.18, TXCF

CM-7

A.12.5.1

Section 5.18, TXCF

CM-7

A.12.5.1

Section 5.18, TXCF

CM-7

A.12.5.1

TRANSMISSION INTEGRITY (TXIG)

The ability of the device to ensure the integrity of transmitted data.

IEC TR 80001-2-2:2012

NIST SP 800-53 Rev. 4

ISO 27002:2013

TXIG-1	Does the device support any mechanism (e.g., digital signatures) intended to ensure data is not modified during transmission?	No	---
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Section 5.19, TXIG

SC-8

A.8.2.3, A.13.1.1, A.13.2.1, A.13.2.3, A.14.1.2, A.14.1.3

TXIG-2 Does the device include multiple sub-components connected by external cables? No

	REMOTE SERVICE (RMOT)			IEC TR 80001-2-2:2012	NIST SP 800-53 Rev. 4	ISO 27002:2013
	<i>Remote service refers to all kinds of device maintenance activities performed by a service person via network or other remote connection.</i>					
RMOT-1	Does the device permit remote service connections for device analysis or repair?	No	—		AC-17	A.6.2.1, A.6.2.2, A.13.1.1, A.13.2.1, A.14.1.2
RMOT-1.1	Does the device allow the owner/operator to initiate remote service sessions for device analysis or repair?	No	—			
RMOT-1.2	Is there an indicator for an enabled and active remote session?	No	—			
RMOT-1.3	Can patient data be accessed or viewed from the device during the remote session?	No	—		AC-17	A.6.2.1, A.6.2.2, A.13.1.1, A.13.2.1, A.14.1.2
RMOT-2	Does the device permit or use remote service connections for predictive maintenance data?	No	—			
RMOT-3	functionality (e.g. software updates, remote training)?	No	—			

OTHER SECURITY CONSIDERATIONS (OTHR)

NONE

IEC TR 80001-2-2:2012

NIST SP 800-53 Rev. 4

ISO 27002:2013

Notes:

- Note 1 Device contains a limited amount of ePHI to identify images - typically a name, date of birth, patient ID, and accession number.
- Note 2 Optional printing of patient reports
- Note 3 Factory reset requires Service Personnel to perform
- Note 4 Optional importing and exporting of patient procedures.
- Note 5 Backup/Restore
- Note 6 Typically an RJ45 Ethernet connection.
- Note 7 Product application defaults to never logging out current user. Inactivity timeout configurable. It's strongly recommend to limit policy changes pushed to the device to User related policies only, such as password complexity requirements, forcing passwords to expire, etc. There are certain policy changes that, if pushed, could negatively impact the product application.
- Note 8 Strongly recommend configuring the product in its own Organizational Unit and limiting policy changes pushed to the system.
- Note 9 Option available to install validated CoTS antimalware products. See product support website for list of validated antimalware software solutions and installation guidance. Malware definitions can be updated by customer at will. Hologic suggests keeping antimalware software version at the same major version as what was validated.
- Note 10 Validated security patches for the product are posted to the product support website at regular intervals.
- Note 11

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Note 12	Vulnerability assessments, leveraging industry standard tools, and Windows security patch validation occur at regular intervals.		
Note 13	Hologic strives to evaluate and test Windows security updates for the product as they're released (typically monthly).		
Note 14	Software databases and configurations can be backed up. Patient studies should be stored to long term storage or exported to external media by the customer.		
Note 15	Product not designed for long term storage. Patient studies should be stored to long term storage.		
Note 16	Windows Firewall enabled and configured to allow product application network traffic. Patient data only sent to configured DICOM devices.		
Note 17	Hologic Connect leverages an encrypted TLS tunnel for remote Service connectivity. TLS can, optionally, be configured for the product Service Tools configuration web application. External network traffic can also be blocked for Service Tools. Patient study transmission to external devices is done using DICOM, without TLS. Customer may configure TLS at the network layer.		
Note 18	Use of unique product accounts is the decision of the customer. Generic accounts can be removed.		
Note 19	Enabled by default, locking the user for 15 minutes after 3 failed logon attempts.		
Note 20	Not configured by default		
Note 21	Passwords not configured to automatically expire by default. Configurable by customer.		
Note 22	Product application leverages Windows Operating System for user authentication. Credentials not stored in application databases. Credentials stored/managed securely via Operating System.		
Note 23	SBOM reviewed and updated as required during product update cycles.		
Note 24	Product utilizes role-based privileges for many sensitive areas of the application.		
Note 25	Can be configured, not restricted by default. If configured, communicate change to service representative.		
Note 26	Hardware installation would require tools, software would require OS authentication.		
Note 27	Additional hardening or concerns may be discussed with Hologic. Implementing additional hardening changes may negatively impact the product.		
Note 28	Security documentation available on product support website.		
Note 29	Contact customer support for instructions		

Software Bill of Materials (SBoM)

Component Name	Developer	Version(s)	Product Use
Windows 10 IoT Enterprise x64	Microsoft	LTSC 2019	Operating System
BiopticsVision	Hologic Inc	2.4.1.0	Main system application.
BiopticsVisionDXA	Hologic Inc	2.4.2.0	Main DXA application.
BiCam	Hologic Inc	2.2.1.0	Detector interface DLL.
BiopticsProcessing	Hologic Inc	2.3.1.0	Processing DLL.
DXAProcessing	Hologic Inc	1.0.4.0	DXA processing DLL.
ConfigUtility	Hologic Inc	1.0.1.0	Configuration DLL.
MWLQuery	Hologic Inc	N/A	Modality WorkList query dictionary.
PlxApi	PLX Technology, Inc	7.2.0.0	Detector SDK API DLL.
LeadTools	LEAD Technologies, Inc.	15.0.1	Visualization and DICOM integration.
Adobe Reader XI	Adobe Systems Incorporated	11.0.23	Adobe reader.
Intel(R) Rapid Storage Technology	Intel Corporation	16.7.10.1030	Intel drivers.
Intel(R) Graphics Driver	Intel Corporation	24.20.100.6287	Intel drivers.
Intel(R) Management Engine Components	Intel Corporation	1829.12.0.1154	Intel drivers.
Intel(R) Optane™ Pinning Explorer Extensions	Intel Corporation	16.7.10.1030	Intel drivers.
Microsoft Access database engine 2010	Microsoft Corporation	14.0.7015.1000	Access database engine.
Microsoft Visual C++ 2005 Redistributable	Microsoft Corporation	8.0.61000	Visual C++ redistributable.
Microsoft Visual C++ 2008 Redistributable	Microsoft Corporation	9.0.21022	Visual C++ redistributable.
Microsoft Visual C++ 2010 Redistributable	Microsoft Corporation	10.0.30319	Visual C++ redistributable.
Microsoft Visual C++ 2013 Redistributable	Microsoft Corporation	12.0.21005.1	Visual C++ redistributable.
Microsoft Visual C++ 2013 Redistributable	Microsoft Corporation	12.0.30501.0	Visual C++ redistributable.
Realtek High Definition Audio Driver	Realtek Semiconductor Corp.	6.0.1.8573	Audio driver.
Bioptics USB-VCP Port	FTDI	2.12.0.0	USB-VCP port drivers.

Additional Notes

Many of the software components listed above are covered by Hologic's program to regularly validate latest released security patches. See the product support website for the latest validated patches available or contact support for assistance.

Note 1