

NX (Modality Workstation)

Version X.0.8X00

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Manufacturer Disclosure Statement for Device Security

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Information, Security & Privacy related information worldwide	Manufacturer Disclosure Statement for Device Security	http://www.agfa.com/en/he/support/interoperability/mds2/index.jsp
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REVISION RECORD

Revision in Project Workspace	Date	Author	Changes/remarks
1	28 July 2011	J. Gibbs	Initial version
2	29 July 2011	J. Gibbs	Added LLid, update CT members, version number in header
3	29 July 2011	J. Gibbs	Updated CT members

Revision in Product Security Website	Date	Author	Changes/remarks
1.0			Initial version

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FOREWORD

This document consists of the Manufacturer Disclosure Statement for Device Security (MDS² form). The intent of the MDS² form is to supply healthcare providers with important information to assist them in assessing the VULNERABILITY and risks associated with protecting ELECTRONIC PROTECTED HEALTH INFORMATION (ePHI) transmitted or maintained by devices. Because security risk assessment spans an entire organization, this document focuses on only those elements of the security risk assessment process associated with devices and systems that maintain or transmit ePHI.

The MDS² form should:

- (1) Be useful to healthcare provider organizations worldwide. While the form does supply information important to providers who must comply with HIPAA privacy and security rules, the information presented may be useful for any healthcare provider who aspires to have an effective information security RISK MANAGEMENT program. Outside the US, providers would therefore find the MDS² form an effective tool to address regional regulations such as EU 95/46 (Europe), Act on the Protection of Personal Information (Act No. 57 of 2003, Japan), and PIPEDA (Canada).
- (2) Include device specific information addressing the technical security-related attributes of the individual device model.
- (3) Provide a simple, flexible way of collecting the technical, device-specific elements of the common/typical information needed by provider organizations (device users/operators) to begin device information security (i.e., confidentiality, integrity, availability) risk assessments.
- (4) HIMSS and NEMA grant permission to make copies and use this form.

Using the information in the MDS² form together with information collected about the care delivery environment (e.g., through tools like ACCE / ECRI's Guide for Information Security for Biomedical Technology), the provider's multidisciplinary risk assessment team can review assembled information and make informed decisions on implementing a local security management plan.

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**Section 1
INSTRUCTIONS FOR OBTAINING AND USING THE MDS² FORM**

1.1 OBTAINING THE MDS² FORM (HEALTHCARE PROVIDERS)

Completed MDS² forms for many devices and systems may be available directly from the Agfa HealthCare Product Security Website:

http://www.agfahealthcare.com/global/en/main/products_services/product-info/isp/index.jsp

1.2 USING THE MDS² FORM (HEALTHCARE PROVIDERS)

1.2.1 Section 1 – Questions 1-19

Section 1 of the MDS² form contains information on the type of data maintained / transmitted by the device, how the data is maintained / transmitted, and other security-related features incorporated in the device, as appropriate. The field "Other Security Considerations" allows the manufacturer to add some general security considerations.

PLEASE BE ADVISED—An indication of a device’s ability to perform any listed function (i.e., a “Yes” answer) is not an implicit or explicit endorsement or authorization by the manufacturer to configure the device or cause the device to perform those listed functions.

It is important to distinguish between capability and permission. The questions contained on the MDS² form refer to device capability. Permission is a contractual matter separate from the MDS² form and is not covered by the MDS² form. Making changes to a device without explicit manufacturer authorization may have significant contractual, regulatory and liability issues.

1.2.2 Section 2 – Explanatory notes

The optional section 2 of the MDS² form contains space for explanatory notes if the manufacturer needs more space to explain specific details to the answers on questions 1-19.

NOTE—Agfa HealthCare may elect to attach supplementary material if additional space for recommended practices or explanatory notes is necessary.

1.3 THE ROLE OF HEALTHCARE PROVIDERS IN THE SECURITY MANAGEMENT PROCESS

It is the obligation of the users of the MDS² form (e.g., the healthcare provider) to employ all necessary and appropriate safeguards to meet their regulatory and organizational requirements. The MDS² document is intended to assist healthcare providers in meeting their regulatory obligations regarding device security. The healthcare provider organization (e.g., a hospital) has the ultimate responsibility for providing effective security management. Agfa HealthCare can assist providers in their security management programs by offering information describing:

- the type of data maintained / transmitted by the manufacturer’s product;
- how data is maintained / transmitted by the manufacturer’s product;
- any security-related features incorporated in the manufacturer’s product.

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In order to effectively manage medical information security and comply with relevant regulations, healthcare providers must employ ADMINISTRATIVE, PHYSICAL and TECHNICAL SAFEGUARDS—most of which are extrinsic to the actual device

1.4 DEFINITIONS

Administrative Safeguards: Administrative actions, policies, and procedures to manage the selection, development, implementation, and maintenance of security measures to protect electronic Protected Health Information and to manage the conduct of the covered entity's workforce in relation to the protection of that information. [45 CFR Part 164]

Anti-Virus Software: See VIRUS SCANNER

Audit trail: Data collected and potentially used to facilitate a security audit [45 CFR Part 142]

Biometric ID: A biometric identification system identifies a human from a measurement of a physical feature or repeatable action of the individual (e.g., hand geometry, retinal scan, iris scan, fingerprint patterns, facial characteristics, DNA sequence characteristics, voice prints, handwritten signature). [45 CFR Part 142]

Electronic Media: (1) Electronic storage media, including memory devices in computers (hard drives) and any removable/transportable digital memory media, such as magnetic tapes or disks, optical disks, or digital memory cards. (2) Transmission media used to exchange information already in electronic storage media, including, for example, the Internet (wide open), extranet (using Internet technology to link a business with information accessible only to collaborating parties), leased lines, dial-up lines, and private networks, and the physical movement of removable/transportable electronic storage media. Certain transmissions, including of paper via facsimile and of voice via telephone, are not considered to be transmissions via electronic media because the information being exchanged did not exist in electronic form before the transmission. [45 CFR Part 160.103]

Electronic Protected Health Information (ePHI): individually identifiable health information (IIHI) that is (1) transmitted by or (2) maintained in electronic media. [45 CFR Part 160.103]

Individually Identifiable Health Information (IIHI): Individually identifiable health information is information that is a subset of health information, including demographic information collected from an individual, and: (1) Is created or received by a health care provider, health plan, employer, or health care clearinghouse; and (2) Relates to the past, present, or future physical or mental health or condition of an individual; the provision of health care to an individual; or the past, present, or future payment for the provision of health care to an individual; and (i) That identifies the individual; or (ii) With respect to which there is a reasonable basis to believe the information can be used to identify the individual. [45 CFR Part 160.103].

Personal Identification Number (PIN): A number or code assigned to an individual and used to provide verification of identity. [45 CFR Part 142]

Physical Safeguards: The physical measures, policies, and procedures to protect a covered entity's electronic information systems and related buildings and equipment from natural and environmental hazards and unauthorized intrusion. [45 CFR Part 164]

Remote Service: A support service (e.g., testing, diagnostics, software upgrades) while not physically or directly connected to the device (e.g., remote access via modem, network, Internet).

Removable Media: See ELECTRONIC MEDIA

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Security Risk Analysis: Conducting an accurate and thorough assessment of the potential risks and vulnerabilities to the integrity, availability, and confidentiality of electronic protected health information. [45 CFR Part 164]

Security Risk Management: (1) The ongoing process of assessing risk, taking steps to reduce risk to an acceptable level, and maintaining that level of risk. [NIST SP 800-26] (2) Security measures sufficient to reduce risks and vulnerabilities to a reasonable and appropriate level. [45 CFR Part 164]

Technical Safeguards: The technology, policies, and procedures to protect electronic Protected Health Information and control access to it. [45 CFR Part 164]

Token: A physical authentication device that the user carries (e.g., smartcard, SecureID[™], etc.). Often combined with a PIN to provide a two-factor authentication method that is generally thought of as superior to simple password authentication.

Virus: In general, computer code that is either:

- (1) A type of programmed threat—a code fragment (not an independent program) that reproduces by attaching to another program. It may damage data directly, or it may degrade system performance by taking over system resources, which are then not available to authorized users.
- (2) Code embedded within a program that causes a copy of itself to be inserted in one or more other programs; in addition to propagation, the virus usually performs some unwanted function. [45 CFR Part 164]

Virus scanner: A computer program (“ANTI-VIRUS SOFTWARE”) that detects a VIRUS computer program, or other kind of malware (e.g., worms and Trojans), warns of its presence, and attempts to prevent it from affecting the protected computer. Malware often results in undesired side effects generally unanticipated by the user.)

Vulnerability: A flaw or weakness in system procedures, design, implementation, or internal controls that could be exercised (accidentally triggered or intentionally exploited) and result in a security breach or a violation of the system’s security policy. [NIST SP 800-30]

ARCRONYMS

CD:	Compact Disk
CF:	Compact Flash
DVD:	Digital Versatile Disk
IP:	Internet Protocol
LAN:	Local Area Network
ROM:	Read Only Memory
SD:	Secure Digital

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USB:	Universal Serial Bus
VPN:	Virtual Private Network
WAN:	Wide Area Network
WiFi:	Wireless Fidelity

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Section 2 MDS² FORM

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SECTION 1

Device Category 17904 Digital Imaging	Manufacturer: Agfa Healthcare N.V.	Document ID: LLid_34019267	Document Release Date: August 15 2011
Device Model: NX (Modality Workstation)	Software Revision: X.0.8X00	Software Release Date 18 July 2011	
Manufacturer or Representative Contact Information:	Company Name: Agfa Healthcare N.V. Representative Name/ Position: John Gibbs / Product Manager modality Workstations	Manufacturer Contact Information Agfa Healthcare N.V Septestraat 27, 2640 Mortsel, Belgium Department: HE/CR-DR Solutions e-mail: John.Gibbs@agfa.com Telephone #:003234447560	

MANAGEMENT OF ELECTRONIC PROTECTED HEALTH INFORMATION (ePHI) *As defined by HIPAA Security Rule, 45 CFR Part 164* Yes No N/A Note #

1. Can this device transmit or maintain *electronic Protected Health Information* (ePHI)? Yes ___
2. Types of ePHI data elements that can be maintained by the device:
 - a. Demographic (e.g., name, address, location, unique identification number)? Yes ___ 1
 - b. Medical record (e.g., medical record #, account #, test or treatment date, device identification number)? Yes ___ 1
 - c. Diagnostic/therapeutic (e.g., photo/radiograph, test results, or physiologic data with identifying characteristics)? .. Yes ___ 2
 - d. Open, unstructured text entered by device user/operator? Yes ___
3. Maintaining ePHI - *Can the device*
 - a. Maintain ePHI temporarily in volatile memory (i.e., until cleared on by power-off or reset)? Yes ___
 - b. Store ePHI persistently on local media? Yes ___ 3
 - c. Import/export ePHI with other systems? Yes ___
4. Mechanisms used for the transmitting, importing/exporting of ePHI - *Can the device*
 - a. Display ePHI (e.g., video display)? Yes ___
 - b. Generate hardcopy reports or images containing ePHI? Yes ___
 - c. Retrieve ePHI from or record ePHI to removable media (e.g., disk, DVD, CD-ROM, tape, CF/SD card, memory stick)?.. Yes ___ 4
 - d. Transmit/receive or import/export ePHI via dedicated cable connection (e.g., IEEE 1073, serial port, USB, FireWire)? Yes ___
 - e. Transmit/receive ePHI via a network connection (e.g., LAN, WAN, VPN, intranet, Internet)?..... Yes ___
 - f. Transmit/receive ePHI via an integrated wireless connection (e.g. WiFi, Bluetooth, infrared)?..... Yes ___ 5
 - g. Other? No ___

ADMINISTRATIVE SAFEGUARDS Yes No N/A Note #

5. Does manufacturer offer operator and technical support training or documentation on device security features?..... No ___
6. What underlying operating system(s) (including version number) are used by the device?WIN XP ProSP3 / WIN7pro 32bit ___

PHYSICAL SAFEGUARDS Yes No N/A Note #

7. Are all device components maintaining ePHI (other than removable media) physically secure (i.e. cannot remove without tools)? Yes ___
8. Does the device have an integral data backup capability (i.e., backup onto removable media such as tape, disk)? Yes ___ 6
9. Can the device boot from uncontrolled or removable media (i.e., a source other than an internal drive or memory component)? Yes ___

TECHNICAL SAFEGUARDS Yes No N/A Note #

10. Can software or hardware not authorized by the device manufacturer be installed on the device? Yes ___ 7
11. Can the device be serviced remotely (i.e., maintenance activities performed by service person via network or remote connection)? .Yes ___
 - a. Can the device restrict remote access to specific devices or network locations (e.g., specific IP addresses)? Yes ___
 - b. Can the device log provide an audit trail of remote-service activity? Yes ___
 - c. Can security patches or other software be installed remotely? Yes ___
12. Level of owner/operator service access to device operating system: *Can the device owner/operator*
 - a. Apply device manufacturer-validated security patches?..... Yes ___

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Manufacturer or Representative Contact Information:	Company Name: Agfa Healthcare N.V.	Manufacturer Contact Information Agfa Healthcare N.V. Septestraat 27, 2640 Mortsel, Belgium	
	Representative Name/ Position: John Gibbs / Product Manager modality Workstations	Department: HE/CR-DR Solutions e-mail: John.Gibbs@agfa.com Telephone #:003234447560	

b. Install or update antivirus software?	Yes ___
c. Update virus definitions on manufacturer-installed antivirus software?	No ___ 8 ___
d. Obtain administrative privileges (e.g. access operating system or application via local root or admin account)?	Yes ___
13. Does the device support user/operator specific ID <i>and</i> password.....	Yes ___ 9 ___
14. Are access sessions terminated after a predetermined length of inactivity (e.g., auto logoff)?	Yes ___ 10 ___
15. Events recorded in device audit trail (e.g., user, date/time, action taken): <i>Can the audit trail record</i>	
a. Login and logout by users/operators?	Yes ___ 11 ___
b. Viewing of ePHI?	Yes ___
c. Creation, modification or deletion of ePHI?	Yes ___
d. Import/export or transmittal/receipt of ePHI?	Yes ___
16. Does the device incorporate an emergency access ("break-glass") feature that logs each instance of use?	No ___ 12 ___
17. Can the device maintain ePHI (e.g., by internal battery) during power service interruptions?	Yes ___ 13 ___
18. Controls when exchanging ePHI with other devices:	
a. Transmitted only via a point-to-point dedicated cable?	No ___
b. Encrypted prior to transmission via a network or removable media?	Yes ___ 14 ___
c. Restricted to a fixed list of network destinations (i.e., host-based access control list)?	Yes ___
19. Does the device ensure the integrity of the ePHI data with implicit or explicit error detection/correction technology?	No ___

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	Representative Name/ Position: John Gibbs / Product Manager modality Workstations	Agfa Healthcare N.V Septestraat 27, 2640 Mortsel, Belgium Department: HE/CR-DR Solutions e-mail: John.Gibbs@agfa.com Telephone #:003234447560	

Other Security Considerations

NX software uses Win XP/ WIN7 provided security features, a default security set-up is provided at installation. Security can be increased or decreased on request of the customer.

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SECTION 2

EXPLANATORY NOTES (from questions 1 – 19)

IMPORTANT: Refer to Section 1.2.2 of the Instructions for this form for the proper interpretation of information requested in this form).

1. Information is entered via keyboard or from patient data base (i.e. Radiology Information System).
2. Medical X-ray Images acquired from CR or DR X-ray modality.
3. Local media is internal hard drive.
4. Import/export via CD/DVD
5. Information is transferred to imaging cassettes via Radio Frequency communication. Also WLAN communication is possible.
6. PC can be delivered with RAID 1 disc mirroring, semi automated back-up to DVD is also possible.
7. Software can be installed by administrator level login. However, software not approved by Agfa, compromises Agfa warranty.
8. Anti-virus software is not provided during installation, but Agfa does make recommendations regarding the use of virus scan software.
9. Using Windows XP/WIN7 provided user log-in.
10. Windows auto log-off and other security features can be used.
11. Audit log reader is not part of the NX software.
12. Local procedures need to be put in place to access an emergency log-in/password;
13. Optional UPS can be provided
14. Secure communication to PACS or Printer via SLL (Secure Socket Layer) is possible.

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