

Exa® PACS/RIS

Antivirus Exclusions Reference Guide

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Introduction

Symbols

The following symbols may appear in the product documentation or on the product.

Symbol	Symbol Name	Symbol Description	Standard Number and Name	Symbol Reference Number
	Manufacturer	Indicates the name and address of the manufacturer	ISO 15223-1:2021	5.1.1
EC REP	Authorized Representative in the European Economic Area (EEA)	Indicates the Authorized Representative, responsible for the device in the European Economic Area (EEA).	ISO 15223-1:2021	5.1.2
	Date of Manufacture	Indicates the date when the device was manufactured.	ISO 15223-1:2021	5.1.3
	Caution	Indicates information that is important for preventing loss of data or misuse of the software.	ISO 15223-1:2021	5.4.4
LOT	Batch Code	Indicates the full Software Release / Version number	ISO 15233-1:2021	5.1.5
SN	Serial number	Indicates the manufacturer's serial number so that a specific medical device can be identified	ISO 15233-1:2021	5.1.7
REF	Catalogue Number	Indicates the manufacturer's catalogue number so that the device can be identified	ISO 15233-1:2021	5.1.6
i	Consult instructions for use	Indicates the need for the user to consult the instructions for use	ISO 15233-1:2021	5.4.3
R Only	Prescription Device	Caution: Federal law restricts this device to sale by or on the order of a licensed healthcare practitioner	21 CFR 801.109(b)(1) Prescription Devices	N/A

BS EN ISO 15223-1:2021 Medical devices - Symbols to be used with information to be supplied by the manufacturer - Part 1: General requirements

ex



Indications for use

EXA[™] is a software device that receives digital images and data from various sources (i.e. CT scanners, MR scanners, ultrasound systems, R/F Units, computed & direct radiographic devices, secondary capture devices, scanners, imaging gateways or other imaging sources). Images and data can be stored, communicated, processed, and displayed within the system and or across computer networks at distributed locations. Lossy compressed mammographic images are not intended for diagnostic review. Mammographic images should only be viewed with a monitor cleared by FDA for viewing mammographic images. For primary diagnosis, post process DICOM "for presentation" images must be used. Typical users of this system are trained professionals, nurses, and technicians.

Training

Users of this software must have received adequate training on its safe and effective use before attempting to operate the product described in this Instructions for Use. Users must make sure they receive adequate training in accordance with local laws or regulations.

Regulatory and compliance



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Items to exclude

The following lists the antivirus scanning and live scanning exceptions for the Exa platform, supported services, and their functions. Consult with your assigned project manager if you have any questions regarding program or service locations, as they may vary between servers or installations.

Apps (typically C:)

C: \Program Files\PostgreSQL\

• Exclude this entire folder.

C:\Program Files (x86)\pgAAdmin 4\

• Exclude this entire folder.

C:\Program Files\nodejs\

• node.exe

C: \opal\bin

• Exclude this entire folder.

Exa platform application drive (typically H: or C:)

%Application Drive%\Viztek\EXA\

- If possible, exclude this entire folder. Otherwise, exclude the following.
 - Nginx81\nginx.exe
 - Nginx82\nginx.exe
 - Nginx83\nginx.exe
 - Nginx84\nginx.exe
 - Redis\redis-server.exe
 - resx\nssm.exe
 - TxTranscription\bin\TXTextControl.Web.Server.exe

%Application Drive%\Viztek\EXA\bin\

• Exclude this entire folder.

Image drive (typically F:)

F:\EXA

• Exclude this entire folder.

Cache drive (typically G:)

%Image Cache Drive%\EXA\Cache\

• Exclude this entire folder.

PostgreSQL database (typically I:)*

[pgdata drive]:\pgdata

%PostgreSQL Drive%\PostgreSQL\%pssql-version%\

• Data Database tables/files.





Scanning or live scanning PostgreSQL data may decrease performance of Exa applications due to database read/write delays, and could lead to files becoming corrupted.