

DEC 11 1998



PHILIPS

K983821

Philips Medical Systems

510(k) Summary

Company name: Philips Medical Systems North America Company

Address: 710 Bridgeport Avenue, Shelton, CT 06484

Contact person: P. Altman

Telephone number: 203-926-7031

Prepared: October 28, 1998

Device name: **Philips Inturis Cardio Image Viewing Package**

Classification name: Image Processing System
Class II (90 LLZ)

Common/Usual name: Workstation

Predicate Device(s): Philips CD-Medical (K945460)

Intended use:

The PHILIPS Inturis Cardio Image Viewing Package is intended for use in processing and viewing cardiac or cardiovascular images, resulting in quantitative and/or qualitative data that can be used to estimate cardiac or cardiovascular function.

System description:

PHILIPS Inturis Cardio Image Viewing Package is software that can be loaded on a workstation which has a Windows NT (4.0) Operating System. The Inturis Cardio Image Viewing workstation provides for viewing and analysis of DICOM Cardiac images which are transported to the workstation via CD-Rom. Quantitative Coronary and Left Ventricular analysis may be performed on these images.

Substantial equivalence Information:

The PHILIPS Inturis Cardio Image Viewing Package is substantially equivalent to the Philips CD-Medical Viewstation (a.k.a. CDM-3500), K945460.

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DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

DEC 11 1998

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Peter Altman
Director of Regulatory Affairs
Philips Medical Systems
North America Company
710 Bridgeport Avenue
P.O. Box 860
Shelton, Connecticut 06484-0917

Re: K983821
Philips Inturis Cardio Image Viewing Package
Dated: October 28, 1998
Received: October 29, 1998
Regulatory class: II
21 CFR 892.2050/Procode: 90 LLZ

Dear Mr. Altman:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4613. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address "<http://www.fda.gov/cdrh/dsmr/dsmamain.html>".

Sincerely yours,

Lillian Yin, Ph.D.
Director, Division of Reproductive,
Abdominal, Ear, Nose and Throat
and Radiological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

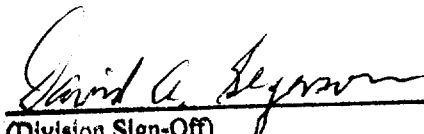
510(k) Number (if known): UnknownDevice Name : Philips Inturis Cardio Image Viewing Package

Indications For Use :

The **PHILIPS Inturis Cardio Image Viewing Package** is intended for use in processing and viewing cardiac or cardiovascular images, resulting in quantitative and/or qualitative data that can be used to estimate cardiac or cardiovascular function.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)


(Division Sign-Off)
Division of Reproductive, Abdominal, ENT,
and Radiological Devices
510(k) Number K983821

Prescription Use ☒
(Per 21 CFR 801.109

OR

Over-The-Counter Use ☐

(Optional Format 1-2-96)