

3/15/99



PHILIPS

K990423

510 (k) Summary

Philips Medical Systems

Company Name: Philips Medical Systems North America Company

Address: 710 Bridgeport Avenue
Shelton, CT 06484

Contact Person: Peter Altman

Telephone Number: 203-926-7031

Prepared (date): February 3, 1999

Device Name: Philips NICOL Collimator Family

Classification Name: Diagnostic X-ray Beam Limiting Device (90 IZW)

Common/Usual Name: Automatic Radiographic Collimator

Predicate Device: Philips Galileo Automatic Collimator

System Description:

The NICOL collimator family is a family of Beam Limiting Devices which may be used in all Philips Radiographic, Universal Radiographic/Fluoroscopic, Cardio and Vascular systems. It will provide rectangular shuttering, circular shuttering, or a combination of both. It will provide a light simulation of the X-ray field, spectral filtering, wedge filtering, dose measurement, and mechanical or electronic SID measurement.

Intended Use:

The Philips NICOL Collimator is intended for use in diagnostic x-ray systems during radiographic and fluoroscopic examinations. The collimator restricts the dimensions of the diagnostic x-ray field by limiting the size of the primary x-ray beam.

Safety Information:

The collimator has been designed to comply with the X-ray performance standard (21 CFR 1020.30, .31, and .32) and with UL 2601, Standard for Safety, Medical Electrical Equipment.

Philips Medical Systems
North America Company
710 Bridgeport Avenue
P.O. Box 860
Shelton, Connecticut 06484-0917
Tel: (203) 926-7674
Fax: (203) 929-6099



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Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850Peter Altman
Director of Regulatory Affairs
Philips Medical Systems
North America Company
710 Bridgeport Avenue
P.O. Box 860
Shelton, CT 06484-0917Re: K990423
NICOL Collimator Family
Dated: February 4, 1999
Received: February 11, 1999
Regulatory class: II
21 CFR 892.1610/Procode: 90 IZW

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 legally marketed predicate 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/dsma/dsmamain.html>".

Sincerely yours,

CAPT Daniel G. Schultz, M.D.
Acting 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): _____

Device Name : Philips NICOL Collimator Family

Indications For Use :

The Philips NICOL Collimator is intended for use in diagnostic x-ray systems during radiographic and fluoroscopic examinations. The collimator restricts the dimensions of the diagnostic x-ray field by limiting the size of the primary x-ray beam.

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

David A. Segman
(Division Sign-Off)

Division of Reproductive, Abdominal, ENT,
and Radiological Devices

510(k) Number K990423

Prescription Use
(Per 21 CFR 801.109

OR

Over-The-Counter Use _____

(Optional Format 1-2-96)