

APR 16 2004

K033103

### 510(K) SUMMARY

Manufacturer: Barco NV Barcoviev  
Theodoor Sevenslaan 106  
8500 Kortrijk  
Belgium

Submitted By: Ferguson Medical  
Consultant to Barco NV

Contact Information: Phone: +32(0) 56 23 32 11  
FAX: +32(0) 56 23 3 74

Classification Name: System, image processing

Common/Usual Name: Medical flat panel grayscale display, monitor,  
display and others

Proprietary Name: MGD 2621L

Classification Number: 21 CFR 892.2050/Procode 90LLZ

Substantial Equivalence: Barco MGD 221 2 Megapixel Diagnostic Display  
(K000293)

Device Description: The MGD 2621L is a diagnostic display

Intended Use: The MGD 2621L device is intended to be used  
in displaying and viewing digital images for  
review by trained medical practitioners.

Technological Characteristics: The MGD 2621L device is a high resolution  
monitor with electronic capabilities for  
evaluation of high resolution medical images.



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

APR 16 2004

BARCO NV BARCOVIEW  
% Mr. Frank Ferguson  
Official Correspondent  
Ferguson Medical  
12200 Academy Road NE, # 931  
ALBUQUERQUE NM 87111

Re: K033103  
Trade/Device Name: MGD 2621L Medical  
Grayscale Display  
Regulation Number: 21 CFR 892.2050  
Regulation Name: Picture archiving and  
communications system  
Regulatory Class: II  
Product Code: 90 LLZ  
Dated: July 14, 2003  
Received: January 26, 2004

Dear Mr. Ferguson:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and 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) that do not require approval of a premarket approval application (PMA). 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 (PMA), 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 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807), labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

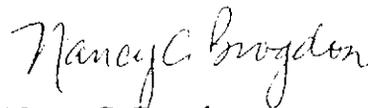
This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to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

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at one of the following numbers, based on the regulation number at the top of the letter:

8xx.1xxx	(301) 594-4591
876.2xxx, 3xxx, 4xxx, 5xxx	(301) 594-4616
884.2xxx, 3xxx, 4xxx, 5xxx, 6xxx	(301) 594-4616
892.2xxx, 3xxx, 4xxx, 5xxx	(301) 594-4654
Other	(301) 594-4692

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" (21CFR Part 807.97) you may obtain. Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer 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,



Nancy C. Brogdon  
Director, Division of Reproductive,  
Abdominal and Radiological Devices  
Office of Device Evaluation  
Center for Devices and Radiological Health

Enclosure

510(k) Number (If known): K033103

Device Name: MGD 2621L

Indications For Use:

The MGD 2621L Medical Grayscale Display is intended to be used as a tool in displaying and viewing digital images (excluding digital mammography) for review and analysis by trained medical practitioners.

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

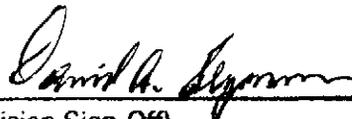
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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use XX  
(Per 21 CFR 801.109)

OR

Over-The-Counter Use \_\_\_\_\_



(Division Sign-Off)  
Division of Reproductive, Abdominal,  
and Radiological Devices

510(k) Number K033103