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K022322

GE Medical Systems

P.O. Box 414, W-709
Milwaukee, WI 53201
USA

510(K) SUMMARY OF SAFETY AND EFFECTIVENESS

This 510(k) summary of safety and effectiveness information is submitted in accordance with the requirements of 21 CFR Part 807.87(h).

Identification of Submitter: Larry A. Kroger, Ph.D.
Senior Regulatory Programs Manager
GE Medical Systems
Tel. ~~(414)~~ 262 544-3894
Summary prepared: 5 June, 2002

Identification of Product(s): Innova 2000 and Innova 2000 S
Classification Name: Solid State X-ray Imaging System
Manufacturer: GE Medical Systems Europe
283, rue de la Minière
78530 Buc Cedex, France
Distributed by: GE Medical Systems, Milwaukee, WI

Marketed Devices: The Digital Fluoroscopic Imaging Systems are substantially equivalent to the currently marketed cardiographic system **LCV+ Version 2** (K993037) that complies with the same or equivalent standards. -
The optional Fast Spin rotational angiography feature in the Innova 2000 system is substantially equivalent to the LCA system (K945375) for this feature.

Device Description: The **Digital Fluoroscopic Imaging System** is designed to perform fluoroscopic x-ray examinations. The detector is comprised of amorphous silicon with a cesium iodide scintillator. The resulting digital image can be sent through a Fiber Channel link to an acquisition equipment then to network (in using DICOM) for applications such as post-processing, printing, viewing and archiving. **Digital Fluoroscopic Imaging System** consists of an angiographic monoplane positioner, a table, an X-RAY system and a digital detector.
The Innova 2000 S system is identical to the Innova 2000 system but without the optional features.

New optional features : The offset C-arm permits rotational cardiac angiography over a total 200° at variable speed from 20° to 40°/sec with Cranio/Caudal angulation.
The SuperFast Gantry (SFG) includes capacitive sensor technology and optimized collision avoidance software that permits an increase of pivot and C-arm speed of up to 20°/sec.

Materials: All construction and materials are compliant with UL 2601.

Design: There are hardware and software redundancies to prevent from single point failures that could cause unintended motion.

Energy Source: 480 VAC 50/60Hz.

Indications for Use: The **Digital Fluoroscopic Imaging Systems** are indicated for use in generating fluoroscopic images of human anatomy for diagnostic and interventional cardiac angiography, and optionally, rotational cardiac angiography procedures. It is intended to replace fluoroscopic images obtained through the image intensifier technology.

Comparison with predicate: These systems are substantially equivalent to the Fluoroscopic system LCV+ Version 2 (K993037).
The optional Fast Spin rotational angiography feature in the Innova 2000 system is substantially equivalent to the LCA system (K945375) for this feature.

Summary of Studies: Not applicable as the digital technology (SSXD, i.e., flat panel detectors) has already been clinically compared in the submission of LCV+ Version 2 (K993037) with the image intensified technology system LC introduced in 1989 (cleared under K890348) and found substantially equivalent.

Conclusions: GE considers the systems to be equivalent with the predicate devices. The systems provide fluoroscopic images that are equivalent to the diagnostic capabilities of the predicate device images. The potential hazards, e.g., wrong measurements, misdiagnosis and increased gantry speeds are controlled by a risk management plan including:

- A hazard identification
- A risk evaluation
- A Software Development and Validation Process

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
10903 New Hampshire Avenue
Document Control Room – WO66-G609
Silver Spring, MD 20993-0002

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Re: K022322
Trade/Device Name: Innova 2000 and Innova 2000S
Regulation Number: 21 CFR 892.1650
Regulation Name: Image-intensified fluoroscopic x-ray system
Regulatory Class: II
Product Code: OWB and JAA
Dated: July 16, 2002
Received: July 17, 2002

Dear Dr. Kroger:

This letter corrects our substantially equivalent letter of October 8, 2002.

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 class II (Special Controls), it may be subject to such additional controls. Existing major regulations affecting your device can be found in Title 21, Code of Federal Regulations (CFR), Parts 800 to 895. 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 Parts 801 and 809); medical device reporting (reporting of

medical device-related adverse events) (21 CFR 803); and good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820). 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 Parts 801 and 809), please contact the Office of *In Vitro* Diagnostic Device Evaluation and Safety at (301) 796-5450. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely Yours,

Janine M. Morris
Acting Director
Division of Radiological Devices
Office of In Vitro Diagnostic Device
Evaluation and Safety
Center for Devices and Radiological Health

Enclosure

STATEMENT OF INTENDED USE

510(k) Number (if known): K022322

Device Name: **Digital Fluoroscopic Imaging System – Innova 2000 and Innova 2000 S**

Indications for Use

The Digital Fluoroscopic Imaging Systems are indicated for use in diagnostic and interventional cardiac angiography and optionally, rotational cardiac angiography procedures of human anatomy. It is intended to replace image intensifier fluoroscopic systems in all diagnostic or interventional cardiac angiography procedures.

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use √
(Per 21 CFR 801-109)

OR Over-The-Counter Use_____

David G. Lyon

(Division Sign-Off)
Division of Reproductive, Abdominal,
and Radiological Devices
510(k) Number K022322