

Attachment 1

OCT 27 2006

Summary of Safety and Effectiveness

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

General Information:

Product Name: Superopen 0.23T

Product Model: NAM-P023A

CFR Section: 21 CFR Part 892.1000
Magnetic resonance diagnostic device

Classification Name: System, Magnetic Resonance Imaging

Product Code: LNH

Device Class: Class II

Applicable Standard: IEC60601-1, Medical electrical equipment - Part 1: General Requirements for Safety
IEC60601-2-33, Medical electrical equipment – Part 2-33: Particular requirements for the safety of magnetic resonance equipment for medical diagnosis
21 CFR Subchapter J, Radiological Health
IEC60825-1, Safety of laser products-Part1:Equipment classification, requirement and user's guide
DICOM 3.0
NEMA MS Series (MS1 – MS8)

Manufacture and Distributor: Neusoft Medical Systems Co., Ltd.
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Shenyang, China
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Summary prepared : August 15th, 2006

Safety and Effectiveness information**Intended Uses:**

The Superopen 0.23T(Modified) is intended to produce images that reflect the spatial distribution of protons (hydrogen nuclei) exhibiting magnetic resonance. The NMR properties that determine the image appearance are proton density, spin-lattice relaxation time (T1), spin-spin relaxation time (T2) and flow. When interpreted by a trained physician, these images provide information that can be useful in determining a diagnosis.

Device Description:

The Superopen 0.23T(Modified) is a 0.35T permanent magnet MRI system. The magnet is mainly made of NdFeB material. The system software based on Windows (TM) is an interactive program with user-friendly interface. Its functions cover scanning control, image reconstruction and image/archive management and maintenance.

Predicated Device:

K033315 : Superopen 0.23T

Statement of Substantial Equivalence:

The Superopen 0.23T(Modified) is of comparable type and substantially equivalent to the Superopen 0.23T (K033315) in that they are similar in technology and intended uses. It is a modified product based on the Superopen 0.23T. Both of these systems are open-permanent-magnet MR Imaging System, use Gradient Subsystem to provide controlled and uniform gradient magnet fields in the X, Y and Z planes, and use RF Subsystem to complete the function of RF signal transmitting/receiving and processing. Image reconstruction is controlled by console's computer that has an interactive user interface, and the system produces 2D and 3D image that can be filmed or electronically stored for future review. Both of these systems have the traditional MRI unit.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Food and Drug Administration
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OCT 27 2006

Re: K062860
Trade/Device Name: Superopen 0.23T
Regulation Number: 21 CFR 892.1000
Regulation Name: Magnetic resonance diagnostic device
Regulatory Class: II
Product Code: LNH
Dated: September 20, 2006
Received: September 25, 2006

Dear Mr. Yanfang:

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 (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 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.



Protecting and Promoting Public Health

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 this letter:

21 CFR 876.xxx	(Gastroenterology/Renal/Urology)	240-276-0115
21 CFR 884.xxx	(Obstetrics/Gynecology)	240-276-0115
21 CFR 894.xxx	(Radiology)	240-276-0120
Other		240-276-0100

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 from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.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) K062860

Device Name: Superopen 0.23T

Indications for use:

The Superopen 0.23T(Modified) is an imaging device, and is intended to provide the physician with physiological and clinical information obtained non-invasively and without the use of ionizing radiation. The MRI system produces transverse, coronal, sagittal, oblique, and curved cross-sectional images that display the internal structure of the head, body, or extremities. The images produced by MRI system reflect the spatial distribution of protons (hydrogen nuclei) exhibiting magnetic resonance. The NMR properties that determine the image appearance are proton density, spin-lattice relaxation time (T1), spin-spin relaxation time (T2), and flow. When interpreted by a trained physician, these images provide information that can be useful in diagnosis determination.

The indications for use are as follows:

Anatomical Region: Head, Body, Spine, Extremities

Nucleus excited: Proton

Diagnostic uses: T1,T2, proton density weighted imaging

Diffusion weighted imaging

MR Angiography

Imaging processing

Imaging capabilities: 2D, 3D Spin Echo(SE)

Short time inversion recovery (STIR)

Fluid attenuated inversion recovery (FLAIR)

2D,3D Field Echo (FE)

2D, 3D Field Echo with Spoiler (FESP)

2D FESP Multi-Slice (FESP-MS)

2D and 3D Field Echo Steady State FID with rephasing gradient (FESS-FID)

2D, 3D Fast Spin Echo (FSE)

2D, 3D MRCP

MR Angiography

2D, 3D TOF

MTC

Echo Planar Imaging (EPI)

Multi-shot SE / FE

Diffusion

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

David H. Seymour

(Division Sign-Off)

Division of Reproductive, Abdominal,
and Radiological Devices

510(k) Number K062860

Prescription Use ✓