K071415

pg 1 of 2

SECTION 2. SUMMARY AND CERTIFICATION

A. 510(k) Summary

Submitter: Nonin Medical, Inc.

Contact Person: Lori M. Roth
Clinical/Regulatory Specialist
Nonin Medical, Inc.
13700 1st Ave. North
Plymouth, MN 55441-5443

AUG - 8 2007

Date Prepared: May 18, 2007

Trade Name: Model 7500FO Digital Pulse Oximeter

Classification Name: and Number: Class II, 21 CFR 870.2700

Product Code: DQA

Predicate Device(s): Nonin's Model 7500FO is substantially equivalent to the Model 8604FO Pulse Oximeter and Fiber Optic Sensor manufactured by Nonin Medical, Inc. that was cleared by the FDA under K910001 on 7/1/91.

Device Description: The Nonin® Model 7500FO Digital Pulse Oximeter is a portable, tabletop device indicated for use in measuring, displaying, and recording functional oxygen saturation of arterial hemoglobin (SpO_2) and pulse rate of adult, pediatric, and infant patients in an MR environment. Testing was performed in MR conditional environments at 1.5T and 3T. It is indicated for spot-checking and / or continuous monitoring of patients who are well or poorly perfused.

The Model 7500FO display uses light-emitting diodes (LED) components to present patient's SpO_2 and pulse rate values, as well as alarm limit and volume settings. The Model 7500FO can be powered externally with a 12 VDC 1.5A AC adapter or with an integral sealed 7.2-volt rechargeable NiMH battery pack.

The Model 7500FO includes adjustable audible and visual pulse rate, oxygen saturation, and perfusion alarms. It also includes a variety of advanced features, including low

	battery alarms, sensor alarm, user defined defaults, real-time data outputs, and patient security mode. Nonin's fiber optic sensors and cables contain no conductive components, they can safely be placed on the patient's finger while inside an MR (magnetic resonance) environment.
Intended Use:	The Nonin® Model 7500FO Digital Pulse Oximeter is a portable, tabletop device indicated for use in measuring, displaying, and recording functional oxygen saturation of arterial hemoglobin (SpO_2) and pulse rate of adult, pediatric, and infant patients in an MR environment. Testing was performed in MR conditional environments at 1.5T and 3T. It is indicated for spot-checking and / or continuous monitoring of patients who are well or poorly perfused.
Functional and Safety Testing:	Nonin's Model 7500FO Pulse Oximeter and fiber optic sensor have successfully undergone both bench and clinical testing in order to demonstrate that it has appropriate functional features and is substantially equivalent to the predicate device.
Conclusion:	Nonin's Model 7500FO is substantially equivalent to the Model 8604FO Pulse Oximeter manufactured by Nonin Medical, Inc. and cleared by the FDA under K910001 on 7/01/91.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

AUG - 8 2007

Ms. Lori M. Roth
Clinical Regulatory Specialist
Nonin Medical, Incorporated
13700 1st Avenue North
Plymouth, Minnesota 55441-5443

Re: K071415
Trade/Device Name: Model 7500FO Digital Pulse Oximeter
Regulation Number: 870.2700
Regulation Name: Oximeter
Regulatory Class: II
Product Code: DQA
Dated: May 18, 2007
Received: May 21, 2007

Dear Ms. Roth:

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.

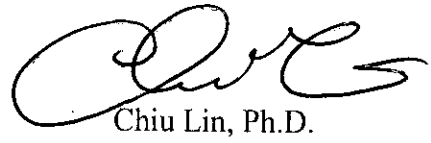
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish 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 (240) 276-0115. 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 (301) 443-6597 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,



Chiu Lin, Ph.D.
Director
Division of Anesthesiology, General Hospital,
Infection Control and Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indications for Use Statement

510(K) Number: K071415

Device Name:

Model 7500FO Digital Pulse Oximeter

Indications for Use:

The Nonin® Model 7500FO Digital Pulse Oximeter is a portable, tabletop device indicated for use in measuring, displaying, and recording functional oxygen saturation of arterial hemoglobin (SpO_2) and pulse rate of adult, pediatric, and infant patients in an MR environment. Testing was performed in MR conditional environments at 1.5T and 3T. It is indicated for spot-checking and / or continuous monitoring of patients who are well or poorly perfused.

(Division Sign-Off) (acting B.C.)
Division of Anesthesiology, General Hospital,
Infection Control, Dental Devices
510(k) Number: K071415

Prescription Use ___X___ AND/OR Over-The-Counter Use _____
(Part 21 CFR 801 Subpart D) (21 CFR 807 Subpart C)

Concurrence of CDRH, Office of Device Evaluation (ODE)