

DEC 03 2001

SUMMARY OF SAFETY AND EFFECTIVENESS FOR
NIPRO SAFELET CATH

K013287

§ 807.92 (a) (1)

Contact Person: Eiji Shinozaki

Date of Summary Preparation:

§ 807.92 (a)(2)

September 27, 2001



NIPRO MEDICAL CORPORATION
3150 N.W. 107 Avenue
Miami, Florida 33172
Tel.: (305) 599-7174
Fax: (305) 599-8454

Trade Name:	Nipro Safelet Cath
Common Name:	Intravascular catheter
Classification Name:	Intravascular catheter (for short-term use) (§ 880.5200)

§ 807.92 (a)(3)

Legally Marketed Substantially Equivalent Device:
Marketed in U.S. under K960051.

§ 908.92 (a)(4)

Description of Device:

The devices we intend to market are intravascular catheters as described in 21 CFR 880.5200. The subject devices are similar to catheters currently marketed by Nipro Medical Corporation under K960051.

Two types of catheters will be available: the container cases types L (long) and S (short). Type L catheters have needles 1 ½ to 2 ½ inches long and Type S have needles ¾ to 1 ¼ inches long. Six gauges of catheters are described here (14, 16, 18, 20, 22 and 24 gauge). The tip of the cannula is beveled. The proximal end of the catheter has a filter cap.

§ 807.92 (a)(5)

Intended Use:

The Nipro Safelet Cath is intended to be inserted into a patient's vascular system for short term use to sample blood, monitor blood pressure or administer fluids intravenously.

§ 807.92 (a)(6)

Comparison of Technical Characteristics:

Nipro Safelet Cath produced in Brazil and in Thailand are similar. Nipro Medical Corporation manufactures the subject devices in Brazil rather than in Thailand. Given the similarities outlined in Table 7, there is no question of their substantial equivalence.



DEC 03 2001

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Ms. Kaelyn B. Hadley
Consultant
Nipro Medical Corporation
1384 Copperfield Court
Lexington, Kentucky 40514

Re: K013287

Trade/Device Name: Nipro Safelet Cath
Regulation Number: 880.5200
Regulation Name: Intravascular Catheter
Regulatory Class: II
Product Code: FOZ
Dated: September 27, 2001
Received: October 2, 2001

Dear Ms. Hadley:

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.

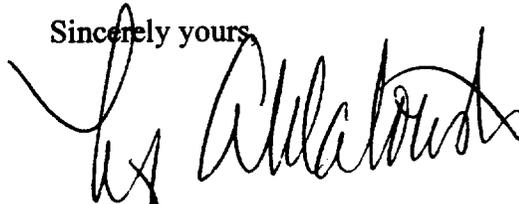
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies.

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 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4618. 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). 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,



Timothy A. Ulatowski
Director

Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indications for Use Statement

510(k) number (if known): K01???? K013287

Device name: Nipro® Safelet Cath

Indications for use: The Nipro® Safelet Cath is intended to be inserted into a patient's vascular system for short-term use to sample blood or administer fluids intravenously.

(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
 (optional Format 1-2-9)

Articia Cucurto
(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(k) Number K013287