K040981

MAY - 7 2004

ATTACHMENT F: 510(k) Summary

SPONSOR:

Wilson-Cook Medical
4900 Bethania Station Road
Winston-Salem, NC 27105

CONTACT/SUBMITTER:

Marge Walls-Walker
Regulatory Affairs Specialist
[336] -744-0157 Ex.290

DATE OF SUBMISSION:

April 14, 2004

DEVICE:

USW Needleknife Papillotome

Trade Name:
Common Name:
Classification:

Wilson-Cook USW Needleknife Papillotome
Needleknife, Precut
Unit, Electrosurgical Endoscopic w,w/o Accessories. 21 CFR § 876.4300, 78 KNS

PREDICATE DEVICES:

Wilson-Cook Needleknife Papillotome (k972674)

INTENDED USE:

Wilson-Cook's USW Needleknife Papillotome is intended for accessing the common bile duct when standard cannulation methods have been exhausted. This device is supplied sterile and intended for single use.

DEVICE DESCRIPTION:

The proposed USW Needleknife Papillotome is a triple-lumen catheter, with one lumen dedicated to the cutting wire, wire guide access and contrast injection respectively. The wire guide lumen is compatible with a .035" wire guide. The 24 GA stainless steel cutting wire allows for the transfer of electric current enhance cannulation efforts when traditional cannulation efforts have been exhausted. The handle contains an electrocautery connection pin, which is compatible with a variety of active cords currently sold separately.

COMPARISON OF CHARACTERISITICS:

We believe the proposed device to be substantially equivalent to the named predicate in terms of Intended Use, Indications for Use, performance characteristics tested, materials of construction, operation and biocompatibility.

PERFORMANCE DATA:

Non-Clinical Testing was performed on characteristics of the USW Needleknife Papillotome deemed necessary to verify safety and performance.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

MAY - 7 2004

Ms. Marge Walls-Walker
Regulatory Affairs Specialist
Wilson-Cook Medical
4900 Bethania Station Road
WINSTON-SALEM NC 27105

Re: K040981
Trade/Device Name: USW Needleknife Papillotome
Regulation Number: 21 CFR §876.4300
Regulation Name: Endoscopic electrosurgical unit and accessories
Regulatory Class: II
Product Code: 78 KNS
Dated: April 14, 2004
Received: April 20, 2004

Dear Ms. Walls-Walker:

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 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

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): K040981

Device Name: Wilson-Cook USW Needleknife Papillotome

Indications for Use:

The Wilson-Cook USW Needleknife Papillotome is intended for accessing the common bile duct when standard cannulation methods have been exhausted. The USW Needleknife Papillotome is supplied sterile and intended for single use only.

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use Only ✓ OR Over-the-Counter ____
(Per 21 CFR § 801.109

David A. Lynn

(Division Sign-Off)
Division of Reproductive, Abdominal,
and Radiological Devices
510(k) Number K040981