

DEC 21 1999

510(k) Summary

1.0 Date Prepared

October 28, 1999

2.0 Submitter (Contact)

Martin D. Sargent
Xomed Surgical Products
Jacksonville, FL
(904) 279-7586

3.0 Device Name

Proprietary Name: Micro-France electrosurgical instruments
Common Name(s): Electrosurgical cutting and coagulation accessories
Classification Name(s): Electrosurgical cutting and coagulation accessories

5.0 Device Classification

Electrosurgical cutting and coagulation accessories:
Procure JOS Class II 21 CFR 878.4400

6.0 Device Description

The various unipolar and bipolar Micro-France electrosurgical instruments consist of scissors, forceps, and probes, and are available in configurations for laparoscopic / endoscopic access and open field surgery. Accessories include unipolar and bipolar cables available in lengths up to 3 M.

7.0 Intended Use

The Electrosurgical instruments are used to remove tissue and control bleeding.

8.0 Substantial Equivalence

The Electrosurgical instruments are substantially equivalent to devices marketed by Richard Wolf Medical Instruments (K980129), Concept (K792365), Instrument Maker (K890972), Linvatec (K944992), Megadyne Medical Products (K903302), New England Surgical Instrument (K895331), and Kirwan Surgical Products (K913514).



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

DEC 21 1999

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Mr. Martin D. Sargent
Senior Regulatory Affairs Specialist
Xomed Surgical Products
6743 Southpoint Drive, North
Jacksonville, Florida 32216-0980

Re: K993655
Trade Name: Micro-France Electrosurgical Instruments, Various
Regulatory Class: II
Product Code: GEI
Dated: October 28, 1999
Received: October 29, 1999

Dear Mr. Sargent:

We have reviewed your Section 510(k) notification of intent to market the devices referenced above and we have determined the devices are substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Devices Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the devices, 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.

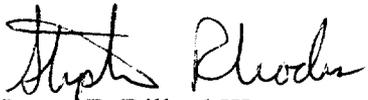
If your devices are classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your devices can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the current Good Manufacturing Practice requirement, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic (QS) inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your devices in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your devices as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your devices to a legally marketed predicate device results in a classification for your devices and thus, permits your devices to proceed to the market.

Page 2 - Mr. Martin D. Sargent

If you desire specific advice for your devices on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4595. Additionally, for questions on the promotion and advertising of your devices, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

Sincerely yours,

for 

James E. Dillard III
Acting Director
Division of General and
Restorative Devices
Office of Devices Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number (if known): _____

Device Name: Electrosurgical Cutting and Coagulation Accessories

Indications for Use:

The electrosurgical instruments are intended to remove tissue and control bleeding.

(Please do not write below this line - continue on another page if needed)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Steph Rhodes

(Division Sign-Off)
Division of General Restorative Devices

510(k) Number K993655

Prescription Use ✓
(Per 21 CFR 801.109)

Or

Over-the-Counter Use _____

(Optional Format 1-2-96)