



Food and Drug-Administration
10903 New Hampshire Avenue
Document Control Room - WO66-G609
Silver Spring, MD 20993-0002

Mr. Peter Kim
Quality Manager
International Medsurg Connection
935 North Plum Grove Road, Suite F
Schaumburg, Illinois 60173

APR 20 2010

Re: K100208
Trade/Device Name: IMC-Surgical Drapes – Various Types, Sizes and Configurations
Regulation Number: 21 CFR 878.4370
Regulation Name: Surgical Drape and Drape Accessories
Regulatory Class: II
Product Code: KXX
Dated: March 8, 2010
Received: March 23, 2010

Dear Mr. Peter Kim:

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. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

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); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); 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.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please go to <http://www.fda.gov/AboutFDA/CentersOffices/CDRH/CDRHOffices/ucm115809.htm> for the Center for Devices and Radiological Health's (CDRH's) Office of Compliance. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

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) 796-7100 or at its Internet address <http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,



Anthony D. Watson, BS, MS, MBA

Director

Division of Anesthesiology, General Hospital,

Infection Control, and Dental Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

Enclosure

4.0 INTENDED USES / INDICATION:

International Medsurg Connection's Surgical Drape is intended to be used as protective coverings used to isolate incision sites and protect against contamination during surgical procedures. The intended use is the same as the intended use on IMC surgical drapes sold by International Medsurg Connection, Reference K052169.

This submission includes drapes that will be sold both sterile and non-sterile. Non-sterile drapes are to be sold to OEMs for EtO sterilization according to ISO 11135. Sterile drapes are to be sold directly to users according to current sterilization process. Related items are indicated at the table below.

Category	Device Name	Catalog #	Sterile/Non-sterile
Surgical Drape	Lithotomy Procedure Drape	21-0043	Sterile
	Lithotomy Procedure Drape	21-0043NS	Non-sterile
	PCNL Procedure Drape	41-0021	Sterile
	PCNL Procedure Drape	41-0021NS	Non-sterile
	GYN Procedure Drape	61-0071	Sterile
	GYN Procedure Drape	61-0071NS	Non-sterile

Indications for Use

510(k) Number :

Device Name: IMC-Surgical Drapes-various types, sizes and configurations (See *

ATTACHMENT)

Indications For Use: This device is intended to be used as protective coverings used to isolate incision sites and protect against contamination during surgical procedures.

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

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

Concurrence of CDRH, Office of Device Evaluation (ODE)



(Division Sign-Off)
Division of Anesthesiology, General Hospital
Infection Control, Dental Devices

510(k) Number: K100208