

9. 510(K) SUMMARY

Date Prepared: November 05, 2007

JAN 16 2008

SUMMARY OF SAFETY AND EFFECTIVENESS

Submitted By: Robert Licha
Director Product and Process Management
Schreiner MediPharm

- 9.1 Trade/Proprietary Name: Needle Trap™ / Needle Safety Label
- 9.2 Common/Usual Name Syringe Anti-Needle Stick Accessory
- 9.3 Classification Name Antistick Syringe
- 9.4 Substantial Equivalence
The Needle Trap™ Needle Safety devices are substantially equivalent to the BD SafetyGlide™ Needle (K951254), Portex® Needle-Pro® EDGE™ Safety Device with Syringe (K061194) and/or the BD Eclipse™ Hypodermic Needle (K010188)
- 9.5 Classification: Class II
Panel: 80
Product Code: 80MEG
Cite: 21CFR 880.5860 Piston syringe
FMF - Piston Syringe
MEG - Antistick Syringe
FMI - Hypodermic Single Lumen Needle
- 9.6 Technological Characteristics
The technological characteristics of the Needle Trap™ Needle Safety devices are similar to the current marketed products
- 9.7 Performance Data
Verification and Validation testing confirmed the products meet their specifications.
- 9.8 Conclusion
Schreiner MediPharm concludes based on the information presented that the new products lines are substantially equivalent to products currently legally marketed in the USA



Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JAN 16 2008

Schreiner MediPharm
C/O Mr. Lee Leichter
President
P/L Biomedical
7609 Cameron Circle
Fort Myers, Florida 33912

Re: K073206
Trade/Device Name: Needle-Trap™
Regulation Number: 21 CFR 880.5860
Regulation Name: Piston Syringe
Regulatory Class: II
Product Code: MEG
Dated: November 8, 2007
Received: November 13, 2007

Dear Mr. Leichter:

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.

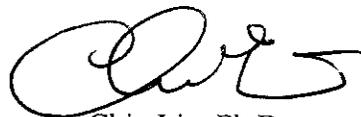
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish 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

510(k) Number: K073206

Device Name: Needle-Trap™

Indications for Use: The Needle-Trap™ contains a mechanism that is used to cover the needle point after use of the syringe and needle for their intended functions. In the activated position the Needle-Trap™ guards against accidental needle sticks during normal handling and disposal of the used needle/syringe combination.

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use
(Per 21 CFR 801.109)

or

Over-the-Counter Use


(Division Sign-Off)

Division of Anesthesiology, General Hospital
Infection Control, Dental Devices

510(k) Number: K073206