

AUG 29 2002

K022063

SECTION VIII

**510(K) SUMMARY OF SAFETY AND EFFECTIVENESS
for 1cc SafePro* Safety Syringe**

1. REGULATORY AUTHORITY

Safe Medical Device Act of 1990, CFR 807.92.

2. CONTACT PERSON

Joseph J. Chang, Ph.D., PE
FORMOSA MEDICAL DEVICES, INC.
U.S. Liaison Office
11497 Columbia Park Drive West, Suite #9
Jacksonville, FL 32258

3. NAME OF MEDICAL DEVICE

Classification Name:	Syringe, Antistick
Classification Code:	MEG
Common/Usual Name:	Syringe
Proprietary Name:	1cc SafePro* Safety Syringe

4. DEVICE CLASSIFICATION

The General Hospital Panel has classified Antistick Syringes (21CFR880.5860) into Class II, Special Controls under section 513 of the Act.

5. STATEMENT OF SUBSTANTIAL EQUIVALENCE

The 1cc SafePro* Safety Syringe is substantially equivalent to:

A) B-D Luer Lok syringe in terms of normal syringe function for fluid transport, and

B) Retractable Technology, Inc. VanishPoint, and SafePro* Safety Syringe for needlestick protection feature. See also the attached table.

*Trademark

6. INTENDED USE

The primary intended use for the 1cc SafePro* Safety Syringe is for general purpose use in aspirating and injecting fluid.

The secondary intended use for the 1cc SafePro* Safety Syringe is for needlestick protection; the device may aid in the reduction of needlestick injuries.

7. DESCRIPTION OF DEVICE

The 1cc SafePro* Safety Syringe consists of a syringe assembly and a needle assembly. The device has a built-in safety feature which may aid in the reduction of the risk of accidental needlestick injuries.

8. SUMMARY OF MATERIAL TESTING

The entire SafePro* Safety Syringe was already tested for material safety and biocompatibility as indicated in a previous 510(K) submission, K012726. Since the 1cc SafePro* Safety Syringes have different gauge size needles, the colorant used are different. Therefore, the hubs molded with all of the colorants for all gauges were tested for biocompatibility. Test results indicated that the hub and thus the entire 1cc SafePro* Safety Syringe meets ISO 10993-1 and US FDA G-95 requirements.

9. SUMMARY OF SIMULATED USE STUDY

A total of 500 1cc SafePro* Safety Syringes were evaluated by 50 participants. No sharps injuries or failure of the safety mechanism occurred. Successful completion of the study supports the claim that 1cc SafePro* Safety Syringe can reduce the risk of accidental needlestick injuries. The positive responses from the Evaluators regarding functional and performance aspects also indicated that the 1cc SafePro* Safety Syringe meets customer requirements.

10. CONCLUSION

The results obtained from bench testing, material safety, and simulated use tests indicate that the 1cc SafePro* Safety Syringe is safe and effective for its intended use.



AUG 29 2002

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Formosa Medical Devices, Incorporated
C/O Mr. Joseph J. Chang
SafePro USA, Incorporated
11497 Columbia Park Drive West, Suite #9
Jacksonville, Florida 32258

Re: K022063

Trade/Device Name: 1cc SafePro* Safety Syringe

Regulation Number: 880.5860

Regulation Name: Piston Syringe

Regulatory Class: II

Product Code: MEG

Dated: June 24, 2002

Received: June 25, 2002

Dear Mr. Chang

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.

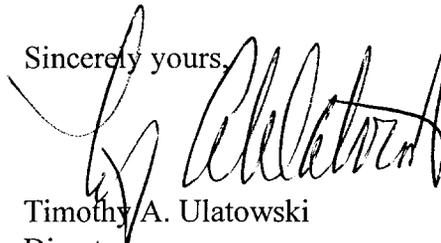
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You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 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 Anesthesiology, General Hospital,
Infection Control and Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

K 022063

INDICATIONS FOR USE

Device Name

1cc SafePro* Safety Syringe

Indication for Use

The primary indicated use for the 1cc SafePro* Safety Syringe is for general purpose use in aspirating and injecting fluid.

The secondary indicated use for the 1cc SafePro* Safety Syringe is for needlestick protection; the device may aid in the reduction of needlestick injuries.

(Please 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
(Optional Format 1-2-96)

OR Over-the-Counter _____

Leticia Cuervo

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

510(k) Number: K022063

*Trademark