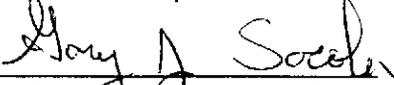


MAY - 6 2004

**510(k) Summary of  
Safety and Effectiveness**

K041099

**Submitter:**

- SPSmedical Supply Corp.  
6789 West Henrietta Road  
Rush, NY 14543 U.S.A.  
Phone: (585)-359-0130 Fax: (585)-359-0167
- Establishment FDA Registration No.: 1319130
- Date Summary was Prepared April 21<sup>st</sup>, 2004
- Gary J. Socola  
Printed name of person submitting for 510(k)
-   
Signature of person submitting for 510(k)
- Director of Quality Assurance  
Title of person submitting for 510(k)

**Device Name and Classification**

Trade Name: SPSmedical SporView<sup>®</sup> Bacterial Spore Strip  
Classification Name: Sterilization Process Biological Indicator  
Common Name: Spore Strip Biological Indicator  
Device Classification: Class II, Regulation Number 880.2800  
Product Code: FRC  
Predicate Device: SPSmedical SporView<sup>®</sup> Bacterial Spore Strip K023716 & K905425

#### Device Description:

The SPSmedical SporView<sup>®</sup> Bacterial Spore Strip is inoculated with either *Geobacillus stearothermophilus* ATCC# 7953 or *Bacillus atrophaeus* ATCC# 9372 bacterial spores. A dual species spore strip containing both the *Geobacillus stearothermophilus* ATCC# 7953 and *Bacillus atrophaeus* ATCC# 9372 organisms is also manufactured. The strip is packaged in a blue, glassine envelope.

#### Intended Use:

The SPSmedical SporView<sup>®</sup> Bacterial Spore Strip is designed for routine monitoring of Steam, Dry Heat, EO Gas, and Chemiclave<sup>®</sup> sterilization processes. The indicator should be placed in the most difficult to sterilize location of the Chemiclave<sup>®</sup> sterilization load. The spore strip is aseptically transferred into SPSmedical's SporView<sup>®</sup> Culture Media (Tryptic Soy Broth with Bromocresol Purple) and incubated at 60°C. The combination of SPSmedical's spore strip and media has been validated for a reduced incubation time of 72 hours for the Chemiclave<sup>®</sup> chemical vapor sterilization process.

#### Statement of Similarity to the Legally Marketed Predicate Device:

- Have the same indicated use
- Incorporate the same materials
- Have the same shelf life
- Packaged using the same materials and processes

#### Non-Clinical Testing:

Testing was conducted following the FDA guidance for the validation of reduced incubation of biological indicators. Testing was performed for the Chemiclave<sup>®</sup> chemical vapor sterilization process using three lots of biological indicators; this included a newly manufactured lot, mid-range expiration lot and a lot nearing its expiration date. All lots were tested using SPSmedical's SporView<sup>®</sup> Culture Media (Tryptic Soy Broth, Modified with Bromocresol Purple). All lots tested resulted in a 72 hour reduced incubation time for the chemical vapor sterilization process when using the combination of the SPSmedical SporView<sup>®</sup> Bacterial Spore Strip and the SPSmedical's SporView<sup>®</sup> Culture Media.

#### Conclusion:

Supportive data has demonstrated that the SPSmedical SporView<sup>®</sup> Bacterial Spore Strip is equivalent to the legally marketed predicate device. Results of performance testing validate that the SPSmedical SporView<sup>®</sup> Bacterial Spore Strip provides a 72 hour reduced incubation time when used with the SPSmedical SporView<sup>®</sup> Culture Media in monitoring the Chemiclave<sup>®</sup> chemical vapor sterilization process.



MAY - 6 2004

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Mr. Gary J. Socola  
Director of Quality Assurance & Sterilization Projects  
SPSmedical Supply Corporation  
6789 West Henrietta Road  
Rush, New York 14543

Re: K041099  
Trade/Device Name: SPSmedical SporView<sup>®</sup> Bacterial Spore Strip  
Regulation Number: 880.2800  
Regulation Name: Sterilization Process Indicator  
Regulatory Class: II  
Product Code: FRC  
Dated: April 21, 2004  
Received: April 26, 2004

Dear Mr. Socola:

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 (301) 594-4618. 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/dsma/dsmamain.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 STATEMENT

510(k) Number (if known): K041099

Device Name: SPSmedical SporView® Bacterial Spore Strip

### Indications For Use:

The SPSmedical SporView® Bacterial Spore Strip is designed to monitor Chemiclave® chemical vapor sterilizers. The combination of SPSmedical's spore strip and SporView® Culture Media (Tryptic Soy Broth, Modified with Bromocresol Purple) has been validated at 60°C for a reduced incubation time of 72 hours for the Chemiclave® chemical vapor sterilization process.

Prescription Use \_\_\_\_\_  
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use X  
(21 CFR 807 Subpart C)

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

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Concurrence of CDRH, Office of Device Evaluation (ODE)

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

510(k) Number: K041099