

K071957

**510(k) Summary for the Pinnacle™ Pelvic Floor Repair Mesh**

**A. Sponsor**

Boston Scientific Corporation  
Urology and Gynecology Division  
100 Boston Scientific Way  
Marlborough, MA 01756

**B. Contact**

Michelle M. Berry  
Senior, Regulatory Affairs Specialist  
508-683-4941  
or  
Donna Gardner  
Director, Regulatory Affairs  
508-683-4398

NOV 08 2007

**C. Device Name**

Tradename: Pinnacle™ Pelvic Floor Repair Kits  
Common/usual name: Surgical Mesh  
Classification Name: FTL – Mesh, Surgical, Polymeric  
21 CFR 878.3300, Class II

**D. Predicate Device(s)**

Tradename: Polyform Synthetic Mesh  
Common/usual name: Surgical Mesh  
Classification Name: ~~OTF~~- Mesh, Surgical, Polymeric  
21 CFR 878.3300, Class II  
Premarket Notification: Proxy Biomedical, Ltd., K051245

**E. Device Description**

The proposed device is a sterile, single use device, consisting of a synthetic mesh assembly and needle holder. The mesh assembly consists of a polypropylene knitted mesh body with integrated legs that are protected by disposable polymer sleeves. At the distal end of the disposable polymer sleeve is a lead with needle designed for use with the currently legally marketed Capio™ Open Access Suture Capturing Device. The disposable lead was designed to facilitate the passage of the proposed mesh through bodily tissues for placement. The proposed mesh will be offered in three mesh models: Total, Anterior/Apical and Posterior designed for performing total vaginal repair, anterior vaginal defects and posterior and/or apical vaginal vault defects respectively.



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

Ms. Michelle M. Berry  
Senior Regulatory Affairs Specialist  
Boston Scientific Corporation  
100 Boston Scientific Way  
MARLBOROUGH MA 01752

SEP 28 2012

Re: K071957  
Trade/Device Name: Undetermined  
Regulation Number: 21 CFR 878.3300  
Regulation Name: Surgical mesh  
Regulatory Class: II  
Product Code: OTP  
Dated: October 3, 2007  
Received: October 4, 2007

Dear Ms. Berry:

This letter corrects our substantially equivalent letter of November 8, 2007.

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 (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 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.

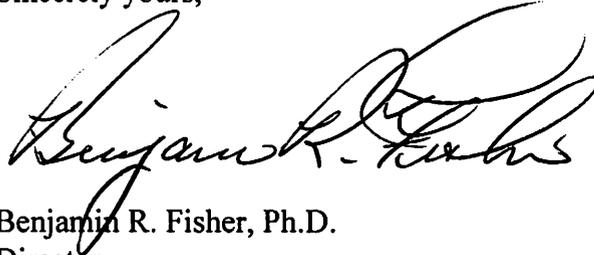
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must

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" (21 CFR 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,

A handwritten signature in black ink, appearing to read "Benjamin R. Fisher". The signature is fluid and cursive, with the first name being the most prominent.

Benjamin R. Fisher, Ph.D.  
Director  
Division of Reproductive, Gastro-Renal,  
and Urological Devices  
Office of Device Evaluation  
Center for Devices and Radiological Health

Enclosure

**Indications for Use Statement**

510(k) Number (if Known): K071957

Device Name: Undetermined

**Indications For Use:**

The Pinnacle™ Pelvic Floor Repair Kits are indicated for tissue reinforcement and stabilization of fascial structures of the pelvic floor for vaginal wall prolapse, where surgical treatment is intended, either as mechanical support or bridging material for the fascial defect.

Prescription Use  X  AND/OR Over-The-Counter Use \_\_\_\_\_  
(21 CFR 801 Subpart D) (21 CFR 801 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)

  
\_\_\_\_\_  
(Division Sign-ature)

**Division of General, Restorative,  
and Neurological Devices**

510(k) Number K071957