

Section 5 - 510(k) Summary

Date Prepared: January 9, 2013 FEB 28 2013
Company: Angiotech
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Contact: Kirsten Stowell
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Device trade name: Quill™ Polypropylene Knotless-Tissue Closure Device, Variable Loop Design

Device Common Name: Nonabsorbable polypropylene surgical suture

Device classification: Nonabsorbable polypropylene surgical suture
Product code, GAW
21 CFR 878.5010
Class II

Legally marketed devices to which the device is substantially equivalent:

K052373	Quill™ Nonabsorbable Polypropylene Barbed Suture
K113744:	Quill™ PDO Knotless Tissue-Closure Device (Polydioxanone), Variable Loop Design
K123409:	Quill™ Monoderm™ Knotless Tissue-Closure Device, Variable Loop Design

Description of the device: The Quill™ Polypropylene Knotless Tissue-Closure Device, Variable Loop Design is a sterile, synthetic nonabsorbable tissue-closure device that is intended for use in the closure of soft tissue, excluding closure of the epidermis. It is comprised of high molecular weight, isotactic polypropylene, undyed, or dyed blue with Phthalocyaninato (2-) Copper. The device is designed with small uni-directional barbs along the long axis of the suture monofilament which contains a welded primary loop and secondary loop design at the distal end. It is available in diameter Size 0 in various lengths affixed to various needle types.

Indications for Use: Quill™ Knotless Tissue-Closure Device comprised of Polypropylene is indicated for soft tissue approximation excluding closure of the epidermis.

Substantial Equivalence: The Quill™ Polypropylene Knotless Tissue-Closure device, Variable Loop Design is identical in material composition and intended use to the Quill™ Polypropylene Barbed Suture predicate. The proposed device is identical in design to the Quill™ PDO and Quill™ Monoderm™ Knotless Tissue-Closure Device, Variable Loop Design predicate devices.

Performance tests: Non-clinical laboratory performance testing was conducted to confirm that the Quill™ Polypropylene Knotless Tissue-Closure device, Variable Loop Design, conforms to the USP monograph for nonabsorbable sutures for tensile strength (as applicable) and needle attachment. This testing was performed in accordance with FDA's Class II Special Controls Guidance Document: Surgical Sutures, Issued June 3, 2003.

The results of this testing demonstrates that the Quill™ Polypropylene Knotless Tissue-Closure device, Variable Loop Design, is substantially equivalent to the predicate devices.



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

Surgical Specialties Corporation, DBA Angiotech
% Ms. Kirsten Stowell
Regulatory Affairs Manager
100 Dennis Drive
Reading, Pennsylvania 19606

February 28, 2013

Re: K130078

Trade/Device Name: Quill™ Polypropylene Knotless Tissue-Closure Device, Variable Loop Design

Regulation Number: 21 CFR 878.5010

Regulation Name: Nonabsorbable poly-propylene surgical suture

Regulatory Class: Class II

Product Code: GAW

Dated: January 09, 2013

Received: January 15, 2013

Dear Ms. Stowell:

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/ucml15809.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, . FOR

Peter  -S

Mark N. Melkerson
Acting Director
Division of Surgical Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Section 4 – Indications for Use Statement

510k number if known: K130078

Device Name: Quill™ Polypropylene Knotless Tissue-Closure Device, Variable Loop Design

Indications for Use:

Quill™ Knotless Tissue-Closure Device comprised of Polypropylene is indicated for soft tissue approximation excluding closure of the epidermis.

Prescription Use X
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use _____
(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)

David Krause

(Division Sign-Off)
Division of Surgical Devices
510(k) Number: K130078