DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

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

May 31, 2016

Spectranetics, Inc.
Kimberley Kline
Senior Manager, Regulatory Affairs
5055 Brandin Court
Fremont, CA 94538

Re: K142983
Trade/Device Name: AngioSculpt® PTA Scoring Balloon Catheter with HydroCross™ Coating
Regulation Number: 21 CFR 870.1250
Regulation Name: Percutaneous Catheter
Regulatory Class: Class II
Product Code: PNO
Dated: February 5, 2015
Received: February 6, 2015

Dear Ms. Kline:

This letter corrects our substantially equivalent letter of March 4, 2015.

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 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 contact the Division of Industry and Consumer Education 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. 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.

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Sincerely yours,

Misti L. Malone -S

for Bram D. Zuckerman, M.D.
Director
Division of Cardiovascular Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

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Sincerely yours,

Misti L. Malone -S

for Bram D. Zuckerman, M.D.
Director
Division of Cardiovascular Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

510(k) Summary for the AngioSculpt Scoring Balloon Catheter with HydroCross™ Coating

1. Submitter's Name / Contact Person

Submitter:	Spectranetics, Inc. 5055 Brandin Court Fremont, CA 94538
Contact Person:	Kimberley Kline Senior Manager, Regulatory Affairs Phone: 510-933-7989 Fax: 510-933-7994
Summary Preparation Date:	February 23, 2015

2. General Information

Trade Name:	AngioSculpt® PTA Scoring Balloon Catheter with HydroCross™ Coating
Common / Usual Name:	Angioplasty catheter
Classification Name:	Percutaneous catheter
Product Codes:	LIT
Predicate Devices:	AngioSculpt® Scoring Balloon Catheter (K112182/K122685/K133998)

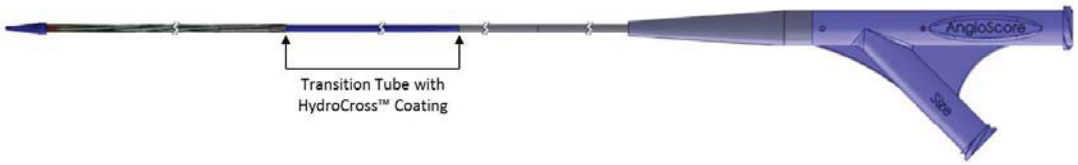
3. Indications for Use

The AngioSculpt PTA Scoring Balloon Catheter is intended for dilatation of lesions in the iliac, femoral, ilio-femoral, popliteal, infra popliteal, and renal arteries, and for the treatment of obstructive lesions of native or synthetic arteriovenous dialysis fistulae. Not for use in the coronary or neuro-vasculature.

4. Device Description

The AngioSculpt PTA Scoring Balloon Catheter with HydroCross™ Coating is a standard two-lumen catheter with a scoring balloon near the distal tip. The distal end of the catheter has a conventional nylon-blend balloon with a scoring element that wraps around the balloon. The scoring element creates focal concentrations of dilating force which minimizes balloon slippage and assists with luminal expansion of stenotic arteries. The balloon has radiopaque markers to aid in positioning the balloon in the stenosis, and is designed to provide an expandable segment of known diameter and length at a specific pressure.

As shown below, the catheter has a segment which is coated with a hydrophilic coating (HydroCross™ Coating).



5. Technological Characteristics

The 2.0-6.0mm x 100mm AngioSculpt catheters with HydroCross™ Coating incorporate substantially equivalent design, dimensional, and performance specifications when compared to the 510(k) cleared AngioSculpt catheters (K112182; 4.0-6.0x100/K122685; 2.0-3.5x100/K133998; 4.0-6.0x200). The following minor differences are noted:

- The scoring element incorporates the same type of inner rings that may aid in scoring element deployment as the catheters cleared under K133998.
- The hydrophilic coating cleared under K133998 was added to the transition tube of the catheters to improve lubricity and deliverability.
- The transition tube material was changed from Pebax 25 to Pebax 55 for manufacturability. This is the same material as cleared under K133998.
- The inner member of the subject catheters is of the same design as the currently marketed 2.0 – 6.0mm x 100mm product; however, the middle layer of the subject catheters will consist of two material options: plexar or orevac. The inner member outer layer of the 4.0 - 6.0mm x 100mm product will also consist of two material options: Pebax 6333 with black colorant or Vestamid Care ML21 2%-3% Clariant Blue (AF53676007). These materials are currently used in cleared AngioSculpt Scoring Balloon Catheters.

6. Summary of Bench Testing

Mechanical testing of the AngioSculpt catheter with HydroCross™ Coating was conducted in accordance with AngioScore's Risk Analysis and all applicable FDA guidance documents and relevant standards.

The following subset of bench tests were conducted to verify that design outputs met design requirements and to confirm proper function and durability. Test articles consisted of finished sterilized catheters.

- Catheter Diameter and Balloon Profile,
- Device Preparation, Deployment and Retraction
- Balloon Inflation and Deflation Time
- Catheter Bond (Tensile) Strength
- Tip Pull Strength

- Catheter Diameter and Balloon Profile (with Scoring Element)
- Torque Strength
- Pushability, Trackability and Secure Edges
- Guidewire Compatibility
- Catheter Effective Length
- Catheter Surface Appearance
- Luer Compatibility
- Coating Length
- Coating Thickness
- Coating Lubricity
- Coating Integrity
- Particulate Evaluation
- Rated Burst Pressure
- Compliance
- Device Fatigue
- Flexibility and Kink

7. Summary of Biocompatibility Testing

The materials of construction for the2.0-6.0mm x 100mm AngioSculpt catheters with HydroCross™ Coating are the same materials as those of the predicate AngioSculpt catheters. Biocompatibility testing of predicate devices cleared under K122685 and K133998 is applicable to the 2.0-6.0mm x 100mm AngioSculpt catheters with HydroCross™ Coating; the following is a summary.

The AngioSculpt catheter with HydroCross™ Coating is categorized as an "External communicating device in contact with circulating blood with limited exposure time". The biocompatibility of the device was assessed in accordance with ISO 10993-1:2009 – *Biological evaluation of medical devices, Part 1 – Evaluation and tests within a risk management process*.

The biocompatibility tests listed below, except Thrombosis (*in-vivo*), were conducted in accordance with the provisions of the FDA GLP regulations 21 CFR Part 58. Thrombogenicity was evaluated as part of a GLP animal study.

- Cytotoxicity - MEM Elution Test using L-929 Mouse Fibroblast cells (ISO)
- Sensitization - Guinea Pig Maximization Sensitization Test
- Irritation - Intracutaneous Reactivity (Irritation) Test
- Systemic Toxicity - Acute Systemic Injection Test (ISO)
- Systemic Toxicity - Material Mediated Pyrogens
- Hemocompatibility - Partial Thromboplastin Time (PTT)
- Hemocompatibility - Hemolysis Direct Contact/Extract Method (ASTM)
- Hemocompatibility - Direct Contact Complement Activation Testing With C3a and SC5b
- Hemocompatibility - Thrombosis (in vivo)
- Genotoxicity - Reverse Bacterial Mutation
- Genotoxicity - In vitro Mouse Lymphoma Assay-Extended Treatment
- Genotoxicity - Rodent bone Marrow Micronucleus Assays

The biocompatibility test results confirm that the AngioSculpt catheter with HydroCross™ Coating, including the minor material changes, is non-cytotoxic, non-sensitizing, non-irritating, not systemically toxic, non-hemolytic, and non-mutagenic , when evaluated under the respective test conditions. As shown in the GLP animal study, no thrombo-embolism was observed when the subject catheters were evaluated under simulated use conditions.

8. Substantial Equivalence Comparison

The subject catheters share the same intended use, principles of operation, overall technical and functional capabilities, packaging and sterilization process, and similar design and materials as the predicate AngioSculpt catheters and are therefore substantially equivalent.

Although there are minor differences between the subject catheters and its predicate devices those differences do not raise new questions of safety or efficacy. Design verification and validation testing demonstrated adequate device performance and confirmed that no new questions of safety or effectiveness for peripheral balloon angioplasty devices were raised. The changes to the subject catheters do not affect the intended use of the device, alter the fundamental scientific technology of the device, or raise new issues of safety and effectiveness. The subject AngioSculpt PTA Scoring Balloon Catheters with HydroCross™ Coating are therefore, substantially equivalent to the predicate catheters.