FEB 20 2014

NEOMETRICS

510(k) Summary

Submitter:	NeoMetrics, Inc. 2605 Fernbrook Lane North, Suite J Plymouth, MN 55447
Contact Person:	Eugene Champeau, President 2605 Fernbrook Lane North, Suite J Plymouth, MN 55447
Date Prepared:	November 8, 2013
Trade Name:	NovaGold™ High Performance Guidewire
Common Name	ERCP Guidewire
Classification Name	Endoscope and accessories (21 CFR 876.1500, Product Code OCY)
Classification:	Class II
Predicate Device:	The subject device is equivalent to K122816; Endscopic Wire Guide manufactured by Wilson-Cook Medical.
Device Description:	The NovaGold Guidewire is constructed from a steerable, metallic core with a PTFE polymer coating over the shaft. A hydrophilic coating is applied over the distal portion of the device. The guidewire has a radiopaque, floppy tip.
Indication for Use:	The NovaGold Guidewire is intended for use in selective cannulation of the biliary ducts including the common bile, pancreatic, cystic, right and left hepatic ducts, and to aid in the placement of diagnostic and therapeutic devices during endoscopic procedures.

Functional and Safety Testing:	To verify that device design met functional and performance requirements, representative samples of the device underwent bench testing in accordance to applicable standards and guidances. These data provides an acceptable assurance of the safety and effectiveness of the NovaGold guidewire and demonstrated the device is equivalent to the predicate.

Comparative Technology Characteristics	A comparison of the characteristics of the proposed device and the predicate device shows the NovaGold guidewire to have the same or similar technological characteristics to the predicate which has received 510(k) clearance.
	
	• Same intended use
	• Same operating principle
	• Same packaging and sterilization process
	• Similarities in design, material types, and technology include
	  o Nominal diameter: .018"
	  o Similar lengths: 260 and 480 cm
	  o Nitinol alloy core wires
	  o Distal radiopaque tip
	  o Lubricious coatings

Non-Clinical Tests Submitted	The following tests were performed to support NovaGold's substantial equivalence. Biocompatibility testing • Cytotoxicity, L-929 MEM Elution • Cytotoxicity, Colony Microassay by Elution • Guinea Pig Maximization Sensitization • Irritation, Intracutaneous Reactivity • Chemical Characterization Bench testing • Coating adherence • Corrosion resistance • Dimensional measurements • Fracture resistance • Tensile strength • Torqueability Shelf Life Testing • 13 month accelerated aging

Conclusion:	NeoMetrics Inc. considers the NovaGold guidewire to be equivalent to the predicate device. This conclusion is based upon the fact that device has an equivalent intended use, and there are no differences that raise new types of questions of safety and effectiveness.

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

February 20, 2014

NeoMetrics, Inc.
David Liebl
President and Chief Technology Officer
2605 Fernbrook Lane North, Suite J
Plymouth, MN 55447

Re: K133076
Trade/Device Name: NovaGold™ High Performance Guidewire
Regulation Number: 21 CFR§ 876.1500
Regulation Name: Endoscope and accessories
Regulatory Class: II
Product Code: OCY
Dated: January 21, 2014
Received: January 24, 2014

Dear David Liebl,

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/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" (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,

Benjamin R. Fisher -S

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

5.0 Indications for Use Statement

510(k) Number K133076

Device Name: NovaGold™ High Performance Guidewire

Indications for Use: The NovaGold Guidewire is intended for use in selective cannulation of the biliary ducts including the common bile, pancreatic, cystic, right and left hepatic ducts, and to aid in the placement of diagnostic and therapeutic devices during endoscopic procedures.

Prescription Use __X_____ (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)

Benjamin R. Fisher -S
2014.02.20 15:08:06 -05'00'