CRESCENT™ Spinal System
510(k) Summary
April 23, 2010

I. **Company:** **Medtronic Sofamor Danek**
1800 Pyramid Place
Memphis, TN 38132
(901) 396-3133

APR 2 6 2010

Contact: **Jennifer Hackney**
Regulatory Affairs Specialist

II. **Proprietary Trade Name: CRESCENT™ Spinal System**

III. **Classification Name:** Intervertebral Body Fusion Device (21 CFR 888.3080)

IV. **Product Code: MAX**

V. **Product Description**

The subject CRESCENT™ Spinal System consists of a variety of hollow intervertebral body spacers featuring a bullet-nosed, anatomically shaped design with axial voids designed to hold autogenous bone graft. The subject devices are designed with diamond V teeth across both superior and inferior surfaces to allow the implant to grip the superior and inferior end plates, thus allowing expulsion resistance. The devices are designed to be implanted through the transforaminal and direct lateral approaches. The devices range from 7mm to 15mm in height and from 25mm to 36mm in length. The devices are manufactured from Medical Grade PEEK (polyetheretherketone). These devices also contain Tantalum markers used for imaging purposes.

V. **Indications**

The CRESCENT™ Spinal System is indicated for interbody fusion with autogenous bone graft in patients with degenerative disc disease (DDD) at one or two contiguous levels from L2 to S1. These DDD patients may also have up to Grade 1 spondylolisthesis or retrolisthesis at the involved levels. DDD is defined as discogenic back pain with degeneration of the disc confirmed by history and radiographic studies. These patients should be skeletally mature and have had six months of non-operative treatment. These devices are intended to be used with supplemental fixation instrumentation which has been cleared by the FDA for use in the lumbar spine.

VI. Identification of the Legally Marketed Predicate Devices Used to Claim Substantial Equivalence

Documentation was provided which demonstrated that the subject CRESCENT™ Spinal System is substantially equivalent to several interbody cages, including the CAPSTONE® Spinal System (K073291, SE 04/24/08), the LT-CAGE® PEEK Lumbar Tapered Fusion Device (P970015, Medtronic Sofamor Danek, Approved 9/10/03), the RAY® Threaded Fusion Cage (P950019, Stryker, Approved 9/4/03), and the BRANTIGEN I/F CAGE® (DePuy, P960025), as well as to the VERTE-STACK® Spinal System (K052261, SE 10/07/05).

VII. Brief Discussion of the Non-Clinical Tests Submitted

The following mechanical tests of the subject CRESCENT™ Spinal System were performed:

- static axial compression testing in accordance with ASTM 2077-03;
- dynamic axial compression testing in accordance with ASTM 2077-03;
- static compression shear testing in accordance with ASTM 2077-03;
- compression shear fatigue testing in accordance with ASTM 2077-03; and
- subsidence testing in accordance with ASTM F2267-04.

VIII. Conclusions Drawn from the Non-Clinical Tests

Results of mechanical testing indicated that all acceptance criteria were met, demonstrating substantial equivalence to the listed predicate devices.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

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

Medtronic Sofamor Danek
% Ms. Jennifer Hackney
Regulatory Affairs Specialist
1800 Pyramid Place
Memphis, Tennessee 38132

APR 2 6 2010

Re: K094025
Trade/Device Name: CRESCENT™ Spinal System
Regulation Number: 21 CFR 888.3080
Regulation Name: Intervertebral body fusion device
Regulatory Class: Class II
Product Code: MAX
Dated: April 15, 2010
Received: April 16, 2010

Dear Ms. Hackney:

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.

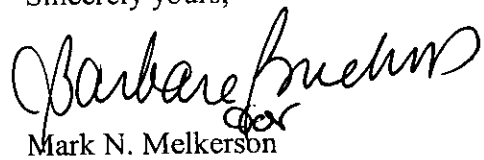
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's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



Mark N. Melkerson
Director
Division of Surgical, Orthopedic
And Restorative Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number (if known): K094025

Device Name: CRESCENT™ Spinal System

Indications for Use:

The CRESCENT™ Spinal System is indicated for interbody fusion with autogenous bone graft in patients with degenerative disc disease (DDD) at one or two contiguous levels from L2 to S1. These DDD patients may also have up to Grade 1 spondylolisthesis or retrolisthesis at the involved levels. DDD is defined as discogenic back pain with degeneration of the disc confirmed by history and radiographic studies. These patients should be skeletally mature and have had six months of non-operative treatment. These devices are intended to be used with supplemental fixation instrumentation which has been cleared by the FDA for use in the lumbar spine.

Prescription Use ____X__ (Part 21 CFR 801 Subpart D) AND/OR Over-The-Counter Use _____ (21 CFR 807 Subpart C)

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of Surgical, Orthopedic,
and Restorative Devices

510(k) Number K094025