



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

Nexxt Spine LLC  
% Ms. Karen Warden  
President  
Backroads Consulting  
P.O. Box 566  
Chesterland, Ohio 44026-2141

April 26, 2016

Re: K153453  
Trade/Device Name: Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System  
Regulation Number: 21 CFR 888.3070  
Regulation Name: Pedicle Screw Spinal System  
Regulatory Class: Class III  
Product Code: NKB, KWP, MNI, MNH, OSH  
Dated: March 25, 2016  
Received: March 28, 2016

Dear Ms. Warden:

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 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" (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 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>.

Sincerely yours,

**Mark N. Melkerson -S**

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

Enclosure

## Indications for Use

510(k) Number (*if known*)

K153453

Device Name

Inertia® Pedicle Screw and Deformity Correxxion™ System

Indications for Use (*Describe*)

The Inertia® Pedicle Screw and Deformity Correxxion™ System is intended for pedicle and non-pedicle immobilization and stabilization of the posterior non-cervical spine (T1-S2/Ilium) in skeletally mature patients as an adjunct to fusion for the following indications: degenerative disc disease (defined as back pain of discogenic origin with degeneration of the disc confirmed by history and radiographic studies), spondylolisthesis, trauma (i.e., fracture or dislocation), spinal stenosis, curvatures (i.e., scoliosis, kyphosis, or lordosis), tumor, pseudarthrosis, and/or failed previous fusion.

When used for posterior non-cervical pedicle screw fixation in pediatric patients, the Inertia® Pedicle Screw and Deformity Correxxion™ System implants are indicated as an adjunct to fusion to treat adolescent idiopathic scoliosis. The Inertia® Pedicle Screw and Deformity Correxxion™ System is to be used with autograft and/or allograft. Pediatric pedicle screw fixation is limited to a posterior approach.

Type of Use (*Select one or both, as applicable*)

Prescription Use (Part 21 CFR 801 Subpart D)

Over-The-Counter Use (21 CFR 801 Subpart C)

**PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON A SEPARATE PAGE IF NEEDED.**

**FOR FDA USE ONLY**

Concurrence of Center for Devices and Radiological Health (CDRH) (*Signature*)

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This section applies only to requirements of the Paperwork Reduction Act of 1995.

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## **510(k) Summary**

**Date:** 30 November 2015

**Sponsor:** Nexxt Spine  
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Noblesville, IN 46060  
Office: 317.436.7801  
Fax: 317.245.2518

**Sponsor Contact:** Andy Elsbury, President

**510(k) Contact:** Karen E. Warden, PhD  
BackRoads Consulting Inc.  
PO Box 566  
Chesterland, OH 44026  
Office: 440.729.8457

**Proposed Trade Name:** The Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System

**Common Name:** Posterior pedicle screw & hook system

**Device Classification:** Class III

**Classification Name:** Pedicle screw spinal system, Spinal interlaminar fixation orthosis

**Regulations:** 888.3070, 888.3050

**Device Product Codes:** NKB, KWP, MNI, MNH, OSH

**Device Description:** The Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System consists of longitudinal members (rods), anchors (screws and hooks), interconnections (offset, rod-rod and crosslink) and fasteners in a variety of sizes to accommodate differing anatomic requirements.  
The Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System implants are sold sterile and non-sterile.

**Indications for Use:** The Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System is intended for pedicle and non-pedicle immobilization and stabilization of the posterior non-cervical spine (T1-S2/Ilium) in skeletally mature patients as an adjunct to fusion for the following indications: degenerative disc disease (defined as back pain of discogenic origin with degeneration of the disc confirmed by history and radiographic studies), spondylolisthesis, trauma (i.e., fracture or dislocation), spinal stenosis, curvatures (i.e., scoliosis, kyphosis, or lordosis), tumor, pseudarthrosis, and/or failed previous fusion.  
When used for posterior non-cervical pedicle screw fixation in pediatric patients, the Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System implants are indicated as an adjunct to fusion to treat adolescent idiopathic scoliosis. The Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System is to be used with autograft and/or allograft. Pediatric pedicle screw fixation is limited to a posterior approach.

**Materials:** Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System from Ti-6Al-4V ELI titanium alloy (ASTM F136) and Cobalt Chrome (per ASTM F1537).

**Predicate Devices:** Primary: CD Horizon<sup>®</sup> Spinal System (Medtronic Sofamor Danek – K152457)  
Additional: Tiger<sup>®</sup> Spine System (Corelink, LLC – K133369) and Inertia<sup>®</sup> Pedicle Screw System (Nexxt Spine – K141376)

**Reference Devices:** Moss Miami<sup>™</sup> Spinal System (DePuy AcroMed, Inc. – K022623), Synergy VLS Open (Interpore Cross International – K011437) and Inertia<sup>®</sup> Pedicle Screw System (Nexxt Spine – K101278)

**Performance Data:** Mechanical testing of worst case Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System constructs included static and dynamic compression bending and static torsion according to ASTM F1717.

The mechanical test results demonstrate that Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System performance is substantially equivalent to the predicate devices.

**Technological Characteristics:** The Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System possesses the same technological characteristics as one or more of the predicate devices. These include:

- intended use (as described above)
- basic design (rod and screw configuration),
- material (titanium alloy),
- sizes (dimensions are comparable to those offered by the predicate systems) and

The fundamental scientific technology of the Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System is the same as previously cleared devices.

**Conclusion:** The Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System possesses the same intended use and technological characteristics as the predicate devices. Therefore Inertia<sup>®</sup> Pedicle Screw and Deformity Correxxion<sup>™</sup> System is substantially equivalent for its intended use.