

510(k) Summary

[in accordance with SMDA of 1990, 21 CFR 807.92(c)]

Contact: Mr. Hartmut Loch
Vice President, Regulatory Affairs
Allez Spine, LLC.
2301 Dupont Drive, Suite 510
Irvine CA 92612

Trade name: Allez Spine *Del Mar*® Polyaxial Pedicle Screw System

Common name: Spinal Fixation System

Classification name: Appliance, Fixation, Spinal Interlaminar - § 888.3050 (KWP) – class II
Orthosis, Spinal Pedicle Fixation - § 888.3070 (MNI) – class II
Orthosis, Spondylolisthesis Spinal Fixation - § 888.3070 (MNH) – class II
Orthosis, Spinal Pedicle Fixation, For Degenerative Disc Disease
- § 888.3070 (NKB) – class III

All Orthopedic Device Panel 87

Product Code: KWP, MNI, MNH & NKB

Device Description and Characteristics: The *Del Mar*® Monoaxial Pedicle Screw System is intended to help provide correction, immobilization and stabilization of spinal segments as an adjunct to fusion of the thoracic, lumbar and/or sacral space.

The DEL MAR® Monoaxial Pedicle Screw System consists of a variety of rods and screws, which can be rigidly locked into a variety of configurations, with each construct being tailor made for the individual case. Monoaxial screws are supplied in winged and non-winged configurations, in a variety of different length, and in 5mm, 6mm, and 7mm diameter sizes. All sizes are able to receive 5.5mm connecting rods only.

The DEL MAR® Monoaxial Pedicle Screw System implant components are fabricated from medical grade titanium alloy per ASTM F136.

Equivalence: The modified *Del Mar*® Monoaxial Pedicle Screw System is substantially equivalent to the original LAGUNA® Monoaxial Pedicle Screw System, which is manufactured and marketed by Allez Spine, LLC.

Indications: The DEL MAR® Monoaxial Pedicle Screw System is intended for posterior, non-cervical fixation for the following conditions: 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 and/or lordosis); tumor pseudarthrosis; and/or failed previous fusion. The DEL MAR® Monoaxial Pedicle Screw System is intended to be used with autograft and/or allograft as an adjunct to fusion.

Performance data: Biomechanical tests have been performed. The test results were equivalent to other similar implants and are sufficient for *in vivo* loading.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Allez Spine, LLC
% Mr. Hartmut Loch
Vice President, Regulatory Affairs
2301 Dupont Drive, Suite 510
Irvine, California 92612

JUL - 9 2009

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Re: K091219

Trade/Device Name: DEL MAR[®] Monoaxial Pedicle Screw System
Regulation Number: 21 CFR 888.3070
Regulation Name: Pedicle screw spinal system
Regulatory Class: III
Product Code: NKB, MNH, MNI, KWP
Dated: June 4, 2009
Received: June 9, 2009

Dear Mr. Loch:

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.

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.

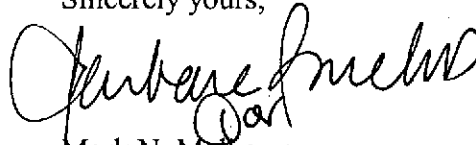
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's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

Page 2 – Mr. Hartmut Loch

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/cdrh/mdr/> 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 (240) 276-3150 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,

A handwritten signature in black ink, appearing to read "Mark N. Meikerson". The signature is written in a cursive style with a large initial "M".

Mark N. Meikerson

Director

Division of Surgical, Orthopedic,
and Restorative Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

Enclosure

