

K082509

510(k) Summary**Submitter:**

Korea Bone Bank Co., Ltd.
Shim Young Bok
#402 Acetechnotower 9th,
345-30, Gasandong, Keumcheongu,
Seoul, South Korea
Phone: 82-02-2404-0475
Fax: 82-02-2104-0478

Contact:**DEC 15 2008**

Kodent Inc.
Jung Bae Bang
13340 E. Firestone Blvd. Suite J
Santa Fe Springs, CA 90670
Email: kodentinc@kodent.co.kr
Phone: 562-404-8466
Fax: 562-404-2757

Device Information

Trade Name: EOS Spinal System
Common Name: Pedicle Screw Spinal Fixation System
Classification Name: Spinal Pedicle Fixation
Spondylolisthesis Spinal Fixation
Product Code: MNH, MNI
Regulation Number: 888.3070

Indication for Use:

The EOS Spinal System is a posterior pedicle screw system indicated for the treatment of severe Spondylolisthesis (Grade 3 and 4) of the L5-S1 vertebra in skeletally mature patients receiving fusion by autogenous bone graft having implants attached to the lumbar and sacral spine (L3 to sacrum) with removal of the implants after the attainment of a solid fusion.

In addition, the EOS Spinal System is intended to provide immobilization and stabilization of spinal segments in skeletally mature patients as an adjunct to fusion in the treatment of the following acute and chronic instabilities or deformities of the thoracic lumbar and sacral spine: degenerative Spondylolisthesis with objective evidence of neurological impairment, fracture, dislocation, scoliosis, kyphosis, spinal tumor and failed previous fusion (pseudarthrosis).

Device Description:

The EOS Spinal System is a top-loading multiple component, posterior spinal fixation system which consists of pedicle screws, rods, set screws, connectors, and a transverse (cross) linking mechanism.

The EOS Spinal System will allow surgeons to build a spinal implant construct to stabilize and promote spinal fusion. The EOS Spinal System implant components are supplied non-sterile, single use and fabricated from titanium alloy (Ti-6Al-4V ELI) that conforms to ASTM F136. Various sizes of these implants are available. Specialized instruments are available for the application and removal of the EOS Spinal System.

Materials:

The devices are manufactured from Ti6Al-4V ELI alloy per ASTM and ISO Standards.

Performance Data:

Performance data per ASTM F1717 were submitted to characterize the subject EOS Spinal System components addressed in this notification.

Predicate Devices:

The subject device is substantially equivalent to the following predicate devices:

- * 4CIS® Vane Spine System (Solco Biomedical Co., Ltd.; K060702)
- * GSS Pedicle Screw System (GS Medical Co., Ltd.; K053573)

Comparison to Predicate Devices:

Testing and other comparisons have established that the subject of EOS Spinal System is substantially equivalent in design, materials, indications and intended use, packaging, labeling, and performance to other predicate devices of the type currently marketed in the U.S.

This summary of 510(k) safety and effectiveness information is being submitted in accordance with the requirements of 21 CFR & 807.93



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Korea Bone Bank, Co., Ltd.
% Kodent Inc.
Jung Bae Bang
13340 E. Firestone Boulevard, Suite J
Santa Fe Springs, California 90670

DEC 15 2008

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Re: K082509

Trade/Device Name: EOS Spinal System
Regulation Number: 21 CFR 888.3070
Regulation Name: Pedicle screw spinal system
Regulatory Class: II
Product Code: MNI, MNH
Dated: December 5, 2008
Received: December 5, 2008

Dear Jung Bae Bang:

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 such 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); 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.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Center for Devices and Radiological Health's (CDRH's) Office of Compliance at (240) 276-0120. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding postmarket surveillance, please contact CDRH's Office of Surveillance and Biometric's (OSB's) Division of Postmarket Surveillance at (240) 276-3474. For questions regarding the reporting of device adverse events (Medical Device Reporting (MDR)), please contact the Division of Surveillance Systems at (240) 276-3464. You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at toll-free number (800) 638-2041 or (240) 276-3150 or the Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,



Mark N. Melkerson
Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indication for Use

510(K) Number (if known): _____

Device Name: EOS Spinal System

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Prescription Use _____

AND/OR

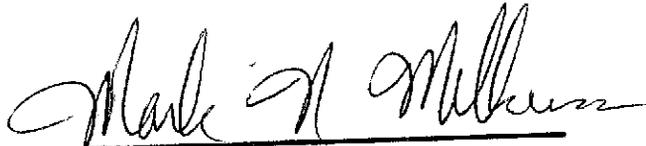
Over-The-Counter _____

(Part 21 CFR 801 Subpart D)

(Per 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)



**(Division Sign-Off)
Division of General, Restorative,
and Neurological Devices**

510(K) Number K082509

