zimmer
dental

Zimmer Dental
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510k No.: K071439

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Traditional 510(k)
PRE-MARKET NOTIFICATION

SEP 20 2007

510(k) SUMMARY (21CFR807.92(a))

1. Submitter's Information:

Name:	Zimmer Dental Inc.
Address:	1900 Aston Ave. Carlsbad, CA 92008-7308 USA
Phone:	760-929-4300
Contact:	Erin L. McVey
Date Prepared:	September 12, 2007

2. Device Name*: Patient-Specific Abutment, Internal Hex, Titanium

**Device trade name not available at time of submission.*

Device Classification Name: Endosseous Dental Implant Abutment

3. Predicate Device: Zimmer® Hex-Lock Abutment, 3.5mm (cat. No. HLA3/3)

4. Device Description:

The Zimmer® Patient-Specific Abutment is a titanium alloy abutment for use with endosseous dental implants to provide support for prosthetic devices. The abutment is manufactured using individual patient-specific requirements to create a design that facilitates functional, as well as, esthetic restoration. Ensure the implant size and abutment angulation are appropriate for the occlusal load. Highly angulated abutments should be avoided in the posterior region.

5. Intended Use:

The Zimmer® Patient Specific Abutment is used as a terminal or intermediate abutment for a cemented prosthesis. The abutment can be used for a single or multiple-unit restoration.

6. Device Comparison:

The new device is substantially equivalent to the predicate relative to material and general design features. In addition, the new device is substantially equivalent to the predicate as evidenced in mechanical testing.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

SEP 20 2007

Ms. Erin L. McVey, RAC(US)
Senior Regulatory Affairs Specialist
Zimmer Dental, Incorporated
1900 Aston Avenue
Carlsbad, California 92008

Re: K071439
Trade/Device Name: Zimmer® Patient-Specific Abutment, Internal Hex, Titanium
Regulation Number: 21 CFR 872.3630
Regulation Name: Endosseous Dental Implant Abutment
Regulatory Class: II
Product Code: NHA
Dated: September 10, 2007
Received: September 12, 2007

Dear Ms. McVey:

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 Office of Compliance at (240) 276-0115. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). 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) 443-6597 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,

Chiu Lin, Ph.D.
Director
Division of Anesthesiology, General Hospital,
Infection Control and Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

K071439

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Indications for Use

510(k) Number (if known): K071439

Device Name: **Zimmer® Patient-Specific Abutment, Internal Hex, Titanium**

Indications For Use:

The Zimmer® Patient Specific Abutment, Internal Hex, Titanium is used as a terminal or intermediate abutment for a cemented prosthesis. The abutment can be used for a single or multiple-unit restoration.

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

Susan Runner

(Division Sign-Off)
Division of Anesthesiology, General Hospital
Infection Control, Dental Devices

510(k) Number: K071439