Zimmer Dental
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Carlsbad, CA 92008
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Traditional 510(k)
PRE-MARKET NOTIFICATION 510(k)

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

1. Submitter's Information:

Name:	Zimmer Dental Inc.
Address:	1900 Aston Ave. Carlsbad, CA 92008
Phone:	760-929-4300
Contact:	Julie Lamothe
Date Prepared:	April 28, 2014

2. Device Name:

Trade Name:	Zimmer Zfx Titanium Abutment for Straumann Bone Level Implant System
Regulation Number:	872.3630
Classification Code:	NHA
Device Classification Name:	Abutment, Implant, Dental, Endosseous

3. Predicate Device(s):

Predicate Device No. 1

Trade Name:	Zimmer Zfx Abutment for NobelReplace Implant System
510(k) Number:	K120873
Regulation Number:	872.3630
Classification Code:	NHA
Device Classification Name:	Abutment, Implant, Dental, Endosseous

Predicate Device No. 2

Trade Name:	Straumann Cares Abutment, Titanium
510(k) Number:	K072151 and K081005
Regulation Number:	872.3630
Classification Code:	NHA
Device Classification Name:	Abutment, Implant, Dental, Endosseous

Predicate Device No. 3

Trade Name:	Straumann Bone Level Anatomic Abutment
510(k) Number:	K071357
Regulation Number:	872.3630
Classification Code:	NHA
Device Classification Name:	Abutment, Implant, Dental, Endosseous

4. Device Description:

The Zimmer Zfx Titanium Abutment for the Straumann Bone Level implant system is designed for use with Straumann Bone Level implants to support single or multi tooth restorations. The abutment/implant interface is a conical connection with 4 grooves for insertion guidance.

The Zimmer Zfx Titanium Abutment for Straumann Bone Level Implant System is a patient specific dental implant abutment with a competitor compatible interface. The purpose of a Patient-Specific abutment is to satisfy customer needs that are otherwise difficult to meet with off-the-shelf abutments. They can be manufactured in multiple sizes, shapes, and angles within the limits established in this submission. They frequently incorporate the modifications typically done at a dental laboratory or "chair-side" by a dentist. Traditional methodologies require the customer (dentist/laboratory technician) to begin with a "stock" abutment and use manual subtractive techniques to remove material from this original "stock" design. However, a Patient-Specific abutment will incorporate these same modifications desired by the customer (dentist/laboratory technician) at the time of fabrication at the manufacturing facility.

The engineering drawings list ranges in areas (attributes) of the abutment that may be modified depending upon patient-specific needs.

The abutment is composed of Titanium alloy (Ti6Al4V), and secured to the implant with a separate Titanium alloy screw for retention.

The new abutment will be offered in Narrow CrossFit (NC) Ø 3.3mm and Regular CrossFit (RC) Ø 4.1mm or Ø 4.8mm implant connection sizes.

5. Indications for Use:

The Zimmer Zfx Titanium Abutment for Straumann Bone Level Implant System is designed for use as a terminal or intermediate abutment for cement retained prostheses. The abutment can be used with Straumann Bone Level implants with a Narrow Crossfit Connection (NC) Ø 3.3mm or a Regular Crossfit Connection (RC) Ø 4.1mm or Ø 4.8mm.

6. Device Comparison:

The new abutment device is substantially equivalent to the predicates relative to material, manufacturing process and general design features. It is fabricated from Titanium alloy and utilizes a conical connection implant/abutment interface with 4 grooves for insertion guidance, which is identical in size and

shape (for a given platform diameter) to the predicate device. The new abutment and base device will be affixed to the implant by a retaining screw, the same manner as the predicate.

7. Technological Characteristics

Feature	**New Device** Zimmer Zfx Titanium Abutment for Straumann Bone Level Implant System	**Predicate #1** Zimmer Zfx Abutment for NobelReplace Implant System	**Predicate #2** Straumann Cares Abutment, Titanium	**Predicate #3** Straumann Bone Level Anatomic Abutment
Material	Titanium 6Al-4V	Titanium 6Al-4V	Titanium	Titanium
Implant Interface	Conical Connection with 4 grooves	Internal Tri-lobe	Conical Connection with 4 grooves	Conical Connection with 4 grooves
Emergence	Contoured, curved	Contoured, curved	Contoured, curved	Contoured, curved
Margin	Pre-machined	Pre-machined	Pre-machined	Pre-machined
Platform Diameter	3.3 (NC), 4.1/4.8 (RC)	3.5mm, 4.3mm, 5.0mm, 6.0mm	3.3 (NC), 4.1/4.8 (RC)	3.3 (NC), 4.1/4.8 (RC)
Cuff Width	2.8mm-8mm (NC) 3.3mm-10.0mm (RC)	3.5mm-12.0mm	10mm MAX (NC) 13mm MAX (RC)	4.0mm (NC) 6.5mm (RC)
Cone Angle	0-30°	0-20°	0-30°	30°
Minimum Height	3.0mm MIN Cone 3.0mm-12.0mm Overall	3.0- 11.5mm Cone 3.5-12.0mm Overall	17mm MAX Overall	5.5mm Cone 7.5mm and 9.0mm overall
Retaining Screw	New device	ZFX09000642	Cat No. 025.2900 and 025.4900	Cat No. 025.2900 and 025.4900

8. Non-Clinical Testing:

Non-clinical test data was used to support the decision of substantial equivalence. This consisted of reverse engineering, compatibility analysis and mechanical fatigue testing. The Straumann Bone Level interface was reverse engineered based on actual measurements taken from Straumann Bone Level implants, abutments and retaining screws in order to assure that the Zimmer Zfx Titanium Abutment is compatible with Straumann Bone Level Implant Systems. Dimensional specifications were developed for the Zimmer fabricated components based on the reverse engineering results. A tolerance analysis as well as a rotational analysis was conducted to illustrate the nature of fit between the Zimmer fabricated parts and the OEM implant. To verify the compatibility of the Zimmer device to the OEM device, fatigue testing was completed using Zimmer fabricated abutments assembled to OEM implants in accordance with the FDA guidance Class II Special Controls Guidance Document: Root-form Dental Implants and Endosseous Dental Implant Abutments. The results were compared to fatigue testing data of the predicate #3 device. Based on the reverse

engineering process, as well as verification of the final connection dimensions and tolerances, the Zimmer Zfx Titanium Abutment for Straumann Bone Level Implant System can be deemed compatible with the Straumann Bone Level implant interface.

The Zimmer Zfx Titanium Abutment for the Straumann Bone Level Implant System will be sold non-sterile and will be sterilized by the end user. The sterilization procedures listed in the Instruction For Use were validated to provide a minimum sterility assurance level of 10^{-6}.

Additionally, Zimmer Dental implant systems were evaluated for interactions with magnetic fields during Magnetic Resonance Imaging (MRI) in accordance with the FDA Guidance: Establishing Safety and Compatibility of Passive Implants in the Magnetic Resonance (MR) Environment. This was done to determine that the presence of the abutment poses no additional restrictions on MRI beyond those that would otherwise occur for the patient.

9. Clinical Testing

No clinical testing was performed. Non-clinical testing was used to support the decision of safety and effectiveness.

10. Conclusion

Based on our analysis, the device is substantially equivalent to the predicate.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

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

July 29, 2014

Zimmer Dental, Inc.
Julie Lamothe, Ph.D
Regulatory Affairs Manager
1900 Aston Ave.
Carlsbad, CA 92008-7308

Re: K141120
Trade/Device Name: Zimmer Zfx Titanium Abutment for Straumann Bone Level Implant System
Regulation Number: 21 CFR 872.3630
Regulation Name: Endosseous Dental Implant Abutment
Regulatory Class: II
Product Code: NHA
Dated: April 21, 2014
Received: April 30, 2014

Dear Dr. Lamothe:

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 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. 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,

Mary S. Runner -
S

Erin I. Keith, M.S.
Director
Division of Anesthesiology, General Hospital,
Respiratory, Infection Control and
Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

zimmer
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Indications for Use

510(k) Number (if known): K141120

Device Name: ***Zimmer Zfx Titanium Abutment for Straumann Bone Level Implant System***

Indications For Use:

The Zimmer Zfx Titanium Abutment for Straumann Bone Level Implant System is designed for use as a terminal or intermediate abutment for cement retained prostheses. The abutment can be used with Straumann Bone Level implants with a Narrow CrossFit Connection (NC) Ø3.3mm or Regular Crossfit Connection (RC) Ø4.1mm or Ø4.8mm.

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)

Sheena A. Green -S
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