K072057

1 of 1

510(k) SUMMARY

AUG - 6 2007

DENTSPLY International
Susquehanna Commerce Center West
221 West Philadelphia Street, Suite 60
York, PA 17405-0872

CONTACT: Helen Lewis

DATE PREPARED: July 18, 2007

TRADE OR PROPRIETARY NAME: OsteoGraf/LD-300

CLASSIFICATION NAME: Bone Grafting Material 21 CFR 872.3930

PREDICATE DEVICES: OsteoGraf/LD-300, (K960353, K942212)

DEVICE DESCRIPTION: The OsteoGraf/LD-300 material is a high purity, high density, non-resorbable, radiopaque, polycrystalline particulate form of hydroxylapatite, the major mineral phase of bone and dental enamel.

INTENDED USE: Treatment of intrabony periodontal defects, augmentation of bony defects in the alveolar ridge, and filling of extraction sites.

TECHNOLOGICAL CHARACTERISTICS: All of the components found in OsteoGraf/LD-300 have been used in legally marketed devices and/or were found safe for dental use. The modifications made to the legally marketed device do not affect biocompatibility. Therefore, it was determined that biocompatibility testing was not necessary. OsteoGraf/LD-300 conforms to applicable industry standards.

We believe that the prior use of the components of OsteoGraf/LD-300 in legally marketed devices, the performance data provided, and biocompatibility support the safety and effectiveness of OsteoGraf/LD-300 for the indicated uses.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

AUG - 6 2007

Ms. Helen Lewis
Director of Corporate Compliance and Regulatory Affairs
DENTSPLY International, Incorporated
Susquehanna Commerce Center
221 West Philadelphia Street, Suite 60
York, Pennsylvania 17405-0872

Re: K072057
Trade/Device Name: OsteoGraf/LD-300
Regulation Number: 21 CFR 872.3930
Regulation Name: Bone Grafting Material
Regulatory Class: II
Product Code: LYC
Dated: July 18, 2007
Received: July 26, 2007

Dear Ms. Lewis:

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" (21 CFR 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

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K072057

INDICATIONS FOR USE STATEMENT

510(k) Number (if known): K072057

Device Name: OsteoGraf/LD-300

Indications for Use:

OsteoGraf/LD-300 is indicated for treatment of intrabony periodontal defects, augmentation of bony defects in the alveolar ridge, and filling of extraction sites.

These are the same indications for use previously cleared for K960353 and K942212.

Prescription Use __X__ (Part 21 CFR 801 Subpart D)	AND/OR	Over-The-Counter Use _____ (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 Anesthesiology, General Hospital
Infection Control, Dental Devices

510(k) Number: K072052

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