K072342

510(k) SUMMARY

OCT 3 2007

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

CONTACT: Helen Lewis DATE PREPARED: August 17, 2007

TRADE OR PROPRIETARY NAME: SULTAN TOOTH ROOT DESENSITIZER

CLASSIFICATION NAME: Cavity varnish 872.3260

PREDICATE DEVICES: PerioSRP™ Tooth Root Conditioner with NovaMin® K071267
Butler NuCare® Root Conditioner with NovaMin® K033295

DEVICE DESCRIPTION: SULTAN TOOTH ROOT DESENSITIZER is identical to the predicate devices. This product is manufactured by Novamin Technology Inc. and is then relabelled/repackaged. SULTAN TOOTH ROOT DESENSITIZER is a two-phase product designed to minimize sensitivity and promote clinical healing following periodontal debridement.

INTENDED USE: SULTAN TOOTH ROOT DESENSTIZER is intended for the rapid relief of hypersensitivity associated with exposed tooth root dentin.

TECHNOLOGICAL CHARACTERISTICS: All of the components found in SULTAN TOOTH ROOT DESENSITIZER have been used in the legally marketed predicate devices.

SULTAN TOOTH ROOT DESENSITIZER was tested for toxicity, irritation, and sensitization and found to be biocompatible.

SULTAN TOOTH ROOT DESENSITIZER is the same product as the legally marketed devices. We believe this fact supports the safety and effectiveness of SULTAN TOOTH ROOT DESENSITIZER.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Ms. Helen Lewis
Director of Corporate Compliance and Regulatory Affairs
DENTSPLY International, Incorporated
Susquehanna Commerce Center West
221 West Philadelphia Street, Suite 60
York, Pennsylvania 17404

OCT 3 2007

Re: K072342
Trade/Device Name: Sultan Tooth Root Desensitizer
Regulation Number: 21 CFR 872.3260
Regulation Name: Cavity Varnish
Regulatory Class: II
Product Code: LBH
Dated: August 17, 2007
Received: August 21, 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" (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

INDICATIONS FOR USE STATEMENT

510(k) Number (if known): K072342

Device Name: SULTAN TOOTH ROOT DESENSITIZER

Indications for Use:

SULTAN TOOTH ROOT DESENSITIZER is intended for the rapid relief of hypersensitivity associated with exposed tooth root dentin.

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)

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

510(k) Number: K072342