

BioHorizons Implant Systems, Inc.
510(k) Notification
January 28, 1998

12980874

MAR - 4 1998

510(k) Summary of Safety and Effectiveness

Proprietary Name

Implant Site Dilators

Common Name

Osteotomes

Classification Name

Dental Surgical Instruments

Classification

Class III (revised 24Feb98)

Official Contact

**Winston Greer, VP of Operations and Director of Quality Assurance
BioHorizons Implant Systems, Inc.
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Birmingham, Alabama 35209
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Device Description

The Implant Site Dilators are hand-operated instruments which provide an aid in the placement of root form implants in soft maxillary bone. All components are made from surgical grade stainless steel.

Product Evaluation

The BioHorizons Implant Site Dilators have been evaluated in laboratory conditions and by consulting implantologists and oral and maxillofacial surgeons. These evaluations indicate that the Implant Site Dilators should be safe and effective when used as intended.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Mr. Winston Greer
Vice President of Operations & Director of
Quality Assurance
BioHorizons Implant Systems, Incorporated
2129 Montgomery Highway
Birmingham, Alabama 35209

MAR - 4 1998

Re: K980374
Trade Name: Implant Site Dilators
Regulatory Class: III
Product Code: DZE
Dated: January 28, 1998
Received: January 30, 1998

Dear Mr. Greer:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to 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). 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 (Pre-market Approval), 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 895. A substantially equivalent determination assumes compliance with the current Good Manufacturing Practice requirement, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic (QS) inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531

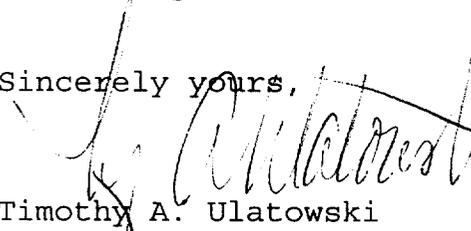
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through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 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 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4618. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

Sincerely yours,



Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

