K072891

Acclarent

Sinus Spacer

APPENDIX A: 510(k) SUMMARY

MAR 19 2008

Sponsor/Submitter:	Acclarent, Inc. 1525-B O'Brien Drive Menlo Park, California 94025
Contact Person:	Keri Yen Regulatory Affairs Specialist Phone: (650) 687-5874 Fax: (650) 687-4449
Date of Submission:	October 9, 2007
Device Trade Name:	TBD
Common Name:	Sinus Spacer
Device Classification:	Class I exempt
Regulation Number:	21 CFR 878.4800
Classification Name:	Manual surgical instrument for general use
Product Code:	KAM
Predicate Devices:	Ethmoid Sinus Spacer (K062458) Rains Frontal Sinus Stent (K951066) Micromedics, Inc. Sphenoid Sinus Stent (K050340) Shikani Middle Meatal Antrostomy Stent (K912418)
Device Description:	The Sinus Spacer is a device that maintains an opening at the intended sinus not to exceed 14 days postoperatively. The subject device can be manually removed during an office follow-up visit at any time.
Indications for Use:	The Sinus Spacer is indicated for use as a postoperative spacer to maintain an opening to the nasal sinuses within the first 14 days following surgery. The Sinus Spacer also helps to prevent obstructions.
Technological Characteristics:	The Sinus Spacer is designed to be implanted into all sinuses and to maintain its position by a self-retention mechanism.
Performance Data:	The Sinus Spacer met all performance acceptance criteria.
Summary of Substantial Equivalence:	The Sinus Spacer is substantially equivalent to the predicate devices as confirmed through relevant performance tests.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

MAR 19 2008

Acclarent, Inc.
c/o Keri Yen
Regulatory Affairs Specialist
1525-B O'Brien Drive
Menlo Park, CA 94025

Re: K072891
Trade/Device Name: Sinus Spacer
Regulation Number: 21 CFR 874.4800
Regulation Name: ENT Manual Surgical Instrument
Regulatory Class: Class I
Product Code: KAM
Dated: March 7, 2008
Received: March 10, 2008

Dear Ms. Yen:

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 Center for Devices and Radiological Health's (CDRH's) Office of Compliance at (240) 276-0115. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding postmarket surveillance, please contact CDRH's Office of Surveillance and Biometric's (OSB's) Division of Postmarket Surveillance at 240-276-3474. For questions regarding the reporting of device adverse events (Medical Device Reporting (MDR)), please contact the Division of Surveillance Systems at 240-276-3464. 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 (240) 276-3150 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,

Malvina B. Eydelman, M.D.

Malvina B. Eydelman, M.D.
Director
Division of Ophthalmic and Ear, Nose
and Throat Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

APPENDIX B: INDICATIONS FOR USE STATEMENT

510(k) Number (if known): K072891

Trade Name: To be determined

Common Name: Sinus Spacer

Indications For Use: The Sinus Spacer is indicated for use as a postoperative spacer to maintain an opening to the nasal sinuses within the first 14 days following surgery. The Sinus Spacer also helps to prevent obstructions.

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 OF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Karen Baker

(Division Sign-Off)
Division of Ophthalmic Ear, Nose and Throat Devises

510(k) Number K072891