

K041887



AUG 19 2004

3.0 510(k) Summary

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**Sponsor:** SYNTHES (USA)  
1690 Russell Road  
Paoli, PA 19301  
(610) 647-9700

**Device Name:** Synthes 1.0/1.2 mm Craniofacial Screws

**Classification:** 21 CFR 872.4880: Screw, Fixation, Intraosseous

**Predicate Device:** KLS Martin 1.0mm System Center-Drive Screw System  
Synthes Maxillofacial Titanium Micro Set

**Device Description:** The 1.0/1.2 mm Craniofacial Screws are either self-drilling or self-tapping, have a flat head profile with rounded edges with a cruciform recess, and are available in various lengths.

**Intended Use:** The Synthes 1.0/1.2 mm Craniofacial screws are intended for use as follows:

Specific Indications:

- Nasoethmoidal fractures
- Infraorbital area fractures
- Frontal sinus wall fractures
- Infant craniofacial surgery

General Indications:

- Maxillofacial surgery

**Substantial Equivalence:** Documentation is provided which demonstrates that the Synthes 1.0/1.2mm Craniofacial Screws are substantially equivalent to other legally marketed devices.

The term "substantial equivalence" as used in this 510(k) notification is limited to the definition of substantial equivalence found in the Federal Food, Drug and Cosmetic Act, as amended and as applied under 21CFR 807, Subpart E under which a device can be marketed without premarket approval or reclassification. A determination of substantial equivalency under this notification is not intended to have any bearing whatsoever on the resolution of patent infringement suits or any other patent matters. No statements related to, or in support of substantial equivalence herein shall be construed as an admission against interest under the US Patent Laws or their application by the courts.



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

AUG 19 2004

Ms. Lisa M. Boyle  
Regulatory Associate  
Synthes (USA)  
1690 Russell Road  
Paoli, Pennsylvania 19301

Re: K041887

Trade/Device Name: Synthes (USA) 1.0/1.2 mm Craniofacial Screws  
Regulation Number: 872.4880  
Regulation Name: Intraosseous Fixation Screw or Wire  
Regulatory Class: II  
Product Code: DZL  
Dated: July 8, 2004  
Received: July 13, 2004

Dear Ms. Boyle:

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 (301) 594-4613. 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/dsma/dsmamain.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

