K020505

APR 2 3 2002

3.0 Summary of Safety and Effectiveness Information

Sponsor:	Synthes (USA) 1690 Russell Road Paoli, PA 19301
Contact:	Matthew M. Hull, RAC Phone: (610) 647-9700 x7191
Device Name:	Translating Maxillary Distractor
Classification:	Class II, 21 CFR 872.4760 (bone plate) and 872.4880 (intraosseous fixation screw or wire)
Substantial Equivalence:	Documentation was provided which demonstrated the Synthes Translating Maxillary Distractor to be substantially equivalent to other legally marketed devices.
Device Description:	The Synthes (USA) Translating Maxillary Distractor is an intra-oral distraction device. It features a distractor body with two adjustable footplate components, each with contourable legs having screw holes that are fixed to the bone via 2.0 mm or 2.4 mm cortex screws.
Intended Use:	The Synthes Translating Maxillary Distractor is intended for use in craniofacial surgery, reconstructive procedures, and selective orthognathic surgery of the maxilla. Specifically it is intended for distraction of the maxilla utilizing a LeFort I osteotmy in adult and pediatric populations.
Material:	Stainless Steel

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

APR 2 3 2002

Mr. Matthew M. Hull
Senior Regulatory Associate
Synthes (USA)
1690 Russell Road
P.O. Box 1766
Paoli, Pennsylvania 19301

Re: K020505
Trade/Device Name: Synthes (USA) Translating Maxillary Distractor
Regulation Number: 872.4760 and 872.4880
Regulation Name: Bone Plate
Regulatory Class: II
Product Code: JEY
Dated: February 14, 2002
Received: February 15, 2002

Dear Mr. Hull:

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 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4613. 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" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained 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,

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

Enclosure

K020505

2.0 Indications for Use Statement

510(k) Number (if known): _____

Device Name: Synthes (USA) Translating Maxillary Distractor

Indications/Contraindications: The Synthes Translating Maxillary Distractor is intended for use in craniofacial surgery, reconstructive procedures, and selective orthognathic surgery of the maxilla. Specifically it is intended for distraction of the maxilla utilizing a LeFort I osteotomy in adult and pediatric populations.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use ✓ OR Over-The-Counter Use_
(Per 21 CFR 801.109)

(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(k) Number K020505