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K040588
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3.0 Summary of Safety and Effectiveness Information - 510(k) Summary

SPONSOR:	Synthes (USA) 1690 Russell Road Paoli, PA 19301 (610) 647-9700 Contact: Lisa M. Boyle
DEVICE NAME:	2.4 mm LCP Wrist Plate
CLASSIFICATION:	Class II, § 888.3030 – Single / multiple component metallic bone fixation appliance and accessories
PREDICATE DEVICE:	• Biomet® Colles Fracture Plate
DEVICE DESCRIPTION:	The 2.4 mm LCP Wrist Plate is a pre-contoured plate which is positioned dorsolaterally on the distal radius and affixed to the second metacarpal to span the wrist. A low profile construct is maintained by incorporating a 2.6 mm thickness, and a width of 7.5 mm. The plate has combination and DCU holes which accept 2.4 mm cortex and 2.4 mm locking screws. The plate is made out of Stainless Steel with tapered ends and is 170 mm in length.
INTENDED USE:	The Synthes (USA) 2.4mm LCP Wrist Plate is intended for Colles' and distal radius fractures with dorsal angulation of the distal fragment.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

MAY - 7 2004

Lisa M. Boyle
Regulatory Associate
Synthes USA
1690 Russel Road
P.O. Box 1766
Paoli, Pennsylvania 19301

Re: K040588
Trade/Device Name: Synthes (USA) 2.4 mm LCP Wrist Plate
Regulation Number: 21 CFR 888.3030
Regulation Name: Single/multiple component metallic bone fixation appliances and accessories
Regulatory Class: II
Product Code: HRS
Dated: March 04, 2004
Received: March 05, 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.

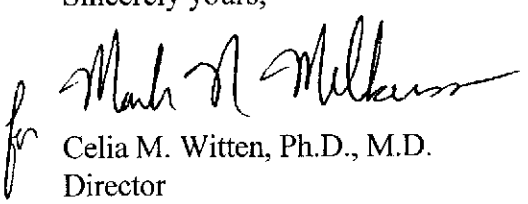
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish 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) 594-4659. 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,



Celia M. Witten, Ph.D., M.D.
Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

2.0 Indications for Use Statement

510(k) Number (if known): K040588

Device Name: Synthes (USA) 2.4 mm LCP Wrist Plate

Indications/Contraindications:

The Synthes (USA) 2.4mm LCP Wrist Plate is intended for colles' and distal radius fractures with dorsal angulation of the distal fragment.

for Mark A. Milhouse

(Division Sign-Off)
Division of General, Restorative, and Neurological Devices

510(k) Number K040588

Prescription Use X OR Over-The-Counter Use _____
(Per 21 CFR 801.109)

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

Concurrence of CDRH, Office of Device Evaluation (ODE)