



Food and Drug Administration
10903 New Hampshire Avenue
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Silver Spring, MD 20993-0002

Newclip Technics
% J.D. Webb
Official Correspondent
The OrthoMedix Group, Inc
1001 Oakwood Blvd
Round Rock, Texas 78681

October 2, 2017

Re: K172596
Trade/Device Name: Large Screws Range
Regulation Number: 21 CFR 888.3040
Regulation Name: Smooth Or Threaded Metallic Bone Fixation Fastener
Regulatory Class: Class II
Product Code: HWC
Dated: August 24, 2017
Received: September 5, 2017

Dear Mr. Webb:

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. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to 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); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); 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.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Industry and Consumer Education (DICE) at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to

<http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Industry and Consumer Education (DICE) at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely,

Mark N. Melkerson -S

Mark N. Melkerson
Director
Division of Orthopedic Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indications for Use

510(k) Number (if known)
K172596

Device Name

Large Screws range

Indications for Use (Describe)

The Large Screws range is indicated for the fixation of bone fractures, pseudarthroses, fusions and osteotomies of foot and ankle as well as medium and large bone fragments including radius, humerus, femur, tibia in adults.

Type of Use (Select one or both, as applicable)

Prescription Use (Part 21 CFR 801 Subpart D)

Over-The-Counter Use (21 CFR 801 Subpart C)

CONTINUE ON A SEPARATE PAGE IF NEEDED.

This section applies only to requirements of the Paperwork Reduction Act of 1995.

DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.

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4. 510 (k) Summary for the LARGE SCREWS RANGE

In accordance with 21 CFR 807.92 of the Federal Code of Regulations, the following 510(k) summary is submitted for the Large Screws Range.

Summary preparation date: August , 2017

1. Submitter:

NEWCLIP TECHNICS
P.A. de la Lande Saint Martin
45 rue des Garottières
F-44115 Haute-Goulaine - France
Telephone: (33) 2 28 21 37 12

Contact Person:

J.D. Webb
The OrthoMedix Group, Inc.
1001 Oakwood Blvd
Round Rock, TX 78681
Telephone: 512-388-0199

2. Trade Name:

Large Screws Range

Common Name:

Large Screws

Product Code:

HWC

Classification Name:

Smooth or threaded metallic bone fixation fastener. (21 CFR part. 888.3040)

3. Primary predicate or legally marketed devices which are substantially equivalent:

- Large Screws Range of Newclip Technics (K160617)

Secondary predicate or legally marketed devices which are substantially equivalent:

- Asnis III Cannulated Screw System of Howmedica Osteonics Corp. (K000080 and K024060)

Reference device:

- Activ Ankle Locking Plating System of Newclip Technics (K143061)



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4. Description of the device:

The Large Screws range consists of screws and washers designed for the fixation of bone fractures, pseudarthroses, fusions and osteotomies of foot and ankle as well as medium and large bone fragments including radius, humerus, femur, tibia in adults.

The implants of the Large Screws Range will be provided non sterile for sterilization by health care professionals prior to use or provided sterile by gamma sterilization.

The instruments of the Large Screws Range will be provided non sterile for sterilization by health care professionals prior to use.

Single use kits (Initial S) containing implants and/or instruments will be provided sterile by gamma sterilization.

Additional sterile instruments will also be provided sterile by gamma sterilization.

Materials:

Titanium alloy Ti-6Al-4V (conform to ASTM F136 and ISO 5832-3).

Function:

The implants of the Large Screws range are indicated for the fixation of bone fractures, pseudarthroses, fusions and osteotomies of foot and ankle as well as medium and large bone fragments including radius, humerus, femur, tibia in adults.

5. Substantial equivalence claimed to predicate devices:

The Large Screws Range is substantially equivalent to the predicate devices in terms of intended use, design, materials used, mechanical safety, and performances.

6. Indications for use:

The Large Screws range is indicated for the fixation of bone fractures, pseudarthroses, fusions and osteotomies of foot and ankle as well as medium and large bone fragments including radius, humerus, femur, tibia in adults.



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7. Non-clinical Test Summary:

Engineering analyses have been conducted in comparison with Large Screws Range of Newclip Technics (K160617).

LAL testing was performed to demonstrate that the subject device meets the endotoxin limit of <20 EU/device.

8. Clinical Test Summary:

No clinical studies were performed.

9. Conclusions Non-clinical and Clinical:

Newclip Technics considers the Large Screws Range to be equivalent to the predicate devices listed above. This conclusion is based upon the device's similarities in principles of operation, technology, materials, and indications for use.