



October 13, 2020

Skeletal Dynamics, Inc.
Diana Taylor
Sr. Regulatory Affairs Specialist
7300 N. Kendall Drive, Suite 400
Miami, Florida 33156

Re: K201962

Trade/Device Name: Skeletal Dynamics Ulnar Shortening Plating System
Regulation Number: 21 CFR 888.3030
Regulation Name: Single/Multiple Component Metallic Bone Fixation Appliances And Accessories
Regulatory Class: Class II
Product Code: HRS
Dated: July 10, 2020
Received: July 15, 2020

Dear Diana Taylor:

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. Although this letter refers to your product as a device, please be aware that some cleared products may instead be combination products. The 510(k) Premarket Notification Database located at <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpmn/pmn.cfm> identifies combination product submissions. 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) for devices or postmarketing safety reporting (21 CFR 4, Subpart B) for combination products (see <https://www.fda.gov/combination-products/guidance-regulatory-information/postmarketing-safety-reporting-combination-products>); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820) for devices or current good manufacturing practices (21 CFR 4, Subpart A) for combination products; and, if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

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 <https://www.fda.gov/medical-devices/medical-device-safety/medical-device-reporting-mdr-how-report-medical-device-problems>.

For comprehensive regulatory information about medical devices and radiation-emitting products, including information about labeling regulations, please see Device Advice (<https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance>) and CDRH Learn (<https://www.fda.gov/training-and-continuing-education/cdrh-learn>). Additionally, you may contact the Division of Industry and Consumer Education (DICE) to ask a question about a specific regulatory topic. See the DICE website (<https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance/contact-us-division-industry-and-consumer-education-dice>) for more information or contact DICE by email (DICE@fda.hhs.gov) or phone (1-800-638-2041 or 301-796-7100).

Sincerely,

Shumaya Ali, MPH
Assistant Director
DHT6C: Division of Restorative, Repair
and Trauma Devices
OHT6: Office of Orthopedic Devices
Office of Product Evaluation and Quality
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known)

K201962

Device Name

Skeletal Dynamics Ulnar Shortening Plating System

Indications for Use (Describe)

The Skeletal Dynamics Ulnar Shortening Plating System is indicated for fractures and osteotomies, in particular for the ulna.

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.

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510(k) SUMMARY
Ulnar Shortening Plating System

Submitter

Skeletal Dynamics, Inc.
7300 N. Kendall Drive, Suite 400
Miami, FL 33156
Phone: 305-596-7585
Facsimile: 305-596-7591
Contact Person: Diana Taylor
Date: September 10, 2020

Name and Classification:

Name: Skeletal Dynamics Ulnar Shortening Plating System
Common Name: Screw, Fixation, Bone
Classification: 21 CFR §888.3030
Regulatory Class: Class II
Product Code(s): HRS

Predicate Devices:

Primary Predicate Device: Medartis APTUS 2.5 Ulna Shortening Plate (K141232, 06/23/2014)
Additional Predicate Device: Medartis APTUS 2.5 TriLock Plate (K103332, 01/24/2011)

Device Description: The Skeletal Dynamics Ulnar Shortening Plating System consists of Distal Ulnar Shortening Plates with left and right configurations in two lengths and Proximal Ulnar Shortening Plates in two lengths. The Ulnar Shortening Plates are from medical grade Titanium Alloy (ASTM F136). Included within the Ulnar Shortening Plating System Set are 3.5mm cortical locking and non-locking bone screws as well as 2.3mm and 2.7mm locking and non-locking pegs from medical grade Titanium Alloy (ASTM F136) for fixation and class 1 surgical instrumentation.

Indications for Use: The Skeletal Dynamics Ulnar Shortening Plating System is indicated for fractures and osteotomies, in particular for the ulna.

Summary of Technological Characteristics: The substantial equivalence of the modifications to the Headless Compression Screw to the cleared predicate system is demonstrated by their identical indications for use, materials, design (fundamental scientific technology) and performance and does not present any new issues of safety or effectiveness.

Performance Testing: Static and dynamic bend strength was submitted with input from ASTM F382, Standard Specification and Test Method for Metallic Bone Plates ASTM F2193, Standard Specification and Test Methods for Components Used in the Surgical Fixation of the Spinal Skeletal System and ASTM F1541, Standard Specification and Test Methods for External Fixation Device. The performance test data as well as engineering analysis provided confirms the Skeletal Dynamics Ulnar Shortening Plating System performance to be substantially equivalent to the predicate devices.

Conclusion: As confirmed within this premarket notification, the Skeletal Dynamics Ulnar Shortening Plating System is equivalent to the legally marketed predicate devices and present no new issues with safety or effectiveness.