March 10, 2023

Arthrex Inc.
Kelsey Roberts
Sr. Regulatory Affairs Specialist
1370 Creekside Blvd
Naples, Florida 34108

Re: K230433
Trade/Device Name: Arthrex Double Loaded Knotless FiberTak® Suture Anchor
Regulation Number: 21 CFR 888.3040
Regulation Name: Smooth Or Threaded Metallic Bone Fixation Fastener
Regulatory Class: Class II
Product Code: MBI
Dated: February 17, 2023
Received: February 17, 2023

Dear Kelsey Roberts:

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,

Sara S. Thompson -S

For

Laurence D. Coyne, Ph.D.
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

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Indications for Use

Form Approved: OMB No. 0910-0120
Expiration Date: 06/30/2023
See PRA Statement below.

510(k) Number *(if known)*
K230433

Device Name

Arthrex Double Loaded Knotless FiberTak® Suture Anchor

Indications for Use *(Describe)*
The Arthrex Double Loaded Knotless FiberTak Suture Anchor is intended to be used for suture (soft tissue) fixation to bone in the shoulder, foot/ankle, knee, hand/wrist, elbow, and hip in the following procedures:

• Elbow: Biceps Tendon Reattachment, Ulnar or Radial Collateral Ligament Reconstruction

• Shoulder: Rotator Cuff Repair, Bankart Repair, SLAP Lesion Repair, Biceps Tenodesis, Acromio-Clavicular Separation Repair, Deltoid Repair, Capsular Shift or Capsulolabral Reconstruction

• Hand/Wrist: Scapholunate Ligament Reconstruction, Repair/Reconstruction of collateral ligaments, Repair of Flexor and Extensor Tendons at the PIP, DIP and MCP joints for all digits, digital tendon transfers, Carpal Ligament Reconstruction and Carpometacarpal joint arthroplasty (basal thumb joint arthroplasty)

• Foot/Ankle: Lateral Stabilization, Medial Stabilization, Achilles Tendon Repair, Metatarsal Ligament Repair, Hallux Valgus reconstruction, digital tendon transfers, Mid-foot reconstruction

• Knee: Medial Collateral Ligament Repair, Lateral Collateral Ligament Repair, Patellar Tendon Repair, Posterior Oblique Ligament Repair, Iliotibial Band Tenodesis, Joint Capsule Closure (Except Canada)

• Hip: Capsular Repair, Acetabular Labral Repair, Gluteal Tendon Repair (Except Canada)

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

Date Prepared	March 7, 2023
510(k) Number	K230433
Submitter	Arthrex Inc. 1370 Creekside Boulevard Naples, FL 34108-1945
Contact Person	Kelsey Roberts Sr. Regulatory Affairs Specialist (239) 598-4302 x 72257 Kelsey.Roberts@Arthrex.com
Trade Name	Arthrex Double Loaded Knotless FiberTak® Suture Anchor
Common Name	Smooth or threaded metallic bone fixation fastener
Product Code	MBI
Classification Name	21 CFR 888.3040: Smooth or threaded metallic bone fixation fastener
Regulatory Class	Class II
Predicate Device	K221396: Arthrex FiberTak Suture Anchor
Reference Devices	K203268: Arthrex FiberTak Suture Anchor K223284: SutureLoc™ Implant K193503: Arthrex SwiveLock Suture Anchor K193575: Arthrex SutureTape K122374: Arthrex Suture K041553: Arthrex Suture Grafting Kit K032245: Arthrex FiberTape™ Family K021434: Arthrex FiberWire™ USP Suture Family
Purpose of Submission	This Special 510(k) premarket notification is submitted as a line extension to the Arthrex FiberTak Suture Anchors cleared via K221396.
Device Description	The Arthrex Double Loaded Knotless FiberTak Suture Anchor is an all-suture knotless anchor intended to be used for fixation of suture (soft tissue) to bone in the shoulder, foot/ankle, knee, hand/wrist, elbow and hip. The Arthrex Double Loaded Knotless FiberTak Suture Anchor is a sterile, single use implant device constructed from a hollow braid of polyester (sheath) preloaded on an inserter with a double loaded suture component composed of UHMWPE and polyester.

Indications for Use	The Arthrex Double Loaded Knotless FiberTak Suture Anchor is intended to be used for suture (soft tissue) fixation to bone in the shoulder, foot/ankle, knee, hand/wrist, elbow, and hip in the following procedures: • Elbow: Biceps Tendon Reattachment, Ulnar or Radial Collateral Ligament Reconstruction • Shoulder: Rotator Cuff Repair, Bankart Repair, SLAP Lesion Repair, Biceps Tenodesis, Acromio-Clavicular Separation Repair, Deltoid Repair, Capsular Shift or Capsulolabral Reconstruction • Hand/Wrist: Scapholunate Ligament Reconstruction, Repair/Reconstruction of collateral ligaments, Repair of Flexor and Extensor Tendons at the PIP, DIP and MCP joints for all digits, digital tendon transfers, Carpal Ligament Reconstruction and Carpometacarpal joint arthroplasty (basal thumb joint arthroplasty) • Foot/Ankle: Lateral Stabilization, Medial Stabilization, Achilles Tendon Repair, Metatarsal Ligament Repair, Hallux Valgus reconstruction, digital tendon transfers, Mid-foot reconstruction • Knee: Medial Collateral Ligament Repair, Lateral Collateral Ligament Repair, Patellar Tendon Repair, Posterior Oblique Ligament Repair, Iliotibial Band Tenodesis, Joint Capsule Closure (Except Canada) • Hip: Capsular Repair, Acetabular Labral Repair, Gluteal Tendon Repair (Except Canada)
Performance Data	Cyclic pull-out testing was conducted on the proposed devices submitted in this Special 510(k). The test data demonstrates that the proposed devices perform statistically equivalent to the predicate device. Bacterial endotoxin per EP 2.6.14/USP <85> was conducted to demonstrate that the device meets pyrogen limit specifications.
Technological Comparison	The proposed devices are a line extension to the predicate device K221396. The proposed and predicate devices have the same basic design, indications for use, intended use, packaging, shelf-life, biocompatibility profile, manufacturing, and sterilization processes. In comparison to the predicate device, the proposed modifications include minor dimensional changes to the sheath and repair suture. Any differences between the proposed and predicate devices are considered minor and do not raise questions concerning safety or effectiveness.

Conclusion	The Arthrex Double Loaded Knotless FiberTak Suture Anchor devices are substantially equivalent to the predicate device in which the basic design features, indications for use, intended use, materials, manufacturing, and sterilization processes are identical. Any differences between the proposed and predicate devices are considered minor and do not raise questions concerning safety or effectiveness. Based on the intended use, technological characteristics, and the test data submitted, Arthrex Inc. has determined that the proposed devices are substantially equivalent to the currently marketed predicate device.