



August 21, 2024

SurGenTec, LLC
Berny Villejeune
Official Correspondent
911 Clint Moore Rd
Boca Raton, Florida 33487

Re: K242141

Trade/Device Name: TiLink-P SI Joint Fusion System
Regulation Number: 21 CFR 888.3040
Regulation Name: Smooth or threaded metallic bone fixation fastener
Regulatory Class: Class II
Product Code: OUR
Dated: July 19, 2024
Received: July 22, 2024

Dear Berny Villejeune:

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 (the 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 available 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.

Additional information about changes that may require a new premarket notification are provided in the FDA guidance documents entitled "Deciding When to Submit a 510(k) for a Change to an Existing Device"

(<https://www.fda.gov/media/99812/download>) and "Deciding When to Submit a 510(k) for a Software Change to an Existing Device" (<https://www.fda.gov/media/99785/download>).

Your device is also subject to, among other requirements, the Quality System (QS) regulation (21 CFR Part 820), which includes, but is not limited to, 21 CFR 820.30, Design controls; 21 CFR 820.90, Nonconforming product; and 21 CFR 820.100, Corrective and preventive action. Please note that regardless of whether a change requires premarket review, the QS regulation requires device manufacturers to review and approve changes to device design and production (21 CFR 820.30 and 21 CFR 820.70) and document changes and approvals in the device master record (21 CFR 820.181).

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 Part 803) for devices or postmarketing safety reporting (21 CFR Part 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 Part 4, Subpart A) for combination products; and, if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR Parts 1000-1050.

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 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,

STEPHANIE SMITH -S

For Ronald P. Jean, Ph.D.

Director

DHT6B: Division of Spinal Devices

OHT6: Office of Orthopedic Devices

Office of Product Evaluation and Quality

Center for Devices and Radiological Health

Enclosure

Indications for Use

Submission Number (if known)

K242141

Device Name

TiLink-P SI Joint Fusion System

Indications for Use (Describe)

The TiLink-P SI Joint Fusion System is intended for sacroiliac joint fusion for conditions including sacroiliac joint disruptions and degenerative sacroiliitis.

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

Submitter: SurGenTec, LLC
911 Clint Moore Rd
Boca Raton, FL 33487
Phone: 561-990-7882
Email: bryan@surgentec.com

Official Correspondent: Mr. Berny Villejeune
Engineering Manager
SurGenTec, LLC
911 Clint Moore Rd
Boca Raton, FL 33487
Phone: 561-409-1494
Email: berny@surgentec.com

Date Prepared: July 22, 2024

Trade Name: TiLink-P SI Joint Fusion System

Common Name: Smooth or threaded metallic bone fixation fastener

Classification: II
21 CFR 888.3040

Product Code: OUR

Predicate Device: TiLink-P Joint Fusion System (K240720)

Device Description:

The purpose of this Special 510(k) is to introduce the TiLink-P with Window variant to the TiLink-P SI Joint Fusion System. The Window feature is designed to promote osteointegration.

The TiLink-P SI Joint Fusion System contains various orthopedic instruments to assist the user in implanting a titanium sacroiliac implant into the sacroiliac joint to transfix the joint. The SurGenTec TiLink-P SI Joint Fixation System includes TiLink Implants and associated Instruments. The TiLink-P implants may be used standalone. The TiLink-P SI Joint Fusion System fixates the sacroiliac joint by transfixing the cortices of the ilium and cortices of the sacrum.

The TiLink-P Compression Anchor is positioned so that it spans the cortices of ilium and cortices of sacrum, the TiLink-P Locking Screw is inserted through the TiLink-P Compression Anchor and along the SI Joint via a posterior approach, in order to compress

the joint. Bone graft material is used with the TiLink-P SI Joint Fusion System to facilitate bone growth.

Indications for Use:

The TiLink-P SI Joint Fusion System is intended for sacroiliac joint fusion for conditions including sacroiliac joint disruptions and degenerative sacroiliitis.

Substantial Equivalence:

The subject TiLink-P with Window SI Joint Fusion System is substantially equivalent to the previously cleared version in K240720 with respect to intended use, materials, design, and function.

Performance Testing:

The following tests have been performed on the TiLink-P with Window SI Joint Fusion System:

- Axial Pullout per ASTM F543
- Static and Dynamic Vertical Shear per ASTM F3574.

Through these mechanical strength tests the TiLink-P SI Joint Fusion System with Window demonstrates substantial equivalence to the predicate device.