DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
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August 7, 2015

Precision Spine, Incorporated
% Mr. Kenneth C. Maxwell II
Regulatory and Quality Specialist
Empirical Testing, Corporation
4628 Northpark Drive
Colorado Springs, Colorado 80918

Re: K151863
Trade/Device Name: Reli SP Spinous Plating System
Regulation Number: 21 CFR 888.3050
Regulation Name: Spinal interlaminal fixation orthosis
Regulatory Class: Class II
Product Code: PEK
Dated: July 7, 2015
Received: July 8, 2015

Dear Mr. Maxwell:

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 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 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 yours,

Lori A. Wiggins -S

for

Mark N. Melkerson
Director
Division of Orthopedic Devices
Office of Device Evaluation
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: January 31, 2017
See PRA Statement on last page.

510(k) Number *(if known)*
K151863

Device Name
Reli SP Spinous Plating System

Indications for Use *(Describe)*

The Reli SP Spinous Plating System of Precision Spine, Inc. is a posterior, non-pedicle supplemental fixation device, intended for use in the non-cervical spine (T1-S1) of skeletally mature patients. It is intended for single level plate fixation/attachment to spinous process for the purpose of achieving supplemental fusion in the following conditions: degenerative disc disease (defined as back pain of discogenic origin with degeneration of the disc confirmed by history and radiographic studies), trauma (i.e. fracture or dislocation), spondylolisthesis, and/or tumor. It is not intended for stand-alone use.

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)

PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON A SEPARATE PAGE IF NEEDED.

FOR FDA USE ONLY

Concurrence of Center for Devices and Radiological Health (CDRH) *(Signature)*

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

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5. 510(K) SUMMARY

Submitter's Name:	Precision Spine
Submitter's Address:	2050 Executive Drive Pearl, MS 39208
Submitter's Telephone:	973-455-7150
Contact Person:	Kenneth C Maxwell II Empirical Testing Corp. 904.392.7576
Date Summary was Prepared:	06 Aug 2015
Trade or Proprietary Name:	Reli SP Spinous Plating System
Common or Usual Name:	Spinous Process Plate
Classification:	Class II per 21 CFR §888.3050 Spinal interlaminal fixation orthosis
Product Code:	PEK
Classification Panel:	87 Orthopedics

DESCRIPTION OF THE DEVICE SUBJECT TO PREMARKET NOTIFICATION:

The Precision Spine Reli SP Spinous Plating System consists of an ISP female plate and an ISP male plate for posterior fixation of the spine in order to achieve fusion. The ISP female plate and an ISP male plate are available in multiple sizes to accommodate various patient anatomies. The ISP female plate and an ISP male plate feature teeth to interface with the bone of the spinous processes. The ISP male plate is passed through the insert such that, in their final position, the ISP female plate and an ISP male plate surround the spinous processes on both sides, and fixation is achieved via compression of the two components onto the spinous processes. The purpose of this submission is to offer an expanded size range for the Reli SP Spinous Plating System.

INDICATIONS FOR USE

The Reli SP Spinous Plating System of Precision Spine, Inc. is a posterior, non-pedicle supplemental fixation device, intended for use in the non-cervical spine (T1-S1) of skeletally mature patients. It is intended for single level plate fixation/attachment to spinous process for the purpose of achieving supplemental fusion in the following conditions: degenerative disc disease (defined as back pain of discogenic origin with degeneration of the disc confirmed by history and radiographic studies), trauma (i.e. fracture or dislocation), spondylolisthesis, and/or tumor. It is not intended for stand-alone use.

TECHNICAL CHARACTERISTICS

The Precision Spine Reli SP Spinous Plating System is manufactured from medical grade Titanium (Ti 6Al-4V) per ASTM F136. The implants are provided non-sterile with instructions for sterilization. The interspinous plates are designed in total heights of 28-55mm.

Reli SP Spinous Plating System is made from material that conforms to ASTM F136. The subject and predicate devices have nearly identical technological characteristics and the minor differences do not raise any new issues of safety and effectiveness. Specifically the following characteristics are identical between the subject and predicates:

- Indications for Use
- Materials of manufacture
- Structural support mechanism

Table 5-1 Predicate Devices

510k Number	Trade or Proprietary or Model Name	Manufacturer	Type
K142378	Interspinous Plate System	Precision Spine	Primary
K130438	Axle Interspinous Fusion System	Spine-X	Additional

PERFORMANCE TESTING SUMMARY

Precision Spine has not conducted mechanical testing as part of this Special 510(k) Device Modification Premarket Notification. An analysis was performed to determine the subject implants were substantially equivalent to the predicate devices, therefore no mechanical testing was conducted as part of this submission.

CONCLUSION

The subject modified is the Reli SP Spinous Plating System and is very similar to previously cleared Reli SP Spinous Plating System. The subject Reli SP Spinous Plating System has the same intended uses, indications, technological characteristics, and principles of operation as the predicate devices. The modifications raise no new types of safety or effectiveness questions. The overall technology characteristics lead to the conclusion that the Reli SP Spinous Plating System is substantially equivalent to the predicate devices.