FDA 510(k) Premarket Notification

K160820 PressON Pro Spinal Fixation System

CLEARED

Jun 28, 2016
96 days to decision

K160820 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k160820/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Mar 24, 2016
Decision date	Jun 28, 2016
Days to decision	96 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nexus Spine, LLC**
Location	Salt Lake City, UT, US
Contact	Jared Crocker
Website	https://nexusspine.com
510(k) history	17 submissions · 17 cleared · 2014-2025