FDA 510(k) Premarket Notification

K173882 SeaSpine Mariner Pedicle Screw System

CLEARED

Feb 6, 2018
47 days to decision

K173882 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k173882/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Dec 21, 2017
Decision date	Feb 6, 2018
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**SeaSpine Orthopedics Corporation**
Location	Carlsbad, CA, US
Contact	Gina Flores
510(k) history	66 submissions · 66 cleared · 2016-2025