FDA 510(k) Premarket Notification

K173702 Juliet Ti LL

CLEARED

Jan 3, 2018
30 days to decision

K173702 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k173702/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 4, 2017
Decision date	Jan 3, 2018
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spineart**
Location	Geneva, CH
Contact	Franck Pennesi
510(k) history	44 submissions · 44 cleared · 2008-2023