FDA 510(k) Premarket Notification

K200312 Tryptik Ti

CLEARED

Apr 6, 2020
60 days to decision

K200312 · Product code: **ODP** · Orthopedic
Source: https://www.510kdatabase.net/k200312/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Feb 6, 2020
Decision date	Apr 6, 2020
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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