FDA 510(k) Premarket Notification

CLEARED

K142347 Amedica Valeo II Interbody Fusion Device

Nov 18, 2014
88 days to decision

K142347 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k142347/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Aug 22, 2014
Decision date	Nov 18, 2014
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Amedica Corp.**
Location	Salt Lake City, UT, US
Contact	William D Jordan
510(k) history	16 submissions · 16 cleared · 2002-2018