FDA 510(k) Premarket Notification

K131671 MECTALIF

CLEARED

Jul 5, 2013
28 days to decision

K131671 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k131671/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jun 7, 2013
Decision date	Jul 5, 2013
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medacta International**
Location	San Diego, CA, US
Contact	ADAM GROSS
510(k) history	39 submissions · 39 cleared · 2011-2016