FDA 510(k) Premarket Notification

K121332 LOOP CAGE

CLEARED

Jun 29, 2012
57 days to decision

K121332 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k121332/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	May 3, 2012
Decision date	Jun 29, 2012
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Advanced Medical Technologies AG**
Location	Round Rock, TX, US
Contact	J.D. WEBB
510(k) history	8 submissions · 8 cleared · 2004-2012