

K121332 LOOP CAGEJun 29, 2012
57 days to decisionK121332 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k121332/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 18, 2026