

K080083 INTREPID SPINAL SYSTEMApr 10, 2008
90 days to decisionK080083 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k080083/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jan 11, 2008
Decision date	Apr 10, 2008
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Sofamor Danek
Location	Memphis, TN, US
Contact	LEE GRANT
510(k) history	154 submissions · 147 cleared · 2002-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k080083/> 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 16, 2026