

K091813 INTREPID SPINAL SYSTEMNov 17, 2009
152 days to decisionK091813 · Product code: **OVD** · Orthopedic
Source: <https://www.510kdatabase.net/k091813/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Lumbar (OVD)
Date received	Jun 18, 2009
Decision date	Nov 17, 2009
Days to decision	152 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091813/> 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