FDA 510(k) Premarket Notification

K081673 DYN-ALIF ANTEROLATERAL INTERVERTEBRAL BODY FUSION DEVICE

CLEARED

Nov 14, 2008
154 days to decision

K081673 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k081673/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jun 13, 2008
Decision date	Nov 14, 2008
Days to decision	154 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ionics Medical Corp.**
Location	Ether, NC, US
Contact	M. LAINE MASHBUM
510(k) history	2 submissions · 2 cleared · 2000-2008

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k081673/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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