FDA 510(k) Premarket Notification

K073440 FOUNDATION CAGE

CLEARED

Apr 24, 2008
139 days to decision

K073440 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k073440/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 7, 2007
Decision date	Apr 24, 2008
Days to decision	139 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Corelink, LLC**
Location	Round Rock, TX, US
Contact	J.D. WEBB
510(k) history	35 submissions · 35 cleared · 2008-2023