FDA 510(k) Premarket Notification

K072289 SPINEFRONTIER DORADO VBR

CLEARED

Nov 1, 2007
77 days to decision

K072289 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k072289/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Aug 16, 2007
Decision date	Nov 1, 2007
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spinefrontier, Inc.**
Location	Beverly, MA, US
Contact	TOM CARLSON
510(k) history	24 submissions · 24 cleared · 2007-2020