

K142504 SpineFrontier Lumbar Interbody Fusion Device System

May 13, 2015
250 days to decisionK142504 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k142504/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Sep 5, 2014
Decision date	May 13, 2015
Days to decision	250 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k142504/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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