

K142026 SPINEFRONTIER ARENA-C HA PEEK CERVICAL INTERVERTEBRAL BODY FUSION DEVICEOct 20, 2014
87 days to decisionK142026 · Product code: **ODP** · Orthopedic
Source: <https://www.510kdatabase.net/k142026/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Jul 25, 2014
Decision date	Oct 20, 2014
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k142026/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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