K132740 SPINEFRONTIER LESPLASTY POSTERIOR CERVICAL LAMINOPLASTY SYSTEM

CLEARED

Nov 26, 2013
84 days to decision

K132740 · Product code: **NQW** · Orthopedic
Source: https://www.510kdatabase.net/k132740/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spine, Plate, Laminoplasty, Metal (NQW)
Date received	Sep 3, 2013
Decision date	Nov 26, 2013
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k132740/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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