CLEARED

K133575 SPINEVISION LUMIS CANNULATED PEDICLE SCREW FIXATION SYSTEM, SPINEVISION U.L.I.S. PEDICLE SCREW FIXATION SYSTEM

Jun 30, 2014
222 days to decision

K133575 · Product code: **MNH** · Orthopedic
Source: https://www.510kdatabase.net/k133575/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Nov 20, 2013
Decision date	Jun 30, 2014
Days to decision	222 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spinevision S.A.**
Location	Douglassville, PA, US
Contact	J.D. WEBB
510(k) history	5 submissions · 5 cleared · 2012-2016