FDA 510(k) Premarket Notification

K140479 COROENT XL-F SYSTEM

CLEARED

Jul 28, 2014
152 days to decision

K140479 · Product code: **OVD** · Orthopedic
Source: https://www.510kdatabase.net/k140479/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Lumbar (OVD)
Date received	Feb 26, 2014
Decision date	Jul 28, 2014
Days to decision	152 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nu Vasive, Incorporated**
Location	San Diego, CA, US
Contact	Cynthia Adams
510(k) history	112 submissions · 112 cleared · 2012-2023