FDA 510(k) Premarket Notification

K143595 NuVasive(R) CCX-L Interbody System

CLEARED

Apr 29, 2015
132 days to decision

K143595 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k143595/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 18, 2014
Decision date	Apr 29, 2015
Days to decision	132 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