FDA 510(k) Premarket Notification

K161230 NuVasive Lumbar Interbody Implants

CLEARED

Aug 25, 2016
115 days to decision

K161230 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k161230/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	May 2, 2016
Decision date	Aug 25, 2016
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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