FDA 510(k) Premarket Notification

CLEARED

K140770 MLX - MEDIAL LATERAL EXPANDABLE LUMBAR INTERBODY SYSTEM

Jul 25, 2014
120 days to decision

K140770 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k140770/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 27, 2014
Decision date	Jul 25, 2014
Days to decision	120 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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