FDA 510(k) Premarket Notification

K112606 ACTIVATOR V-E

CLEARED

Mar 6, 2012
180 days to decision

K112606 · Product code: **LXM** · Physical Medicine
Source: https://www.510kdatabase.net/k112606/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manipulator, Plunger-like Joint (LXM)
Date received	Sep 8, 2011
Decision date	Mar 6, 2012
Days to decision	180 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Activator Methods International, Ltd.**
Location	Phoenix, AZ, US
Contact	ARLAN W FUHR
510(k) history	3 submissions · 3 cleared · 2001-2012