FDA 510(k) Premarket Notification

K072519 ACTIVATOR V SPINAL ADJUSTING INSTRUMENT

CLEARED

Oct 23, 2007
46 days to decision

K072519 · Product code: **LXM** · Physical Medicine
Source: https://www.510kdatabase.net/k072519/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manipulator, Plunger-like Joint (LXM)
Date received	Sep 7, 2007
Decision date	Oct 23, 2007
Days to decision	46 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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