FDA 510(k) Premarket Notification

K091000 KNEEKG

CLEARED

Oct 26, 2009
201 days to decision

K091000 · Product code: **LXJ** · Physical Medicine
Source: https://www.510kdatabase.net/k091000/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Interactive Rehabilitation Exercise Devices (LXJ)
Date received	Apr 8, 2009
Decision date	Oct 26, 2009
Days to decision	201 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Emovi**
Location	Montreal, Quebec, CA
Contact	EMMANUEL MONTINI
510(k) history	1 submissions · 1 cleared · 2009-2009