FDA 510(k) Premarket Notification

K073099 FMRI HARDWARE SYSTEM

CLEARED

Nov 20, 2007
18 days to decision

K073099 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k073099/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Nov 2, 2007
Decision date	Nov 20, 2007
Days to decision	18 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Nordicneurolab**
Location	Minneapolis, MN, US
Contact	CONSTANCE G BUNDY
510(k) history	6 submissions · 6 cleared · 2007-2014