

K073099 FMRI HARDWARE SYSTEMNov 20, 2007
18 days to decisionK073099 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073099/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 16, 2026