

K893881 ACCESS TMJun 23, 1989
28 days to decisionK893881 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k893881/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	May 26, 1989
Decision date	Jun 23, 1989
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Diasonics, Inc.
Location	Mchenry, IL, US
Contact	BEN KHOSRAVI
510(k) history	42 submissions · 41 cleared · 1978-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893881/>, 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 20, 2026