

K923010 MAGNETIC RESONANCE DIAGNOSTIC DEVICEAug 25, 1992
64 days to decisionK923010 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k923010/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jun 22, 1992
Decision date	Aug 25, 1992
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Hitachi Medical Systems America, Inc.
Location	Twinsburg, OH, US
Contact	JOCHEN ROGERS
510(k) history	100 submissions · 100 cleared · 1991-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923010/> 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