

**K890092 HI STANDARD MAGNETIC RESONANCE DIAGNOSTIC
DEVICE**Jun 23, 1989
164 days to decisionK890092 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k890092/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jan 10, 1989
Decision date	Jun 23, 1989
Days to decision	164 days
Third-party review	No

APPLICANT

Company	Health Images, Inc.
Location	Atlanta, GA, US
Contact	RAYMOND F MCCLELLAN
510(k) history	5 submissions · 5 cleared · 1989-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890092/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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