FDA 510(k) Premarket Notification

K923010 MAGNETIC RESONANCE DIAGNOSTIC DEVICE

CLEARED

Aug 25, 1992
64 days to decision

K923010 · 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