FDA 510(k) Premarket Notification

K903319 MAGNETIC RESONANCE DIAGNOSTIC DEVICE

CLEARED

Aug 17, 1990
23 days to decision

K903319 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k903319/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jul 25, 1990
Decision date	Aug 17, 1990
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**GE Medical Systems**
Location	Milwaukee, WI, US
Contact	LARRY A KROGER,
510(k) history	169 submissions · 166 cleared · 1989-2022