FDA 510(k) Premarket Notification

K900889 MAGNETIC RESONANCE DIAGNOSTIC DEVICE ACCESSORIES

CLEARED

Mar 23, 1990
24 days to decision

K900889 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k900889/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Feb 27, 1990
Decision date	Mar 23, 1990
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**Siemens Medical Solutions USA, Inc.**
Location	Hoffman Estates, IL, US
Contact	SCOTT A HENSLEY
510(k) history	778 submissions · 778 cleared · 1980-2026