FDA 510(k) Premarket Notification

CLEARED

K030042 PREMIER III PHASED ARRAY CTL SPINE COIL

Feb 24, 2003
49 days to decision

K030042 · Product code: **MOS** · Radiology
Source: https://www.510kdatabase.net/k030042/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coil, Magnetic Resonance, Specialty (MOS)
Date received	Jan 6, 2003
Decision date	Feb 24, 2003
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Usa Instruments, Inc.**
Location	Aurora, OH, US
Contact	CHRISTIE SHUMAKER
510(k) history	64 submissions · 64 cleared · 1997-2005