FDA 510(k) Premarket Notification

K130706 DUOFLEX COIL SUITE (1.5T)

CLEARED

Jul 25, 2013
132 days to decision

K130706 · Product code: **MOS** · Radiology
Source: https://www.510kdatabase.net/k130706/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coil, Magnetic Resonance, Specialty (MOS)
Date received	Mar 15, 2013
Decision date	Jul 25, 2013
Days to decision	132 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Mr Instruments, Inc.**
Location	Minneapolis, MN, US
Contact	ROBERT BECK
510(k) history	7 submissions · 7 cleared · 2004-2017