

**K130706 DUOFLEX COIL SUITE (1.5T)**Jul 25, 2013  
132 days to decisionK130706 · 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	<b>Mr Instruments, Inc.</b>
Location	Minneapolis, MN, US
Contact	ROBERT BECK
510(k) history	7 submissions · 7 cleared · 2004-2017

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130706/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 16, 2026