

K131382 EXPRESSION MR200 MRI PATIENT MONITORING SYSTEM, EXPRESSION IP5 INFORMATION PORTALJul 15, 2013
62 days to decisionK131382 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k131382/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	May 14, 2013
Decision date	Jul 15, 2013
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Invivo Corporation
Location	Pewaukee, WI, US
Contact	RUSTY KELLY
510(k) history	29 submissions · 29 cleared · 2005-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k131382/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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