

K130290 MODIFICATION TO INTEL-GE CARE IINOVATIONS GUIDE

Jun 4, 2013
118 days to decision

K130290 · Product code: **DRG** · Cardiovascular
Source: <https://www.510kdatabase.net/k130290/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Feb 6, 2013
Decision date	Jun 4, 2013
Days to decision	118 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Intel-Ge Care Innovations, LLC
Location	Roseville, CA, US
Contact	MAUREEN GLYNN
510(k) history	3 submissions · 3 cleared · 2013-2013

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k130290/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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