

**K082185 GE LOGIQ E9 DIAGNOSTIC ULTRASOUND SYSTEM,
MODELS 5205000, 5205000-2**

Aug 15, 2008
14 days to decision

K082185 · Product code: IYO · Radiology
Source: <https://www.510kdatabase.net/k082185/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Aug 1, 2008
Decision date	Aug 15, 2008
Days to decision	14 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ge Medical Systems Ultrasound and Primary Care Dia
Location	Milwaukee, WI, US
Contact	NICOLE LANDREVILLE
510(k) history	7 submissions · 7 cleared · 2004-2011

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k082185/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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