

**K092140 GE VIVID I DIAGNOSTIC ULTRASOUND, GE VIVID Q  
DIAGNOSTIC ULTRASOUND**

Aug 21, 2009  
37 days to decision

K092140 · Product code: IYO · Radiology  
Source: <https://www.510kdatabase.net/k092140/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Jul 15, 2009
Decision date	Aug 21, 2009
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ge Medical Systems Israel, Ultrasound, Ltd.</b>
Location	Tirat Carmel, IL
Contact	JAMES TURNER
510(k) history	3 submissions · 3 cleared · 2008-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k092140/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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