

**K082374 VIVID I AND VIVID Q**Sep 2, 2008  
15 days to decisionK082374 · Product code: IYN · Radiology  
Source: <https://www.510kdatabase.net/k082374/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Aug 18, 2008
Decision date	Sep 2, 2008
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ge Medical Systems Israel, Ultrasound, Ltd.</b>
Location	Tirat Carmel, IL
Contact	ISRAEL CITRON
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/k082374/> 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 27, 2026