

**K070792 QLAB QUANTIFICATION**Apr 6, 2007  
15 days to decisionK070792 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k070792/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Mar 22, 2007
Decision date	Apr 6, 2007
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Ultrasound, Inc.</b>
Location	Santa Ana, CA, US
Contact	LYNN HARMER
510(k) history	46 submissions · 46 cleared · 1985-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070792/> 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 26, 2026