

**K063085 MODIFICATION TO ACUSON SEQUOIA ULTRASOUND SYSTEM**Nov 14, 2006  
35 days to decisionK063085 · Product code: IYO · Radiology  
Source: <https://www.510kdatabase.net/k063085/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Oct 10, 2006
Decision date	Nov 14, 2006
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	SHEILA W PICKERING
510(k) history	778 submissions · 778 cleared · 1980-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063085/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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