

**K900881 OR-340 INTRAOPERATIVE ULTRASOUND SYSTEM**Sep 25, 1990  
211 days to decisionK900881 · Product code: **IYO** · Radiology  
Source: <https://www.510kdatabase.net/k900881/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Feb 26, 1990
Decision date	Sep 25, 1990
Days to decision	211 days
Third-party review	No

**APPLICANT**

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Company	<b>Codman &amp; Shurtleff, Inc.</b>
Location	Mchenry, IL, US
Contact	JAMES R VEALE
510(k) history	152 submissions · 151 cleared · 1976-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900881/>, 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 June 15, 2026