

**K881028 7.5 MHZ ENDORECTAL PROBE**Apr 27, 1988  
48 days to decisionK881028 · Product code: **IYO** · Radiology  
Source: <https://www.510kdatabase.net/k881028/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Mar 10, 1988
Decision date	Apr 27, 1988
Days to decision	48 days
Third-party review	No

**APPLICANT**

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Company	<b>Philips Ultrasound, Inc.</b>
Location	Santa Ana, CA, US
Contact	MARTIN A KAUFMAN
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/k881028/>; 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