

K880839 AI 5200 DIAGNOSTIC ULTRASOUND SYSTEMJun 10, 1988
101 days to decisionK880839 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k880839/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Mar 1, 1988
Decision date	Jun 10, 1988
Days to decision	101 days
Third-party review	No

APPLICANT

Company	Phoenix Imaging
Location	Tempe, AZ, US
Contact	MATT HEDLUND
510(k) history	2 submissions · 2 cleared · 1988-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880839/>; Data sourced from FDA 510(k) public records (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. - Generated May 21, 2026