

K873937 SDR 3500 (PHILIPS PLATINUM +)Mar 23, 1988
177 days to decisionK873937 · Product code: IYO · Radiology
Source: <https://www.510kdatabase.net/k873937/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Sep 28, 1987
Decision date	Mar 23, 1988
Days to decision	177 days
Third-party review	No

APPLICANT

Company	Philips Ultrasound, Inc.
Location	Santa Ana, CA, US
Contact	MARTIN A KAUFMAN
510(k) history	46 submissions · 46 cleared · 1985-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k873937/>; 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 26, 2026