FDA 510(k) Premarket Notification

K873937 SDR 3500 (PHILIPS PLATINUM +)

CLEARED

Mar 23, 1988
177 days to decision

K873937 · 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