FDA 510(k) Premarket Notification

K881028 7.5 MHZ ENDORECTAL PROBE

CLEARED

Apr 27, 1988
48 days to decision

K881028 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k881028/

SUBMISSION DETAILS

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

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