FDA 510(k) Premarket Notification

K885228 RT6800 ECHOCARDIOGRAPHY SYSTEM

CLEARED

Mar 14, 1989
84 days to decision

K885228 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k885228/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Dec 20, 1988
Decision date	Mar 14, 1989
Days to decision	84 days
Third-party review	No

APPLICANT

Company	**General Electric Co.**
Location	Mchenry, IL, US
Contact	LARRY A KROGER
510(k) history	254 submissions · 254 cleared · 1976-2011