

K885228 RT6800 ECHOCARDIOGRAPHY SYSTEMMar 14, 1989
84 days to decisionK885228 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k885228/>; 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 17, 2026