FDA 510(k) Premarket Notification

K871555 RT5000 W/QUANTI. SPEC. DOPPLER, CW DOPP. MODIFICA.

CLEARED

Mar 2, 1988
315 days to decision

K871555 · Product code: **DXK** · Cardiovascular
Source: https://www.510kdatabase.net/k871555/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Apr 22, 1987
Decision date	Mar 2, 1988
Days to decision	315 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k871555/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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