FDA 510(k) Premarket Notification

K851807 RT2600(115V)(220/240V) #H4008A & H4008AE

CLEARED

Sep 19, 1985
142 days to decision

K851807 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k851807/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Apr 30, 1985
Decision date	Sep 19, 1985
Days to decision	142 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k851807/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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