FDA 510(k) Premarket Notification

K842880 MECHANICAL SECTOR PROBE 4 MHZ DFM

CLEARED

Dec 10, 1984
140 days to decision

K842880 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k842880/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Jul 23, 1984
Decision date	Dec 10, 1984
Days to decision	140 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k842880/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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