FDA 510(k) Premarket Notification

K802708 SYSTEM, ECHO, PULSED ULTRASONIC DIAG.

CLEARED

Jan 7, 1981
68 days to decision

K802708 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k802708/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Oct 31, 1980
Decision date	Jan 7, 1981
Days to decision	68 days
Third-party review	No

APPLICANT

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