

**K802708 SYSTEM, ECHO, PULSED ULTRASONIC DIAG.**Jan 7, 1981  
68 days to decisionK802708 · Product code: **IYO** · Radiology  
Source: <https://www.510kdatabase.net/k802708/>**SUBMISSION DETAILS**

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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**

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

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Device record: <https://www.510kdatabase.net/k802708/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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