

K802341 DIOSCANDec 10, 1980
75 days to decisionK802341 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k802341/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Sep 26, 1980
Decision date	Dec 10, 1980
Days to decision	75 days
Third-party review	No

APPLICANT

Company	Sonometrics Systems, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1978-1982

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Device record: <https://www.510kdatabase.net/k802341/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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