FDA 510(k) Premarket Notification

K802341 DIOSCAN

CLEARED

Dec 10, 1980
75 days to decision

K802341 · 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