FDA 510(k) Premarket Notification

K802530 PFIZONIC 250 ULTRASOUND SCANNER

CLEARED

Dec 18, 1980
63 days to decision

K802530 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k802530/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Oct 16, 1980
Decision date	Dec 18, 1980
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Pfizer Medical Systems, Inc.**
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1977-1981