FDA 510(k) Premarket Notification

K802531 PFIZONIC 200 ULTRASOUND SCANNER

CLEARED

Dec 18, 1980
63 days to decision

K802531 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k802531/

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k802531/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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