

**K802530 PFIZONIC 250 ULTRASOUND SCANNER**Dec 18, 1980  
63 days to decisionK802530 · Product code: **IYO** · RadiologySource: <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        | <b>Pfizer Medical Systems, Inc.</b>     |
| Location       | Mchenry, IL, US                         |
| 510(k) history | 12 submissions · 12 cleared · 1977-1981 |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k802530/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 22, 2026