

**K790089 TAPE MEASURE**Mar 6, 1979  
48 days to decisionK790089 · Product code: **FTY** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k790089/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)         |
| Submission type       | Traditional                                |
| Device classification | Tape, Measuring, Rulers And Calipers (FTY) |
| Date received         | Jan 17, 1979                               |
| Decision date         | Mar 6, 1979                                |
| Days to decision      | 48 days                                    |
| Third-party review    | No   |

**APPLICANT**

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|----------------|---|
| Company        | <b>Pfizer, Inc.</b>                     |
| Location       | Mchenry, IL, US                         |
| 510(k) history | 30 submissions · 30 cleared · 1977-2018 |

Pfizer, Inc. is an American multinational pharmaceutical and biotechnology corporation headquartered in Manhattan, New York City. Founded in 1849, Pfizer is one of the oldest pharmaceutical companies in North America. Pfizer's FDA 510(k) regulatory record includes cleared devices from total submissions, spanning 1977 to 2018. The company's device portfolio demonstrates strength in orthopedic devices, including surgical implants and fixation systems. This regulatory activity is now historical, with no clearances recorded in the past five years. The company's cleared device...