

K881378 PFIZER HOSPITAL PRODUCTS GROUP ACL DRILL GUIDEJun 17, 1988
77 days to decisionK881378 · Product code: LXI · Orthopedic
Source: <https://www.510kdatabase.net/k881378/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Drill, Ligament (LXI)
Date received	Apr 1, 1988
Decision date	Jun 17, 1988
Days to decision	77 days
Third-party review	No

APPLICANT

Company	Pfizer, Inc.
Location	Mchenry, IL, US
Contact	JOSEPH E LAPTEWICZ
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...

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881378/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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