

**K851971 SERIES 7000/HOLTER RECORDER**Jun 20, 1985  
44 days to decisionK851971 · Product code: **DSH** · CardiovascularSource: <https://www.510kdatabase.net/k851971/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Magnetic Tape, Medical (DSH)
Date received	May 7, 1985
Decision date	Jun 20, 1985
Days to decision	44 days
Third-party review	No

**APPLICANT**

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Company	<b>Pfizer, Inc.</b>
Location	Mchenry, IL, US
Contact	THOMAS MASSOPUST
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...

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