

**K851024 PLEUR-EVAC AUTOTRANSFUSION SYSTEM  
-TENTATIVE**May 23, 1985  
72 days to decisionK851024 · Product code: **CAC** · Anesthesiology  
Source: <https://www.510kdatabase.net/k851024/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Autotransfusion (CAC)
Date received	Mar 12, 1985
Decision date	May 23, 1985
Days to decision	72 days
Third-party review	No

**APPLICANT**

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Company	<b>Pfizer, Inc.</b>
Location	Mchenry, IL, US
Contact	SHELDON STEINBERG
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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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k851024/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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