

**K770425 NEEDLES, STERILE, AIRWAY**Apr 15, 1977  
39 days to decisionK770425 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k770425/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 7, 1977
Decision date	Apr 15, 1977
Days to decision	39 days
Third-party review	No

**APPLICANT**

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Company	<b>Pfizer Pharmaceuticals</b>
Location	Mchenry, IL, US
510(k) history	6 submissions · 5 cleared · 1977-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k770425/> 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 July 1, 2026