

**K895480 BLUNT NEEDLE PORT/I.V. PORT**Dec 1, 1989  
81 days to decisionK895480 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k895480/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 11, 1989
Decision date	Dec 1, 1989
Days to decision	81 days
Third-party review	No

**APPLICANT**

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Company	<b>Applied Medical Technologies</b>
Location	Mchenry, IL, US
Contact	DEAN SECREST
510(k) history	23 submissions · 20 cleared · 1981-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895480/>; 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