

**K981386 MANAN BLUNT NEEDLE**Jun 29, 1998  
74 days to decisionK981386 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k981386/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 16, 1998
Decision date	Jun 29, 1998
Days to decision	74 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Manan Medical Products, Inc.</b>
Location	Skokie, IL, US
Contact	MICHAEL PLISHKA
510(k) history	39 submissions · 38 cleared · 1988-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k981386/> 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 May 20, 2026