

**K990288 FREE FLOW SAFETY DEVICE, MODEL MG 245052**Feb 24, 1999  
26 days to decisionK990288 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k990288/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 29, 1999
Decision date	Feb 24, 1999
Days to decision	26 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Teva Medical, Ltd.</b>
Location	Kulpsville, PA, US
Contact	MARYANN MELUS
510(k) history	2 submissions · 2 cleared · 1997-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990288/> 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 4, 2026