

**K000369 VACUFLOW+ BLOOD COLLECTION SET, MODEL  
52221/3/5**Mar 30, 2000  
55 days to decisionK000369 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k000369/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Feb 4, 2000
Decision date	Mar 30, 2000
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary
Other names	VACUFLOW+ WITH HOLDER BLOOD COLLECTION SET, MODEL 82021/3/5 WITH HOLDER

**APPLICANT**

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Company	<b>Medpro Technologies, Inc.</b>
Location	Appollo Beach, FL, US
Contact	ART WARD
510(k) history	1 submissions · 1 cleared · 2000-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000369/> 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 June 29, 2026