

**K894335 MODEL #535 PACU STRETCHER/BED & #530  
ED/TRAUMA STR**Sep 12, 1989  
60 days to decisionK894335 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k894335/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jul 14, 1989
Decision date	Sep 12, 1989
Days to decision	60 days
Third-party review	No

**APPLICANT**

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Company	<b>Midmark Corp.</b>
Location	Versailles, OH, US
Contact	JOHN OLDIGES
510(k) history	31 submissions · 31 cleared · 1981-2013

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894335/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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