

**K801839 DISPOSABLE HYPODERMIC NEEDLES**Sep 16, 1980  
46 days to decisionK801839 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k801839/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Aug 1, 1980
Decision date	Sep 16, 1980
Days to decision	46 days
Third-party review	No

**APPLICANT**

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Company	<b>Martin&amp;apos;S Maine Exchange</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1981

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k801839/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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