

**K802400 MS 16 SYRINGE DRIVER**Dec 16, 1980  
75 days to decisionK802400 · Product code: **FRN** · General Hospital  
Source: <https://www.510kdatabase.net/k802400/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Oct 2, 1980
Decision date	Dec 16, 1980
Days to decision	75 days
Third-party review	No

**APPLICANT**

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Company	<b>Princeton Medical Instruments, Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1980

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Device record: <https://www.510kdatabase.net/k802400/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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