

K812888 SYRINGE, PISTONNov 10, 1981
26 days to decisionK812888 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k812888/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 15, 1981
Decision date	Nov 10, 1981
Days to decision	26 days
Third-party review	No

APPLICANT

Company	Frisco Products, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k812888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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