

K833399 SYRINGEJan 10, 1984
97 days to decisionK833399 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k833399/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 5, 1983
Decision date	Jan 10, 1984
Days to decision	97 days
Third-party review	No

APPLICANT

Company	I M, Inc.
Location	Mchenry, IL, US
510(k) history	41 submissions · 40 cleared · 1976-1992

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Device record: <https://www.510kdatabase.net/k833399/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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