

**K833435 MULTI PURPOSE SYRINGE**Dec 12, 1983  
68 days to decisionK833435 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k833435/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 5, 1983
Decision date	Dec 12, 1983
Days to decision	68 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k833435/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026