

**K854877 ACU-SYRINGE W/WO NEEDLES**Feb 19, 1986  
75 days to decisionK854877 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k854877/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Dec 6, 1985
Decision date	Feb 19, 1986
Days to decision	75 days
Third-party review	No

**APPLICANT**

---

Company	<b>Acuderm, Inc.</b>
Location	Mchenry, IL, US
Contact	CHARLES YEH
510(k) history	13 submissions · 13 cleared · 1983-1994

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k854877/>, 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 May 14, 2026