

**K895890 AUTOPEN**Feb 16, 1990  
134 days to decisionK895890 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k895890/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 5, 1989
Decision date	Feb 16, 1990
Days to decision	134 days
Third-party review	No

**APPLICANT**

---

Company	<b>Ulster Scientific, Inc.</b>
Location	Mchenry, IL, US
Contact	F CUNNINGHAM
510(k) history	20 submissions · 20 cleared · 1980-1994

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895890/>, 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 7, 2026