

**K880963 HAC AUTOLOGY CENTERS DISPENSING KIT FOR FIB.CON.**Jun 24, 1988  
108 days to decisionK880963 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k880963/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Mar 8, 1988
Decision date	Jun 24, 1988
Days to decision	108 days
Third-party review	No

**APPLICANT**

---

Company	<b>Buckman Co., Inc.</b>
Location	Mchenry, IL, US
Contact	DAVID W SCHLERF
510(k) history	111 submissions · 104 cleared · 1983-1998

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880963/> 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 21, 2026