

**K011176 REUSABLE GLASS SYRINGES**Aug 10, 2001  
115 days to decisionK011176 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k011176/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Apr 17, 2001
Decision date	Aug 10, 2001
Days to decision	115 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Lockett Medical Corp.</b>
Location	Providence, RI, US
Contact	WILLIAM LOCKETT III
510(k) history	3 submissions · 3 cleared · 1995-2001

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k011176/> 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 July 4, 2026