FDA 510(k) Premarket Notification

K011176 REUSABLE GLASS SYRINGES

CLEARED

Aug 10, 2001
115 days to decision

K011176 · 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	**Lockett Medical Corp.**
Location	Providence, RI, US
Contact	WILLIAM LOCKETT III
510(k) history	3 submissions · 3 cleared · 1995-2001