FDA 510(k) Premarket Notification

K983430 LITE-TOUCH SYRINGE

CLEARED

Feb 9, 1999
133 days to decision

K983430 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k983430/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Sep 29, 1998
Decision date	Feb 9, 1999
Days to decision	133 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medicore, Inc.**
Location	Walker, MI, US
Contact	BONNIE J KAPLAN
510(k) history	3 submissions · 3 cleared · 1981-1999