FDA 510(k) Premarket Notification

K101708 PATIENT SAFE LUER CAP

CLEARED

Feb 24, 2011
252 days to decision

K101708 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k101708/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jun 17, 2010
Decision date	Feb 24, 2011
Days to decision	252 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Retractable Technologies, Inc.**
Location	Lewisville, TX, US
Contact	RHONDA WELLS
510(k) history	13 submissions · 13 cleared · 1995-2021