FDA 510(k) Premarket Notification

K011951 EZ SYRINGE

CLEARED

Aug 2, 2001
42 days to decision

K011951 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k011951/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jun 21, 2001
Decision date	Aug 2, 2001
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Owen Mumford, Ltd.**
Location	Marietta, GA, US
Contact	ROBERT SHAW
Website	http://www.owenmumford.com/us/
510(k) history	13 submissions · 13 cleared · 2000-2023

Owen Mumford, Ltd. is a global medical device manufacturer with over 70 years of experience designing and manufacturing innovative healthcare solutions. The company specializes in drug delivery systems, blood collection devices, and safety lancets for both clinical and home use. Owen Mumford operates with a manufacturing facility in Marietta, US, and serves healthcare professionals and patients worldwide. The company has received FDA 510(k) clearances from total submissions, spanning from 2000 to 2023. Owen Mumford's cleared devices focus primarily on General Hospital app...