FDA 510(k) Premarket Notification

K202432 MiniLoad Syringe

CLEARED

Feb 19, 2021
178 days to decision

K202432 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k202432/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 25, 2020
Decision date	Feb 19, 2021
Days to decision	178 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Ocuject, LLC**
Location	Newport Beach, CA, US
Contact	Rebecca K Pine
510(k) history	8 submissions · 8 cleared · 2017-2024