

K212544 MiniLoad SyringeOct 13, 2021
62 days to decisionK212544 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k212544/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Syringe, Piston (FMF)
Date received	Aug 12, 2021
Decision date	Oct 13, 2021
Days to decision	62 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k212544/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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