

K202432 MiniLoad SyringeFeb 19, 2021
178 days to decisionK202432 · 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 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k202432/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026