FDA 510(k) Premarket Notification

# K882025 MODIFIED EXTRAVAKIT

**CLEARED**

Jul 27, 1988
71 days to decision

K882025 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k882025/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent - SD |
| Submission type | Traditional |
| Device classification | Syringe, Piston (FMF) |
| Date received | May 17, 1988 |
| Decision date | Jul 27, 1988 |
| Days to decision | 71 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Extravakit, Inc.** |
| Location | Tucson, AZ, US |
| Contact | DORR, PHD |
| 510(k) history | 1 submissions · 0 cleared · 1988-1988 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k882025/ 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 July 8, 2026