FDA 510(k) Premarket Notification

# K772129 MEDI-JECTOR

**CLEARED**

Dec 8, 1977
27 days to decision

K772129 · Product code: **KZE** · General Hospital
Source: https://www.510kdatabase.net/k772129/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Injector, Fluid, Non-electrically Powered (KZE) |
| Date received | Nov 11, 1977 |
| Decision date | Dec 8, 1977 |
| Days to decision | 27 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Derata Corp.** |
| Location | Mchenry, IL, US |
| 510(k) history | 6 submissions · 6 cleared · 1977-1988 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k772129/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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