FDA 510(k) Premarket Notification

# K823767 ERG-JET ELECTRODE

**CLEARED**

Jan 8, 1983
25 days to decision

K823767 · Product code: **HLF** · Ophthalmic
Source: https://www.510kdatabase.net/k823767/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Device, Measuring, Lens Radius, Ophthalmic (HLF) |
| Date received | Dec 14, 1982 |
| Decision date | Jan 8, 1983 |
| Days to decision | 25 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Life-Tech Intl., Inc.** |
| Location | Walker, MI, US |
| 510(k) history | 68 submissions · 66 cleared · 1982-2000 |