

K823767 ERG-JET ELECTRODEJan 8, 1983
25 days to decisionK823767 · Product code: **HLF** · Ophthalmic
Source: <https://www.510kdatabase.net/k823767/>**SUBMISSION DETAILS**

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k823767/> 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 21, 2026