

K822923 5012 PARABOLIC STIMULATORDec 28, 1982
88 days to decisionK822923 · Product code: **HLX** · Ophthalmic
Source: <https://www.510kdatabase.net/k822923/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Photostimulator, Ac-powered (HLX)
Date received	Oct 1, 1982
Decision date	Dec 28, 1982
Days to decision	88 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k822923/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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