

K822917 MODEL 5400 OPTOKINETIC STIMULATORNov 1, 1982
31 days to decisionK822917 · Product code: **HOW** · Ophthalmic
Source: <https://www.510kdatabase.net/k822917/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drum, Opticokinetic (HOW)
Date received	Oct 1, 1982
Decision date	Nov 1, 1982
Days to decision	31 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/k822917/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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