

**K822919 5010 VISUSTIM**Dec 28, 1982  
88 days to decisionK822919 · Product code: **HLX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k822919/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k822919/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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