FDA 510(k) Premarket Notification

K822923 5012 PARABOLIC STIMULATOR

CLEARED

Dec 28, 1982
88 days to decision

K822923 · 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