FDA 510(k) Premarket Notification

K822917 MODEL 5400 OPTOKINETIC STIMULATOR

CLEARED

Nov 1, 1982
31 days to decision

K822917 · 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