FDA 510(k) Premarket Notification

K831829 KARICKHOFF FLYING CORPUSCLE VIEWER

CLEARED

Oct 19, 1983
134 days to decision

K831829 · Product code: **HIX** · Ophthalmic
Source: https://www.510kdatabase.net/k831829/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Maxwell Spot, Ac-powered (HIX)
Date received	Jun 7, 1983
Decision date	Oct 19, 1983
Days to decision	134 days
Third-party review	No

APPLICANT

Company	**Surgidev Corp.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1983-1985