FDA 510(k) Premarket Notification

K830008 KARICKHOFF KERATOSCOPE

CLEARED

Jan 26, 1983
23 days to decision

K830008 · Product code: **HLR** · Ophthalmic
Source: https://www.510kdatabase.net/k830008/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratoscope, Battery-powered (HLR)
Date received	Jan 3, 1983
Decision date	Jan 26, 1983
Days to decision	23 days
Third-party review	No

APPLICANT

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