

K830008 KARICKHOFF KERATOSCOPEJan 26, 1983
23 days to decisionK830008 · 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

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Device record: <https://www.510kdatabase.net/k830008/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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