

K842892 NIDEK DIGITAL KERATOMETEROct 5, 1984
74 days to decisionK842892 · Product code: **HQL** · Ophthalmic
Source: <https://www.510kdatabase.net/k842892/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intraocular Lens (HQL)
Date received	Jul 23, 1984
Decision date	Oct 5, 1984
Days to decision	74 days
Third-party review	No

APPLICANT

Company	Nidek, Inc.
Location	Mchenry, IL, US
510(k) history	77 submissions · 77 cleared · 1983-2005

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Device record: <https://www.510kdatabase.net/k842892/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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