

K834305 ARGON/KRYPTON ION LASER PHOTOCOAGULAMar 16, 1984
109 days to decisionK834305 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k834305/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Nov 28, 1983
Decision date	Mar 16, 1984
Days to decision	109 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k834305/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 19, 2026