

K924260 NIDEK CV-12000Mar 26, 1993
214 days to decisionK924260 · Product code: **HQC** · Ophthalmic
Source: <https://www.510kdatabase.net/k924260/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Aug 24, 1992
Decision date	Mar 26, 1993
Days to decision	214 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k924260/> 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