

K936302 NIDEK MODEL VT-4000, MODIFICATIONJul 15, 1994
198 days to decisionK936302 · Product code: **HQE** · Ophthalmic
Source: <https://www.510kdatabase.net/k936302/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Vitreous Aspiration And Cutting, Ac-powered (HQE)
Date received	Dec 29, 1993
Decision date	Jul 15, 1994
Days to decision	198 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/k936302/> 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