FDA 510(k) Premarket Notification

K924260 NIDEK CV-12000

CLEARED

Mar 26, 1993
214 days to decision

K924260 · 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