FDA 510(k) Premarket Notification

K834305 ARGON/KRYPTON ION LASER PHOTOCOAGULA

CLEARED

Mar 16, 1984
109 days to decision

K834305 · 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