FDA 510(k) Premarket Notification

K832912 GELENDER VACUUM FIXATION RING SYS

CLEARED

Jan 10, 1984
134 days to decision

K832912 · Product code: **HNH** · Ophthalmic
Source: https://www.510kdatabase.net/k832912/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ring, Ophthalmic (flieringa) (HNH)
Date received	Aug 29, 1983
Decision date	Jan 10, 1984
Days to decision	134 days
Third-party review	No

APPLICANT

Company	**Miami Eye Technology, Inc.**
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1983-1984