

K832912 GELENDER VACUUM FIXATION RING SYSJan 10, 1984
134 days to decisionK832912 · Product code: **HNH** · Ophthalmic
Source: <https://www.510kdatabase.net/k832912/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ring, Ophthalmic (fleringa) (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

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Device record: <https://www.510kdatabase.net/k832912/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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