

**K922996 MODIFIED SITE DIAPHRAGM CASSETTE FOR
OPHTHALMICS**Jul 27, 1992
35 days to decisionK922996 · Product code: **KYG** · Ophthalmic
Source: <https://www.510kdatabase.net/k922996/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Irrigation, Ocular Surgery (KYG)
Date received	Jun 22, 1992
Decision date	Jul 27, 1992
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Chiron Vision Corp.
Location	Irvine, CA, US
Contact	BRUNT MILLER
510(k) history	34 submissions · 34 cleared · 1980-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k922996/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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