

K821897 LENSJan 5, 1983
191 days to decisionK821897 · Product code: **KYG** · Ophthalmic
Source: <https://www.510kdatabase.net/k821897/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Irrigation, Ocular Surgery (KYG)
Date received	Jun 28, 1982
Decision date	Jan 5, 1983
Days to decision	191 days
Third-party review	No

APPLICANT

Company	Mortan, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1983-1987

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Device record: <https://www.510kdatabase.net/k821897/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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