

K802332 INTRAJECTDec 22, 1980
90 days to decisionK802332 · Product code: **KYG** · Ophthalmic
Source: <https://www.510kdatabase.net/k802332/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Irrigation, Ocular Surgery (KYG)
Date received	Sep 23, 1980
Decision date	Dec 22, 1980
Days to decision	90 days
Third-party review	No

APPLICANT

Company	General Medical Engineering Corp.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k802332/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 1, 2026