

K840590 FLEISCHMAN-SWARTZ ENDO-OCULAR PROBEMay 9, 1984
90 days to decisionK840590 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k840590/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Laser, Ophthalmic (HQF) |
| Date received | Feb 9, 1984 |
| Decision date | May 9, 1984 |
| Days to decision | 90 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Hgm, Inc. |
| Location | Salt Lake City, UT, US |
| 510(k) history | 23 submissions · 23 cleared · 1983-1995 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k840590/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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