

K840854 MEDICAL LASER ENDOCOAGULATOR 5KApr 2, 1984
38 days to decisionK840854 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k840854/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Feb 24, 1984
Decision date	Apr 2, 1984
Days to decision	38 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k840854/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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