

**K931784 HGM ILLUMINATING IMAGING ENDOOCULAR PROBE**Mar 1, 1994  
326 days to decisionK931784 · Product code: **HQF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k931784/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Apr 9, 1993
Decision date	Mar 1, 1994
Days to decision	326 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Hgm, Inc.</b>
Location	Salt Lake City, UT, US
Contact	JOSEPH G LAMBERT
510(k) history	23 submissions · 23 cleared · 1983-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931784/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 21, 2026