FDA 510(k) Premarket Notification

K840590 FLEISCHMAN-SWARTZ ENDO-OCULAR PROBE

CLEARED

May 9, 1984
90 days to decision

K840590 · 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