FDA 510(k) Premarket Notification

K931784 HGM ILLUMINATING IMAGING ENDOCULAR PROBE

CLEARED

Mar 1, 1994
326 days to decision

K931784 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k931784/

SUBMISSION DETAILS

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

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