FDA 510(k) Premarket Notification

K840854 MEDICAL LASER ENDOCOAGULATOR 5K

CLEARED

Apr 2, 1984
38 days to decision

K840854 · 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