

K960522 DIOPEXY PROBEJul 19, 1996
164 days to decisionK960522 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k960522/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Feb 6, 1996
Decision date	Jul 19, 1996
Days to decision	164 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Iriderm Div.
Location	Mountain View, CA, US
Contact	THEODORE A BOUTACOFF
510(k) history	10 submissions · 10 cleared · 1989-1997

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