FDA 510(k) Premarket Notification

K922253 KERATOREF L60

CLEARED

Jun 19, 1992
49 days to decision

K922253 · Product code: **HLQ** · Ophthalmic
Source: https://www.510kdatabase.net/k922253/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratoscope, Ac-powered (HLQ)
Date received	May 1, 1992
Decision date	Jun 19, 1992
Days to decision	49 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Luneau Ophtalmologie SA**
Location	Chartres Cedex, FR
Contact	JEAN-NOEL YOUNG
510(k) history	3 submissions · 3 cleared · 1991-1992