

**K922253 KERATOREF L60**Jun 19, 1992  
49 days to decisionK922253 · Product code: **HLQ** · Ophthalmic  
Source: <https://www.510kdatabase.net/k922253/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k922253/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026