

**K923708 LEMPERT OPTIC DISC ANALYSIS SYSTEM**Mar 19, 1993  
235 days to decisionK923708 · Product code: **HKI** · Ophthalmic  
Source: <https://www.510kdatabase.net/k923708/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Ophthalmic, Ac-powered (HKI)
Date received	Jul 27, 1992
Decision date	Mar 19, 1993
Days to decision	235 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Lempert Ophthalmic Instruments</b>
Location	Ithaca, NY, US
Contact	PHILIP LEMPERT
510(k) history	1 submissions · 1 cleared · 1993-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923708/> 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