

**K000731 MODIFICATION TO RTA RETINAL THICKNESS
ANALYZER**Mar 31, 2000
25 days to decisionK000731 · Product code: **HLI** · Ophthalmic
Source: <https://www.510kdatabase.net/k000731/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Ophthalmoscope, Ac-powered (HLI)
Date received	Mar 6, 2000
Decision date	Mar 31, 2000
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Talia Technology , Ltd.
Location	Washington, DC, US
Contact	JONATHAN S KAHAN
510(k) history	6 submissions · 6 cleared · 1994-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k000731/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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