

**K023249 ALCON LADARWAVE CUSTOMCORNEA WAVEFRONT SYSTEM**Oct 18, 2002  
18 days to decisionK023249 · Product code: **NCF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k023249/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Aberrometer, Ophthalmic (NCF)
Date received	Sep 30, 2002
Decision date	Oct 18, 2002
Days to decision	18 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Alcon Laboratories, Inc.</b>
Location	Fort Worth, TX, US
Contact	CHRISTY STEVENS
Website	<a href="https://www.alcon.com">https://www.alcon.com</a>
510(k) history	43 submissions · 42 cleared · 1996-2026

Alcon Laboratories, Inc. is an eye care company headquartered in Fort Worth, Texas. The company develops innovative vision products and treatments for patients worldwide. Alcon maintains a strong FDA 510(k) regulatory record with FDA 510(k) cleared devices from total submissions. The company specializes exclusively in Ophthalmic devices, a focus reflected across its entire submission portfolio. Alcon's regulatory activity spans from 1996 to 2026, with recent clearances demonstrating continued innovation in vision care technologies. Recent FDA 510(k) clearances include con...

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Device record: <https://www.510kdatabase.net/k023249/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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