

K191795 AOCup Lens Case with AODiscJul 31, 2019
28 days to decisionK191795 · Product code: **LPN** · Ophthalmic
Source: <https://www.510kdatabase.net/k191795/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Accessories, Soft Lens Products (LPN)
Date received	Jul 3, 2019
Decision date	Jul 31, 2019
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Alcon Laboratories, Inc.
Location	Fort Worth, TX, US
Contact	Katryna Warren
Website	https://www.alcon.com
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...

REGULATORY CONSULTANT

Consulting firm	Ciba Vision GmbH
Contact	Ralf Finke

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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Device record: <https://www.510kdatabase.net/k191795/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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