

**K972454 OXYCOR (BALAFILCON A) VISIBILITY TINTED CONTACT LENS**Aug 8, 1997  
38 days to decisionK972454 · Product code: LPL · Ophthalmic  
Source: <https://www.510kdatabase.net/k972454/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lenses, Soft Contact, Daily Wear (LPL)
Date received	Jul 1, 1997
Decision date	Aug 8, 1997
Days to decision	38 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bausch &amp; Lomb, Inc.</b>
Location	Mchenry, IL, US
Contact	DENNIS HAHN
Website	<a href="http://www.bausch.com">http://www.bausch.com</a>
510(k) history	92 submissions · 92 cleared · 1977-2019

Bausch & Lomb, Inc. is a Canadian eye health company founded in 1853. The company is now part of Valeant Pharmaceuticals following a 2013 acquisition. Bausch & Lomb has received FDA 510(k) clearances from total submissions since 1977. The company specializes in Ophthalmic devices, which represent 83% of its regulatory submissions. Recent cleared devices include contact lenses, intraocular lens injectors, lens delivery systems, and care solutions. The company's last FDA 510(k) clearance was in 2019, and this profile reflects its historical regulatory record. Bausch & Lomb ...

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k972454/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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