FDA 510(k) Premarket Notification

CLEARED

K050948 PUREVISION MULTI-FOCAL (BALAFILCON A) VISIBILITY TINTED CONTACT LENS

May 18, 2005
33 days to decision

K050948 · Product code: **LPL** · Ophthalmic
Source: https://www.510kdatabase.net/k050948/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Lenses, Soft Contact, Daily Wear (LPL)
Date received	Apr 15, 2005
Decision date	May 18, 2005
Days to decision	33 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bausch & Lomb, Inc.**
Location	Mchenry, IL, US
Contact	ANNE ZAVERTNIK
Website	http://www.bausch.com
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 ...