FDA 510(k) Premarket Notification

K063694 AKREOS SINGLE USE INSERTION DEVICE, MODEL AI-28

CLEARED

Mar 7, 2007
84 days to decision

K063694 · Product code: **KYB** · Ophthalmic
Source: https://www.510kdatabase.net/k063694/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Lens, Guide, Intraocular (KYB)
Date received	Dec 13, 2006
Decision date	Mar 7, 2007
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bausch & Lomb, Inc.**
Location	Mchenry, IL, US
Contact	NANETTE CANEPA
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 ...