

**K131208 BAUSCH + LOMB PROCOMFORT (SAMFILCON A)
CONTACT LENS, BAUSCH + LOMB PROCOMFORT
(SAMFILCON A) CONTACT LENS FOR PRESBYOPIA,**Sep 11, 2013
135 days to decisionK131208 · Product code: LPL · Ophthalmic
Source: <https://www.510kdatabase.net/k131208/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lenses, Soft Contact, Daily Wear (LPL)
Date received	Apr 29, 2013
Decision date	Sep 11, 2013
Days to decision	135 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bausch & Lomb, Inc.
Location	Mchenry, IL, US
Contact	Barbara Klube-Falso
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 ...