FDA 510(k) Premarket Notification

K862517 BAUSCH & LOMB CONTACT LENS VIEWER

CLEARED

Aug 11, 1986
41 days to decision

K862517 · Product code: **HLF** · Ophthalmic
Source: https://www.510kdatabase.net/k862517/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Measuring, Lens Radius, Ophthalmic (HLF)
Date received	Jul 1, 1986
Decision date	Aug 11, 1986
Days to decision	41 days
Third-party review	No

APPLICANT

Company	**Bausch & Lomb, Inc.**
Location	Mchenry, IL, US
Contact	NICHOLAS A WALP
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 ...