FDA 510(k) Premarket Notification

K890514 PRE-MOISTENED CLEANSING PAD

CLEARED

Apr 10, 1989
68 days to decision

K890514 · Product code: **LKB** · General Hospital
Source: https://www.510kdatabase.net/k890514/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pad, Alcohol, Device Disinfectant (LKB)
Date received	Feb 1, 1989
Decision date	Apr 10, 1989
Days to decision	68 days
Third-party review	No

APPLICANT

Company	**Bausch & Lomb, Inc.**
Location	Mchenry, IL, US
Contact	DAVIES, O.D.
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 ...