

K896685 BAUSCH & LOMB FREQUENT REPLACEMENT CASE SYSTEMDec 26, 1989
28 days to decisionK896685 · Product code: **LRX** · Ophthalmic
Source: <https://www.510kdatabase.net/k896685/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Case, Contact Lens (LRX) |
| Date received | Nov 28, 1989 |
| Decision date | Dec 26, 1989 |
| Days to decision | 28 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Bausch & Lomb, Inc. |
| Location | Mchenry, IL, US |
| Contact | DARLA J ELKIN |
| 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 ...

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