

K173365 Bausch + Lomb Boston Scleral Lens CaseNov 17, 2017
22 days to decisionK173365 · Product code: **LRX** · Ophthalmic
Source: <https://www.510kdatabase.net/k173365/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Case, Contact Lens (LRX) |
| Date received | Oct 26, 2017 |
| Decision date | Nov 17, 2017 |
| Days to decision | 22 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Bausch & Lomb, Incorporated |
| Location | Rochester, NY, US |
| Contact | Nancy Fehrman |
| 510(k) history | 27 submissions · 27 cleared · 2002-2024 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173365/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 31, 2026