

**K220613 Bausch + Lomb (kalifilcon A) Soft (hydrophilic) Contact Lens, Bausch + Lomb (kalifilcon A) Soft (hydrophilic) Contact Lens for Astigmatism, and, Bausch + Lomb (kalifilcon A) Soft (hydrophilic) Contact Lens for Presbyopia**Mar 8, 2023  
370 days to decisionK220613 · Product code: LPL · Ophthalmic  
Source: <https://www.510kdatabase.net/k220613/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)     |
| Submission type       | Traditional                            |
| Device classification | Lenses, Soft Contact, Daily Wear (LPL) |
| Date received         | Mar 3, 2022                            |
| Decision date         | Mar 8, 2023                            |
| Days to decision      | 370 days                               |
| Third-party review    | No                                     |
| Combination product   | No                                     |
| PCCP authorized       | No                                     |
| Summary / Statement   | Summary                                |

**APPLICANT**

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|----------------|---|
| Company        | <b>Bausch &amp; Lomb, Incorporated</b>  |
| Location       | Rochester, NY, US                       |
| Contact        | Barbara Klube-Falso                     |
| 510(k) history | 27 submissions · 27 cleared · 2002-2024 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k220613/> 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](http://accessdata.fda.gov). - Generated May 13, 2026