

K161819 NNR06 Multi-Purpose SolutionNov 25, 2016
147 days to decisionK161819 · Product code: **LPN** · Ophthalmic
Source: <https://www.510kdatabase.net/k161819/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Soft Lens Products (LPN)
Date received	Jul 1, 2016
Decision date	Nov 25, 2016
Days to decision	147 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k161819/> 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 June 2, 2026