

**K002319 PURILENS SALINE SOLUTION**Aug 30, 2000  
30 days to decisionK002319 · Product code: **LPN** · Ophthalmic  
Source: <https://www.510kdatabase.net/k002319/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Soft Lens Products (LPN)
Date received	Jul 31, 2000
Decision date	Aug 30, 2000
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Purilens, Inc.</b>
Location	New Port Riche, FL, US
Contact	ART WARD
510(k) history	2 submissions · 2 cleared · 1999-2000

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k002319/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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 June 29, 2026