FDA 510(k) Premarket Notification

K002319 PURILENS SALINE SOLUTION

CLEARED

Aug 30, 2000
30 days to decision

K002319 · 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	**Purilens, Inc.**
Location	New Port Riche, FL, US
Contact	ART WARD
510(k) history	2 submissions · 2 cleared · 1999-2000