

**K992028 REFRESH CONTACTS LUBRICATING AND
REWETTING DROPS**Oct 7, 1999
113 days to decisionK992028 · Product code: LPN · Ophthalmic
Source: <https://www.510kdatabase.net/k992028/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Accessories, Soft Lens Products (LPN)
Date received	Jun 16, 1999
Decision date	Oct 7, 1999
Days to decision	113 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Allergan, Inc.
Location	Walker, MI, US
Contact	PAUL J NOWACKI
Website	http://www.allergan.com
510(k) history	33 submissions · 33 cleared · 1982-2019

Allergan, Inc. was an American global pharmaceutical company headquartered in Walker, US. The company focused on eye care, neurosciences, medical dermatology, and medical aesthetics before ceasing independent operations in 2015. Allergan received FDA 510(k) clearances from total submissions between 1982 and 2019. The company's regulatory portfolio was dominated by Ophthalmic devices, which accounted for 88% of all submissions. Notable cleared products include the XEN Glaucoma Treatment System, REFRESH rewetting drops, and the Natrelle tissue expander for surgical applicat...