FDA 510(k) Premarket Notification

K821051 LIQUIFILM WETTING SOLUTION

CLEARED

May 13, 1982
28 days to decision

K821051 · Product code: **HPX** · Ophthalmic
Source: https://www.510kdatabase.net/k821051/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Contact (polymethylmethacrylate) (HPX)
Date received	Apr 15, 1982
Decision date	May 13, 1982
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Allergan, Inc.**
Location	Walker, MI, US
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...