

**K991620 VISINE FOR CONTACTS**Sep 24, 1999  
136 days to decisionK991620 · Product code: **LPN** · Ophthalmic  
Source: <https://www.510kdatabase.net/k991620/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Soft Lens Products (LPN)
Date received	May 11, 1999
Decision date	Sep 24, 1999
Days to decision	136 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pfizer, Inc.</b>
Location	Mchenry, IL, US
Contact	JOHN TOMASZEWSKI
510(k) history	30 submissions · 30 cleared · 1977-2018

Pfizer, Inc. is an American multinational pharmaceutical and biotechnology corporation headquartered in Manhattan, New York City. Founded in 1849, Pfizer is one of the oldest pharmaceutical companies in North America. Pfizer's FDA 510(k) regulatory record includes cleared devices from total submissions, spanning 1977 to 2018. The company's device portfolio demonstrates strength in orthopedic devices, including surgical implants and fixation systems. This regulatory activity is now historical, with no clearances recorded in the past five years. The company's cleared device...

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