

**K993039 VISCOUS FLUID INJECTOR SYSTEM MODULE  
(CX5700)**Jan 7, 2000  
119 days to decisionK993039 · Product code: **MRH** · Ophthalmic  
Source: <https://www.510kdatabase.net/k993039/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion, Ophthalmic (MRH)
Date received	Sep 10, 1999
Decision date	Jan 7, 2000
Days to decision	119 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bausch &amp; Lomb, Inc.</b>
Location	Mchenry, IL, US
Contact	VANADA JOHNSON
Website	<a href="http://www.bausch.com">http://www.bausch.com</a>
510(k) history	92 submissions · 92 cleared · 1977-2019

Bausch & Lomb, Inc. is a Canadian eye health company founded in 1853. The company is now part of Valeant Pharmaceuticals following a 2013 acquisition. Bausch & Lomb has received FDA 510(k) clearances from total submissions since 1977. The company specializes in Ophthalmic devices, which represent 83% of its regulatory submissions. Recent cleared devices include contact lenses, intraocular lens injectors, lens delivery systems, and care solutions. The company's last FDA 510(k) clearance was in 2019, and this profile reflects its historical regulatory record. Bausch & Lomb ...

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k993039/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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