

**K131958 BAUSCH & LOMB INJECTOR SYSTEM**Jan 9, 2014  
196 days to decisionK131958 · Product code: **MSS** · Ophthalmic  
Source: <https://www.510kdatabase.net/k131958/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Folders And Injectors, Intraocular Lens (iol) (MSS)
Date received	Jun 27, 2013
Decision date	Jan 9, 2014
Days to decision	196 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bausch &amp; Lomb</b>
Location	Rochester, NY, US
Contact	JASON SMITH
Website	<a href="http://www.bausch.com">http://www.bausch.com</a>
510(k) history	12 submissions · 12 cleared · 2002-2017

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131958/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 1, 2026