FDA 510(k) Premarket Notification

K131958 BAUSCH & LOMB INJECTOR SYSTEM

CLEARED

Jan 9, 2014
196 days to decision

K131958 · Product code: **MSS** · Ophthalmic
Source: https://www.510kdatabase.net/k131958/

SUBMISSION DETAILS

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

Company	**Bausch & Lomb**
Location	Rochester, NY, US
Contact	JASON SMITH
Website	http://www.bausch.com
510(k) history	12 submissions · 12 cleared · 2002-2017