FDA 510(k) Premarket Notification

K173365 Bausch + Lomb Boston Scleral Lens Case

CLEARED

Nov 17, 2017
22 days to decision

K173365 · Product code: **LRX** · Ophthalmic
Source: https://www.510kdatabase.net/k173365/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Case, Contact Lens (LRX)
Date received	Oct 26, 2017
Decision date	Nov 17, 2017
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bausch & Lomb, Incorporated**
Location	Rochester, NY, US
Contact	Nancy Fehrman
510(k) history	27 submissions · 27 cleared · 2002-2024