FDA 510(k) Premarket Notification

K171404 BOSTON XO (hexafocon A), BOSTON XO2 (hexafocon B)

CLEARED

Jul 17, 2017
66 days to decision

K171404 · Product code: **HQD** · Ophthalmic
Source: https://www.510kdatabase.net/k171404/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Contact (other Material) - Daily (HQD)
Date received	May 12, 2017
Decision date	Jul 17, 2017
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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