

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

Jul 17, 2017  
66 days to decision

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

**SUBMISSION DETAILS**

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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**

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Company	<b>Bausch &amp; Lomb, Incorporated</b>
Location	Rochester, NY, US
Contact	Glenn A. Davies
510(k) history	27 submissions · 27 cleared · 2002-2024

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

Device record: <https://www.510kdatabase.net/k171404/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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