

**K781658 PROJECTOR, ACCU-CHART ACUITY**Oct 17, 1978  
19 days to decisionK781658 · Product code: **HOS** · Ophthalmic  
Source: <https://www.510kdatabase.net/k781658/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Projector, Ophthalmic (HOS)
Date received	Sep 28, 1978
Decision date	Oct 17, 1978
Days to decision	19 days
Third-party review	No

**APPLICANT**

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Company	<b>Bausch &amp; Lomb, Inc.</b>
Location	Mchenry, IL, US
Website	<a href="http://www.bausch.com">http://www.bausch.com</a>
510(k) history	92 submissions · 92 cleared · 1977-2019

Bausch & Lomb, Inc. is a Canadian eye health company founded in 1853. The company is now part of Valeant Pharmaceuticals following a 2013 acquisition. Bausch & Lomb has received FDA 510(k) clearances from total submissions since 1977. The company specializes in Ophthalmic devices, which represent 83% of its regulatory submissions. Recent cleared devices include contact lenses, intraocular lens injectors, lens delivery systems, and care solutions. The company's last FDA 510(k) clearance was in 2019, and this profile reflects its historical regulatory record. Bausch & Lomb ...