

**K923735 SOUNDCHOICE K-AMP**Sep 28, 1992  
63 days to decisionK923735 · Product code: **ESD** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k923735/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jul 27, 1992
Decision date	Sep 28, 1992
Days to decision	63 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Bausch &amp; Lomb, Inc.</b>
Location	Mchenry, IL, US
Contact	TERRENCE MARTIN
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 ...

---