

K910610 INTERPLAK HOME PLAQUE REMOVAL INSTRUMENTJul 9, 1991
147 days to decisionK910610 · Product code: **JEQ** · Dental
Source: <https://www.510kdatabase.net/k910610/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Toothbrush, Powered (JEQ)
Date received	Feb 12, 1991
Decision date	Jul 9, 1991
Days to decision	147 days
Third-party review	No

APPLICANT

Company	Bausch & Lomb, Inc.
Location	Mchenry, IL, US
Contact	STEPHENS
Website	http://www.bausch.com
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 ...

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