

**K012308 KAVO QUATTROCARE**Jun 26, 2002  
338 days to decisionK012308 · Product code: **EFB** · Dental  
Source: <https://www.510kdatabase.net/k012308/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Handpiece, Air-powered, Dental (EFB)
Date received	Jul 23, 2001
Decision date	Jun 26, 2002
Days to decision	338 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Kavo America</b>
Location	Washington, DC, US
Contact	JOHN WESTERMEIER
510(k) history	20 submissions · 19 cleared · 1991-2008

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012308/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 14, 2026