

**K961200 BOSWORTH LIQUID RESIN II**May 17, 1996  
51 days to decisionK961200 · Product code: **EBI** · DentalSource: <https://www.510kdatabase.net/k961200/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	Mar 27, 1996
Decision date	May 17, 1996
Days to decision	51 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Harry J. Bosworth Co.</b>
Location	Mchenry, IL, US
Contact	MILDRED M GOLDSTEIN
510(k) history	45 submissions · 45 cleared · 1979-2011

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k961200/> 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 June 19, 2026