FDA 510(k) Premarket Notification

# K961200 BOSWORTH LIQUID RESIN II

**CLEARED**

May 17, 1996
51 days to decision

K961200 · Product code: **EBI** · Dental
Source: 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 | **Harry J. Bosworth Co.** |
| Location | Mchenry, IL, US |
| Contact | MILDRED M GOLDSTEIN |
| 510(k) history | 45 submissions · 45 cleared · 1979-2011 |