

K831647 DENTURE RELINING, REPAIR/REBASING RESINJun 30, 1983
38 days to decisionK831647 · Product code: **EBI** · DentalSource: <https://www.510kdatabase.net/k831647/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	May 23, 1983
Decision date	Jun 30, 1983
Days to decision	38 days
Third-party review	No

APPLICANT

Company	Dentsply Intl.
Location	Walker, MI, US
510(k) history	279 submissions · 279 cleared · 1976-2013

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k831647/> Data sourced from FDA 510(k) public records (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. - Generated May 19, 2026