

K820215 IMPROVED DYCALMar 23, 1982
56 days to decisionK820215 · Product code: **EJK** · DentalSource: <https://www.510kdatabase.net/k820215/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Liner, Cavity, Calcium Hydroxide (EJK)
Date received	Jan 26, 1982
Decision date	Mar 23, 1982
Days to decision	56 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/k820215/> 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