

K001050 CALCIJECTJul 21, 2000
112 days to decisionK001050 · Product code: **EJK** · DentalSource: <https://www.510kdatabase.net/k001050/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Liner, Cavity, Calcium Hydroxide (EJK)
Date received	Mar 31, 2000
Decision date	Jul 21, 2000
Days to decision	112 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Centrix, Inc.
Location	Mchenry, IL, US
Contact	SCOTT COHN
510(k) history	47 submissions · 47 cleared · 1979-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001050/> 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 14, 2026