

K933999 STERIBASE (ACCUBASE)Jan 5, 1994
141 days to decisionK933999 · Product code: **EJK** · Dental
Source: <https://www.510kdatabase.net/k933999/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Liner, Cavity, Calcium Hydroxide (EJK)
Date received	Aug 17, 1993
Decision date	Jan 5, 1994
Days to decision	141 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k933999/> 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