FDA 510(k) Premarket Notification

K933999 STERIBASE (ACCUBASE)

CLEARED

Jan 5, 1994
141 days to decision

K933999 · 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