FDA 510(k) Premarket Notification

K921408 LIGHT CURE DENTIN AND LINER BASE PRODUCT LINE

CLEARED

Nov 24, 1992
245 days to decision

K921408 · Product code: **EJK** · Dental
Source: https://www.510kdatabase.net/k921408/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Liner, Cavity, Calcium Hydroxide (EJK)
Date received	Mar 24, 1992
Decision date	Nov 24, 1992
Days to decision	245 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ultradent Products, Inc.**
Location	Salt Lake City, UT, US
Contact	LINDA CHATWIN
Website	https://www.ultradent.com
510(k) history	103 submissions · 103 cleared · 1992-2026