FDA 510(k) Premarket Notification

K003383 PALMLIGHT

CLEARED

Feb 6, 2001
98 days to decision

K003383 · Product code: **EBZ** · Dental
Source: https://www.510kdatabase.net/k003383/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Activator, Ultraviolet, For Polymerization (EBZ)
Date received	Oct 31, 2000
Decision date	Feb 6, 2001
Days to decision	98 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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