

K003383 PALMLIGHTFeb 6, 2001
98 days to decisionK003383 · Product code: **EBZ** · DentalSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k003383/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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