FDA 510(k) Premarket Notification

K020091 OPTILUX 501

CLEARED

Mar 21, 2002
70 days to decision

K020091 · Product code: **EBZ** · Dental
Source: https://www.510kdatabase.net/k020091/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Activator, Ultraviolet, For Polymerization (EBZ)
Date received	Jan 10, 2002
Decision date	Mar 21, 2002
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Kerr Corporation (Danbury)**
Location	Mchenry, IL, US
Contact	COLLEEN BOSWELL
510(k) history	32 submissions · 32 cleared · 1978-2005

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k020091/ 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 July 1, 2026