

**K993683 PULPDENT UNO-DUO**Dec 13, 1999  
42 days to decisionK993683 · Product code: **KLE** · Dental  
Source: <https://www.510kdatabase.net/k993683/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agent, Tooth Bonding, Resin (KLE)
Date received	Nov 1, 1999
Decision date	Dec 13, 1999
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pulpdent Corp.</b>
Location	Watertown, MA, US
Contact	KENNETH J BERK
510(k) history	78 submissions · 78 cleared · 1990-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k993683/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 2, 2026