

**K905302 DENTHESIVE**Feb 26, 1991  
90 days to decisionK905302 · Product code: **KLE** · DentalSource: <https://www.510kdatabase.net/k905302/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agent, Tooth Bonding, Resin (KLE)
Date received	Nov 28, 1990
Decision date	Feb 26, 1991
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Kulzer, Inc.</b>
Location	Mchenry, IL, US
Contact	SHARON PARKER
510(k) history	31 submissions · 31 cleared · 1981-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905302/>; 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 4, 2026