

**K021083 VERACAM**Aug 30, 2002  
149 days to decisionK021083 · Product code: **EIA** · DentalSource: <https://www.510kdatabase.net/k021083/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Operative Dental (EIA)
Date received	Apr 3, 2002
Decision date	Aug 30, 2002
Days to decision	149 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Lumalite, Inc.</b>
Location	Spring Valley, CA, US
Contact	W. EDWARD JOHANSEN
510(k) history	3 submissions · 3 cleared · 1999-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k021083/> 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 1, 2026