

**K111017 CLEARTEMP**Jun 6, 2011  
55 days to decisionK111017 · Product code: **EMA** · Dental  
Source: <https://www.510kdatabase.net/k111017/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cement, Dental (EMA)
Date received	Apr 12, 2011
Decision date	Jun 6, 2011
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ultradent Products, Inc.</b>
Location	Salt Lake City, UT, US
Contact	DIANE ROGERS
Website	<a href="https://www.ultradent.com">https://www.ultradent.com</a>
510(k) history	103 submissions · 103 cleared · 1992-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k111017/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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