

**K111832 PALADON ULTRA POWDER (PINK), PALADON ULTRA LIQUID**

Sep 29, 2011  
93 days to decision

K111832 · Product code: **EBI** · Dental  
Source: <https://www.510kdatabase.net/k111832/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	Jun 28, 2011
Decision date	Sep 29, 2011
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Heraeus Kulzer, LLC</b>
Location	South Bend, IN, US
Contact	CHERYL ZIMMERMAN
510(k) history	11 submissions · 11 cleared · 2009-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)  
 Device record: <https://www.510kdatabase.net/k111832/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).  
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