

**K110037 PALAXPRESS ULTRA POWDER (PINK)**Apr 29, 2011  
113 days to decisionK110037 · Product code: **EBI** · DentalSource: <https://www.510kdatabase.net/k110037/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	Jan 6, 2011
Decision date	Apr 29, 2011
Days to decision	113 days
Third-party review	No
Summary / Statement	Summary
Other names	PALAXPRESS ULTRA LIQUID

**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/k110037/>; 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 June 9, 2026