

**K132815 PUREGRAFT 50 SYSTEM**Jan 24, 2014  
137 days to decisionK132815 · Product code: **MUU** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k132815/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Suction, Lipoplasty (MUU)
Date received	Sep 9, 2013
Decision date	Jan 24, 2014
Days to decision	137 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cytori Therapeutics, Inc.</b>
Location	San Diego, CA, US
Contact	KENNETH K KLEINHENZ
510(k) history	6 submissions · 6 cleared · 2006-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k132815/> 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 May 25, 2026