

**K132220 P&S SERENITY**Feb 20, 2015  
583 days to decisionK132220 · Product code: **HGX** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k132220/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Breast, Powered (HGX)
Date received	Jul 17, 2013
Decision date	Feb 20, 2015
Days to decision	583 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Limerick, Inc.</b>
Location	Burbank, CA, US
Contact	PATRICIA KELLY
510(k) history	4 submissions · 4 cleared · 2002-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k132220/> 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 5, 2026