

**K051295 I-D GLIDE**Feb 15, 2006  
273 days to decisionK051295 · Product code: **NUC** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k051295/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lubricant, Personal (NUC)
Date received	May 18, 2005
Decision date	Feb 15, 2006
Days to decision	273 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Westridge Laboratories, Inc.</b>
Location	Crofton, MD, US
Contact	YOLANDA SMITH
510(k) history	3 submissions · 3 cleared · 2006-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051295/> 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 26, 2026