

**K952465 PROBE PERSONAL LUBRICANT - SILKY LIGHT**Nov 22, 1995  
180 days to decisionK952465 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k952465/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	May 26, 1995
Decision date	Nov 22, 1995
Days to decision	180 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Davryan Laboratories, Inc.</b>
Location	Portland, OR, US
Contact	BRUCE CAMPBELL
510(k) history	2 submissions · 2 cleared · 1995-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k952465/> 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 29, 2026