

**K983181 CYTOPREP**Nov 16, 1998  
66 days to decisionK983181 · Product code: **HHI** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k983181/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Abortion, Vacuum (HHI)
Date received	Sep 11, 1998
Decision date	Nov 16, 1998
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cytoprep Inc. Est.</b>
Location	Eschen, LI
Contact	K G PAGANIS
510(k) history	1 submissions · 1 cleared · 1998-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k983181/> 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