

**K980212 MITYVAC**Jul 9, 1998  
169 days to decisionK980212 · Product code: **HDB** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k980212/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Extractor, Vacuum, Fetal (HDB)
Date received	Jan 21, 1998
Decision date	Jul 9, 1998
Days to decision	169 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Prism Technologies, Inc.</b>
Location	Chicago, IL, US
Contact	HOWARD M HOLSTEIN, ESQ.
510(k) history	5 submissions · 4 cleared · 1991-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980212/> 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