

**K991963 PATTON SPECULUM**Sep 1, 1999  
83 days to decisionK991963 · Product code: **HIB** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k991963/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Speculum, Vaginal, Nonmetal (HIB)
Date received	Jun 10, 1999
Decision date	Sep 1, 1999
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Patton Medical Corp.</b>
Location	Austin, TX, US
Contact	MICHAEL T PATTON
510(k) history	4 submissions · 4 cleared · 1999-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k991963/> 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