

**K904160 CYTOSMEAR BRUSH**Nov 2, 1990  
52 days to decisionK904160 · Product code: **HHI** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k904160/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Abortion, Vacuum (HHI)
Date received	Sep 11, 1990
Decision date	Nov 2, 1990
Days to decision	52 days
Third-party review	No

**APPLICANT**

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Company	<b>Brooke Intl.</b>
Location	Pasadena, CA, US
Contact	TODD M GATES
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904160/>, 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 30, 2026