

**K051594 MCCARTNEY ACCESS TUBE**Sep 2, 2005  
78 days to decisionK051594 · Product code: **HEW** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k051594/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Culdoscope (and Accessories) (HEW)
Date received	Jun 16, 2005
Decision date	Sep 2, 2005
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Gynetech Pty. , Ltd.</b>
Location	Cincinnati, OH, US
Contact	DENNIS HAHN
510(k) history	4 submissions · 4 cleared · 2005-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051594/> 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 4, 2026