

**K073053 COLPO-MASTER**Nov 15, 2007  
16 days to decisionK073053 · Product code: **HEX** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k073053/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Colposcope (and Colpomicroscope) (HEX)
Date received	Oct 30, 2007
Decision date	Nov 15, 2007
Days to decision	16 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>United Products &amp; Instruments, Inc.</b>
Location	Hasbrouck Heights, NJ, US
Contact	GEORGE MYERS
510(k) history	1 submissions · 1 cleared · 2007-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073053/> 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 26, 2026