

**K991322 ENHANCE-S PLUS**Jul 26, 1999  
98 days to decisionK991322 · Product code: **MQL** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k991322/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Reproductive (MQL)
Date received	Apr 19, 1999
Decision date	Jul 26, 1999
Days to decision	98 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Conception Technology, Inc.</b>
Location	Irvine, CA, US
Contact	GREG HOLLAND
510(k) history	5 submissions · 5 cleared · 1998-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k991322/> 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 28, 2026