

**K980237 GYNEX ENDOSPECULUM**Apr 8, 1998  
75 days to decisionK980237 · Product code: **KNA** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k980237/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, Specialized Obstetric-gynecologic (KNA)
Date received	Jan 23, 1998
Decision date	Apr 8, 1998
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Gynex Corp.</b>
Location	Redmond, WA, US
Contact	DAVID W SCHLERF
510(k) history	5 submissions · 5 cleared · 1998-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980237/> 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