

**K891351 NEWPORT LATERAL VAGINAL RETRACTOR**May 11, 1989  
62 days to decisionK891351 · Product code: **KOH** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k891351/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, General Obstetric-gynecologic (KOH)
Date received	Mar 10, 1989
Decision date	May 11, 1989
Days to decision	62 days
Third-party review	No

**APPLICANT**

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Company	<b>Simpson/Basye, Inc.</b>
Location	Mchenry, IL, US
Contact	ROBERT A BASYE
510(k) history	12 submissions · 12 cleared · 1978-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k891351/>; 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