

**K894264 KDF-2.3**Aug 15, 1989  
56 days to decisionK894264 · Product code: **HDR** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k894264/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cap, Cervical (HDR)
Date received	Jun 20, 1989
Decision date	Aug 15, 1989
Days to decision	56 days
Third-party review	No

**APPLICANT**

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Company	<b>Unimar, Inc.</b>
Location	Mchenry, IL, US
Contact	ANTHONY HEMMING
510(k) history	10 submissions · 9 cleared · 1981-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894264/>; 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