

**K900308 SPERM SELECT SYSTEM**Apr 13, 1990  
81 days to decisionK900308 · Product code: **HDR** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k900308/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cap, Cervical (HDR)
Date received	Jan 22, 1990
Decision date	Apr 13, 1990
Days to decision	81 days
Third-party review	No

**APPLICANT**

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Company	<b>Pharmacia, Inc.</b>
Location	Mchenry, IL, US
Contact	ALBERT P MAYO
510(k) history	129 submissions · 126 cleared · 1976-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900308/>; 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 May 14, 2026