

**K895530 STANDARD CONDOM**Nov 28, 1989  
76 days to decisionK895530 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k895530/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Sep 13, 1989
Decision date	Nov 28, 1989
Days to decision	76 days
Third-party review	No

**APPLICANT**

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Company	<b>Mayer Laboratories</b>
Location	Oakland, CA, US
Contact	DAVID P MAYER
510(k) history	16 submissions · 16 cleared · 1987-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895530/>, 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 20, 2026