

**K874236 APOLLO CONDOM**Jul 1, 1988  
260 days to decisionK874236 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k874236/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Oct 15, 1987
Decision date	Jul 1, 1988
Days to decision	260 days
Third-party review	No

**APPLICANT**

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Company	<b>Apollo Enterprise, Ltd. Co.</b>
Location	Miami, FL, US
Contact	HOWARD DOLLINGER, DDS
510(k) history	3 submissions · 3 cleared · 1988-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874236/>; 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