

**K902936 POLYURETHANE CONDOM**Mar 6, 1991  
246 days to decisionK902936 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k902936/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Jul 3, 1990
Decision date	Mar 6, 1991
Days to decision	246 days
Third-party review	No

**APPLICANT**

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Company	<b>Apex Medical Technologies, Inc.</b>
Location	San Diego, CA, US
Contact	ALICE A DEPAUL
510(k) history	8 submissions · 8 cleared · 1987-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902936/>; 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 14, 2026