

**K905279 CONDOM (RUBBER) CONTRACEPTIVE**Feb 19, 1991  
88 days to decisionK905279 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k905279/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Nov 23, 1990
Decision date	Feb 19, 1991
Days to decision	88 days
Third-party review	No

**APPLICANT**

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Company	<b>Aladan Corp.</b>
Location	Jessup, MD, US
Contact	BRADLEY PUGH
510(k) history	18 submissions · 18 cleared · 1989-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905279/>, 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 29, 2026