

**K896107 RUBBER CONTRACEPTIVE**Jun 29, 1990  
249 days to decisionK896107 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k896107/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Oct 23, 1989
Decision date	Jun 29, 1990
Days to decision	249 days
Third-party review	No

**APPLICANT**

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Company	<b>Banrub Sdn Bhd</b>
Location	Johor Bahru, J.D.T., MY
Contact	GOH SIANG
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896107/>; 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