

**K813172 CONDOM-MULTIPAL STYLES**Mar 5, 1982  
109 days to decisionK813172 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k813172/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Nov 16, 1981
Decision date	Mar 5, 1982
Days to decision	109 days
Third-party review	No

**APPLICANT**

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Company	<b>Circle Rubber Corp.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1982-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k813172/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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