

**K833765 TROJAN PLUS II**Dec 16, 1983  
50 days to decisionK833765 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k833765/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Oct 27, 1983
Decision date	Dec 16, 1983
Days to decision	50 days
Third-party review	No

**APPLICANT**

---

Company	<b>Youngs Rubber Corp.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

---

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

Device record: <https://www.510kdatabase.net/k833765/> 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