FDA 510(k) Premarket Notification

K833765 TROJAN PLUS II

CLEARED

Dec 16, 1983
50 days to decision

K833765 · Product code: **HIS** · Obstetrics & Gynecology
Source: 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	**Youngs Rubber Corp.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983