FDA 510(k) Premarket Notification

K813172 CONDOM-MULTIPAL STYLES

CLEARED

Mar 5, 1982
109 days to decision

K813172 · Product code: **HIS** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k813172/

SUBMISSION DETAILS

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

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