FDA 510(k) Premarket Notification

K895530 STANDARD CONDOM

CLEARED

Nov 28, 1989
76 days to decision

K895530 · Product code: **HIS** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k895530/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Sep 13, 1989
Decision date	Nov 28, 1989
Days to decision	76 days
Third-party review	No

APPLICANT

Company	**Mayer Laboratories**
Location	Oakland, CA, US
Contact	DAVID P MAYER
510(k) history	16 submissions · 16 cleared · 1987-2001