FDA 510(k) Premarket Notification

K905279 CONDOM (RUBBER) CONTRACEPTIVE

CLEARED

Feb 19, 1991
88 days to decision

K905279 · Product code: **HIS** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k905279/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Nov 23, 1990
Decision date	Feb 19, 1991
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Aladan Corp.**
Location	Jessup, MD, US
Contact	BRADLEY PUGH
510(k) history	18 submissions · 18 cleared · 1989-1996