FDA 510(k) Premarket Notification

K952465 PROBE PERSONAL LUBRICANT - SILKY LIGHT

CLEARED

Nov 22, 1995
180 days to decision

K952465 · Product code: **HIS** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k952465/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	May 26, 1995
Decision date	Nov 22, 1995
Days to decision	180 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Davryan Laboratories, Inc.**
Location	Portland, OR, US
Contact	BRUCE CAMPBELL
510(k) history	2 submissions · 2 cleared · 1995-1995