FDA 510(k) Premarket Notification

K051295 I-D GLIDE

CLEARED

Feb 15, 2006
273 days to decision

K051295 · Product code: **NUC** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k051295/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lubricant, Personal (NUC)
Date received	May 18, 2005
Decision date	Feb 15, 2006
Days to decision	273 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Westridge Laboratories, Inc.**
Location	Crofton, MD, US
Contact	YOLANDA SMITH
510(k) history	3 submissions · 3 cleared · 2006-2024