FDA 510(k) Premarket Notification

K013817 GLO-SPEC

CLEARED

Dec 21, 2001
35 days to decision

K013817 · Product code: **HIB** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k013817/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Speculum, Vaginal, Nonmetal (HIB)
Date received	Nov 16, 2001
Decision date	Dec 21, 2001
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sol Weiss MD, Inc.**
Location	Reseda, CA, US
Contact	SOL WEISS
510(k) history	3 submissions · 3 cleared · 2001-2002