

K013817 GLO-SPECDec 21, 2001
35 days to decisionK013817 · Product code: **HIB** · Obstetrics & GynecologySource: <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

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