

K252208 JFord Speculum Sleeve™Mar 6, 2026
234 days to decisionK252208 · Product code: **HIB** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k252208/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Speculum, Vaginal, Nonmetal (HIB)
Date received	Jul 15, 2025
Decision date	Mar 6, 2026
Days to decision	234 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Ford Medtech, LLC
Location	Westbury, NY, US
Contact	Jacqueline Ford
510(k) history	1 submissions · 1 cleared · 2026-2026

REGULATORY CONSULTANT

Consulting firm	The Jans Group, LLC
Contact	Michele Jans

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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