

**K022948 VAGINAL SPECULUM, MODEL 5555S, 5555M, OR 5555L**Nov 4, 2002  
60 days to decisionK022948 · Product code: **HIB** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k022948/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Speculum, Vaginal, Nonmetal (HIB)
Date received	Sep 5, 2002
Decision date	Nov 4, 2002
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medical Action Industries, Inc.</b>
Location	Mchenry, IL, US
Contact	SHARON SUESS GRAHAM
510(k) history	8 submissions · 8 cleared · 1981-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022948/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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