FDA 510(k) Premarket Notification

K022948 VAGINAL SPECULUM, MODEL 5555S, 5555M, OR 5555L

CLEARED

Nov 4, 2002
60 days to decision

K022948 · Product code: **HIB** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k022948/

SUBMISSION DETAILS

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

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