FDA 510(k) Premarket Notification

K891041 MIYA SPECULUM

CLEARED

Mar 22, 1989
22 days to decision

K891041 · Product code: **HDF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k891041/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Speculum, Vaginal, Metal (HDF)
Date received	Feb 28, 1989
Decision date	Mar 22, 1989
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Zinnanti Surgical Instruments, Inc.**
Location	Mchenry, IL, US
Contact	ANNA STRAIGHT
510(k) history	76 submissions · 76 cleared · 1984-1997