FDA 510(k) Premarket Notification

K991963 PATTON SPECULUM

CLEARED

Sep 1, 1999
83 days to decision

K991963 · Product code: **HIB** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k991963/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Speculum, Vaginal, Nonmetal (HIB)
Date received	Jun 10, 1999
Decision date	Sep 1, 1999
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Patton Medical Corp.**
Location	Austin, TX, US
Contact	MICHAEL T PATTON
510(k) history	4 submissions · 4 cleared · 1999-2002