FDA 510(k) Premarket Notification

K900642 SWIVEL ADAPTER

CLEARED

Mar 19, 1990
38 days to decision

K900642 · Product code: **HHI** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k900642/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Abortion, Vacuum (HHI)
Date received	Feb 9, 1990
Decision date	Mar 19, 1990
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Williams Specialty Products, Inc.**
Location	Encino, CA, US
Contact	WILLIAMS, MD
510(k) history	2 submissions · 2 cleared · 1990-1990