FDA 510(k) Premarket Notification

K871683 COLPOSTAR V-6

CLEARED

Aug 5, 1987
98 days to decision

K871683 · Product code: **HEX** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k871683/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Colposcope (and Colpomicroscope) (HEX)
Date received	Apr 29, 1987
Decision date	Aug 5, 1987
Days to decision	98 days
Third-party review	No

APPLICANT

Company	**Wallach Surgical Devices, Inc.**
Location	Walker, MI, US
Contact	VINCENT FRAZZETTA
510(k) history	27 submissions · 27 cleared · 1981-2002

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k871683/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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