FDA 510(k) Premarket Notification

K896065 WALLACH CC/BRUSH

CLEARED

Jan 4, 1990
78 days to decision

K896065 · Product code: **HHT** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k896065/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spatula, Cervical, Cytological (HHT)
Date received	Oct 18, 1989
Decision date	Jan 4, 1990
Days to decision	78 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/k896065/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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