

K897070 ENDOBRUSH(TM) DEVICESep 10, 1990
262 days to decisionK897070 · Product code: **HFE** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k897070/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Brush, Endometrial (HFE)
Date received	Dec 22, 1989
Decision date	Sep 10, 1990
Days to decision	262 days
Third-party review	No
Other names	OMNIBRUSH

APPLICANT

Company	Intl. Cytobrush, Inc.
Location	Hollywood, FL, US
Contact	TSE, PH.D.
510(k) history	4 submissions · 4 cleared · 1986-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k897070/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026