

K881921 C-E BRUSHJul 19, 1988
71 days to decisionK881921 · Product code: **HHT** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k881921/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spatula, Cervical, Cytological (HHT)
Date received	May 9, 1988
Decision date	Jul 19, 1988
Days to decision	71 days
Third-party review	No

APPLICANT

Company	Surgipath Medical Industries, Inc.
Location	Mchenry, IL, US
Contact	JOY F MONEK
510(k) history	30 submissions · 30 cleared · 1977-1988

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881921/>, 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 May 14, 2026