FDA 510(k) Premarket Notification

K881921 C-E BRUSH

CLEARED

Jul 19, 1988
71 days to decision

K881921 · Product code: **HHT** · Obstetrics & Gynecology
Source: 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