FDA 510(k) Premarket Notification

K862476 LS3000 CANNULA UTERINE SUCTION

CLEARED

Jul 18, 1986
18 days to decision

K862476 · Product code: **HHI** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k862476/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Abortion, Vacuum (HHI)
Date received	Jun 30, 1986
Decision date	Jul 18, 1986
Days to decision	18 days
Third-party review	No

APPLICANT

Company	**Engineering, Inc.**
Location	Foster City, CA, US
Contact	ROBERT LASH
510(k) history	5 submissions · 4 cleared · 1986-1988