FDA 510(k) Premarket Notification

K800834 SER-TEX

CLEARED

Jun 30, 1980
77 days to decision

K800834 · Product code: **KOH** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k800834/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, General Obstetric-gynecologic (KOH)
Date received	Apr 14, 1980
Decision date	Jun 30, 1980
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**V.S.S.I., Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980