

K800834 SER-TEXJun 30, 1980
77 days to decisionK800834 · Product code: **KOH** · Obstetrics & GynecologySource: <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.I., Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k800834/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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