

K790851 VAGI-PRESS #401Jul 30, 1979
87 days to decisionK790851 · Product code: **KOH** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k790851/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, General Obstetric-gynecologic (KOH)
Date received	May 4, 1979
Decision date	Jul 30, 1979
Days to decision	87 days
Third-party review	No

APPLICANT

Company	C. Intl.
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1979-1979

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Device record: <https://www.510kdatabase.net/k790851/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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