

K823792 QUARTZ RETRACTOR/PROBEJan 18, 1983
34 days to decisionK823792 · Product code: **KOH** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k823792/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, General Obstetric-gynecologic (KOH)
Date received	Dec 15, 1982
Decision date	Jan 18, 1983
Days to decision	34 days
Third-party review	No

APPLICANT

Company	American V. Mueller
Location	Mchenry, IL, US
510(k) history	31 submissions · 29 cleared · 1980-1987

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Device record: <https://www.510kdatabase.net/k823792/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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