

K802313 UTERINE CURETTEJan 12, 1981
111 days to decisionK802313 · Product code: **HCY** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k802313/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Curette, Uterine (HCY)
Date received	Sep 23, 1980
Decision date	Jan 12, 1981
Days to decision	111 days
Third-party review	No

APPLICANT

Company	Gynometrics, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k802313/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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