FDA 510(k) Premarket Notification

K802313 UTERINE CURETTE

CLEARED

Jan 12, 1981
111 days to decision

K802313 · Product code: **HCY** · Obstetrics & Gynecology
Source: 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	**Gynemetrics, Inc.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1981