

K800162 ONCE TMFeb 11, 1980
17 days to decisionK800162 · Product code: **HDW** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k800162/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Diaphragm, Contraceptive (and Accessories) (HDW)
Date received	Jan 25, 1980
Decision date	Feb 11, 1980
Days to decision	17 days
Third-party review	No

APPLICANT

Company	G.D. Searle and Co.
Location	Mchenry, IL, US
510(k) history	56 submissions · 56 cleared · 1976-1982

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Device record: <https://www.510kdatabase.net/k800162/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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