FDA 510(k) Premarket Notification

K800162 ONCE TM

CLEARED

Feb 11, 1980
17 days to decision

K800162 · Product code: **HDW** · Obstetrics & Gynecology
Source: 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