FDA 510(k) Premarket Notification

K821297 GRAVES SPECULA

CLEARED

May 24, 1982
20 days to decision

K821297 · Product code: **HDF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k821297/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Speculum, Vaginal, Metal (HDF)
Date received	May 4, 1982
Decision date	May 24, 1982
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Conphar, Inc.**
Location	Mchenry, IL, US
510(k) history	122 submissions · 122 cleared · 1979-1982