

K821297 GRAVES SPECULAMay 24, 1982
20 days to decisionK821297 · Product code: **HDF** · Obstetrics & GynecologySource: <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

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Device record: <https://www.510kdatabase.net/k821297/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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