

K820637 CATH-HOLDMar 25, 1982
17 days to decisionK820637 · Product code: **KOD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k820637/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Mar 8, 1982
Decision date	Mar 25, 1982
Days to decision	17 days
Third-party review	No

APPLICANT

Company	Womco Products
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

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Device record: <https://www.510kdatabase.net/k820637/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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