FDA 510(k) Premarket Notification

K820637 CATH-HOLD

CLEARED

Mar 25, 1982
17 days to decision

K820637 · Product code: **KOD** · Gastroenterology & Urology
Source: 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