FDA 510(k) Premarket Notification

K800306 FOLEY CATHETER

CLEARED

Feb 28, 1980
16 days to decision

K800306 · Product code: **KOD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k800306/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Feb 12, 1980
Decision date	Feb 28, 1980
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**Abco Dealers, Inc.**
Location	Mchenry, IL, US
510(k) history	127 submissions · 127 cleared · 1976-1991