

K800306 FOLEY CATHETERFeb 28, 1980
16 days to decisionK800306 · Product code: **KOD** · Gastroenterology & UrologySource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800306/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026