

**K883763 ACME GUIDEWIRES FOR PERCUTANEOUS
INSERTION OF CATH**Nov 18, 1988
73 days to decisionK883763 · Product code: **KOD** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k883763/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Sep 6, 1988
Decision date	Nov 18, 1988
Days to decision	73 days
Third-party review	No

APPLICANT

Company	Acme-Monaco Corp.
Location	Plainville, CT, US
Contact	HERBERT M CARTER
510(k) history	5 submissions · 5 cleared · 1988-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k883763/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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