FDA 510(k) Premarket Notification

K883763 ACME GUIDEWIRES FOR PERCUTANEOUS INSERTION OF CATH

CLEARED

Nov 18, 1988
73 days to decision

K883763 · 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