FDA 510(k) Premarket Notification

K855076 ANGIOMED URETER DIATION SETS

CLEARED

Mar 11, 1986
82 days to decision

K855076 · Product code: **KOD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k855076/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Dec 19, 1985
Decision date	Mar 11, 1986
Days to decision	82 days
Third-party review	No

APPLICANT

Company	**Angiomed U.S., Inc.**
Location	Anaheim, CA, US
Contact	RICHARD P MOHR
510(k) history	24 submissions · 24 cleared · 1986-1988