FDA 510(k) Premarket Notification

K864225 ANGIOMED URETERAL D&M SETS/URETEROSCOPY GUID STENT

CLEARED

Jan 30, 1987
94 days to decision

K864225 · Product code: **FAD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k864225/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Oct 28, 1986
Decision date	Jan 30, 1987
Days to decision	94 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