

**K855076 ANGIOMED URETER DIATION SETS**Mar 11, 1986  
82 days to decisionK855076 · Product code: **KOD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k855076/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k855076/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026