

**K864225 ANGIOMED URETERAL D&M SETS/URETEROSCOPY  
GUID STENT**Jan 30, 1987  
94 days to decisionK864225 · Product code: **FAD** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k864225/>**SUBMISSION DETAILS**

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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**

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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/k864225/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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