

**K854906 PERCUTANEOUS NEPHROSTOMY SETS (OTTO)**

Jan 24, 1986

K854906 · Product code: **GBO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k854906/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Nephrostomy, General & Plastic Surgery (GBO)
Date received	Jan 24, 1986
Decision date	Jan 24, 1986
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/k854906/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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