

K854907 PERCUTANEOUS NEPHROSTOMY SETS (GUNTHER)Jan 24, 1986
46 days to decisionK854907 · Product code: **GBO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k854907/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Nephrostomy, General & Plastic Surgery (GBO)
Date received	Dec 9, 1985
Decision date	Jan 24, 1986
Days to decision	46 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k854907/>; Data sourced from FDA 510(k) public records (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. - Generated June 29, 2026