FDA 510(k) Premarket Notification

CLEARED

K854906 PERCUTANEOUS NEPHROSTOMY SETS (OTTO)

Jan 24, 1986

K854906 · Product code: **GBO** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k854906/

SUBMISSION DETAILS

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

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