

K951738 ARGYLE NEO-SERT TRIPLE LUMEN UMBILICAL VESSEL CATHETERSep 29, 1995
168 days to decisionK951738 · Product code: **FOS** · General Hospital
Source: <https://www.510kdatabase.net/k951738/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Umbilical Artery (FOS)
Date received	Apr 14, 1995
Decision date	Sep 29, 1995
Days to decision	168 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sherwood Medical Co.
Location	Mchenry, IL, US
Contact	DENNIS POZZO
510(k) history	191 submissions · 177 cleared · 1976-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k951738/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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