FDA 510(k) Premarket Notification

K951738 ARGYLE NEO-SERT TRIPLE LUMEN UMBILICAL VESSEL CATHETER

CLEARED

Sep 29, 1995
168 days to decision

K951738 · 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