FDA 510(k) Premarket Notification

K842889 ARGYLE CUSTOM TUBING PACK

CLEARED

Aug 27, 1984
35 days to decision

K842889 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k842889/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jul 23, 1984
Decision date	Aug 27, 1984
Days to decision	35 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k842889/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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