FDA 510(k) Premarket Notification

K860742 CAROTID BYPASS SHUNT

CLEARED

May 28, 1986
90 days to decision

K860742 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k860742/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Feb 27, 1986
Decision date	May 28, 1986
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Clincal Instruments Corp.**
Location	Mchenry, IL, US
Contact	MICHAEL L POLANYI
510(k) history	8 submissions · 8 cleared · 1977-1991

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k860742/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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