FDA 510(k) Premarket Notification

K872788 MODIFIED CAROTID SHUNT W/SPRING REINFORCED LUMEN

CLEARED

Sep 25, 1987
73 days to decision

K872788 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k872788/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jul 14, 1987
Decision date	Sep 25, 1987
Days to decision	73 days
Third-party review	No

APPLICANT

Company	**Uresil Corp.**
Location	Mchenry, IL, US
Contact	MICHAEL JARON
510(k) history	45 submissions · 44 cleared · 1981-2001