

K791858 CARDIOPLEGIA ADMIN. SETSOct 26, 1979
37 days to decisionK791858 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k791858/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Sep 19, 1979
Decision date	Oct 26, 1979
Days to decision	37 days
Third-party review	No

APPLICANT

Company	Travenol Laboratories, S.A.
Location	Mchenry, IL, US
Website	https://www.baxter.com
510(k) history	206 submissions · 206 cleared · 1976-1988

Travenol Laboratories, S.A. is a medical device manufacturer based in McHenry, US. The company specializes in infusion, dialysis, and hospital care devices. Travenol Laboratories received FDA 510(k) clearances from total submissions between 1976 and 1988. The company's cleared devices span general hospital and gastroenterology/urology categories, including infusion systems, dialysis equipment, and administration sets. This regulatory record reflects the company's historical focus on critical care and renal therapy technologies. The company is inactive and represents a his...
