FDA 510(k) Premarket Notification

K792202 TROCAR CATHETER/THORACIC CATHETER

CLEARED

Nov 20, 1979
18 days to decision

K792202 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k792202/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Nov 2, 1979
Decision date	Nov 20, 1979
Days to decision	18 days
Third-party review	No

APPLICANT

Company	**Portex, Inc.**
Location	Walker, MI, US
510(k) history	20 submissions · 20 cleared · 1977-2004