FDA 510(k) Premarket Notification

K001961 STOCKERT A242 AND A252 SERIES ARTERIAL FEMORAL CANNULAE

CLEARED

Jan 25, 2001
212 days to decision

K001961 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k001961/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jun 27, 2000
Decision date	Jan 25, 2001
Days to decision	212 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stoeckert Instrumente**
Location	Mchenry, IL, US
Contact	LYNNE LEONARD
510(k) history	13 submissions · 13 cleared · 1982-2001