FDA 510(k) Premarket Notification

K020784 JOSTRA SINGLE STAGE VENOUS RETURN CATHETERS

CLEARED

Jan 9, 2003
304 days to decision

K020784 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k020784/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 11, 2002
Decision date	Jan 9, 2003
Days to decision	304 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Jostra AG**
Location	Newark, DE, US
Contact	KATHLEEN JOHNSON
510(k) history	13 submissions · 13 cleared · 2000-2004