FDA 510(k) Premarket Notification

K915480 B-D ULTRA-VUE PROCEDURE NEEDLE W/ SPINAL TYPE PT.

CLEARED

Jan 7, 1992
32 days to decision

K915480 · Product code: **DWD** · Cardiovascular
Source: https://www.510kdatabase.net/k915480/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suction Control, Intracardiac, Cardiopulmonary Bypass (DWD)
Date received	Dec 6, 1991
Decision date	Jan 7, 1992
Days to decision	32 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Bd Becton Dickinson Vacutainer Systems Preanalytic**
Location	Washington, DC, US
Contact	RUSSELL ARNSBERGER
510(k) history	632 submissions · 625 cleared · 1976-2001