

**K915480 B-D ULTRA-VUE PROCEDURE NEEDLE W/ SPINAL
TYPE PT.**Jan 7, 1992
32 days to decisionK915480 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k915480/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026