

**K915482 B-D ULTRA-VUE PROCEDURE NEEDLE W/  
WESTCOTT TYPE PT**Jan 7, 1992  
32 days to decisionK915482 · Product code: **DWD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k915482/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k915482/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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