

K860136 BARD-PARKER CAROTID SHUNTJun 25, 1986
162 days to decisionK860136 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k860136/>**SUBMISSION DETAILS**

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| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF) |
| Date received | Jan 14, 1986 |
| Decision date | Jun 25, 1986 |
| Days to decision | 162 days |
| Third-party review | No |

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/k860136/>; 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