

K994209 STOCKERT V142 SERIES VENOUS CANNULAE WITH LIGHTHOUSE TIPJun 14, 2000
183 days to decisionK994209 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k994209/>**SUBMISSION DETAILS**

| | |
|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Abbreviated |
| Device classification | Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF) |
| Date received | Dec 14, 1999 |
| Decision date | Jun 14, 2000 |
| Days to decision | 183 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Cobe Cardiovascular, Inc. |
| Location | Arvada, CO, US |
| Contact | LYNNE LEONARD |
| 510(k) history | 43 submissions · 43 cleared · 1992-2005 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k994209/> 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 25, 2026