

K881074 SHILEY FEMORAL VENOUS CANNULAMay 31, 1988
78 days to decisionK881074 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k881074/>**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 | Mar 14, 1988 |
| Decision date | May 31, 1988 |
| Days to decision | 78 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|-------------------------------------------|
| Company | Shiley, Inc. |
| Location | Mchenry, IL, US |
| Contact | RADINE POBUDA |
| 510(k) history | 174 submissions · 174 cleared · 1976-1993 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881074/>; 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 26, 2026