

K802543 RIDGID VENA CAVA CANNULAEJan 16, 1981
93 days to decisionK802543 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k802543/>**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 | Oct 15, 1980 |
| Decision date | Jan 16, 1981 |
| Days to decision | 93 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Texas Medical Products, Inc. |
| Location | Mchenry, IL, US |
| 510(k) history | 30 submissions · 30 cleared · 1980-1989 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k802543/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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