

**K801398 AORTIC AND FEMORAL PERFUSION CANNULAE**Jun 30, 1980  
17 days to decisionK801398 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k801398/>**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         | Jun 13, 1980   |
| Decision date         | Jun 30, 1980   |
| Days to decision      | 17 days  |
| Third-party review    | No   |

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k801398/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026