

K801401 CARDIOPULMONARY BYPASS TUBINGJun 30, 1980
17 days to decisionK801401 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k801401/>**SUBMISSION DETAILS**

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

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

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Device record: <https://www.510kdatabase.net/k801401/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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