

**K801396 SAPHENOUS VEIN CANNULAE**Jun 30, 1980  
17 days to decisionK801396 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k801396/>**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/k801396/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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