

**K802261 PRESSURE GAUGE ACCESSORY KIT/DILATATION**Oct 3, 1980  
17 days to decisionK802261 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k802261/>**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	Sep 16, 1980
Decision date	Oct 3, 1980
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>Stanco Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1979-1981

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k802261/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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