

**K060101 ESTECH AORTIC EASY FLOW CANNULA**May 10, 2006  
118 days to decisionK060101 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k060101/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jan 12, 2006
Decision date	May 10, 2006
Days to decision	118 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Endoscopic Technologies, Inc.</b>
Location	Norcross, GA, US
Contact	CRAIG COOMBS
510(k) history	8 submissions · 8 cleared · 1993-2006

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k060101/> 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 30, 2026