

**K010253 ANGIO FLOW THERMODILUTION CATHETER, MODEL AF-CATH-01**Mar 19, 2002  
417 days to decisionK010253 · Product code: **DYG** · Cardiovascular  
Source: <https://www.510kdatabase.net/k010253/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Flow Directed (DYG)
Date received	Jan 26, 2001
Decision date	Mar 19, 2002
Days to decision	417 days
Third-party review	No
Summary / Statement	Summary
Other names	ANGIOFLOW METER, MODELS HQ100 AND HQ101

**APPLICANT**

---

Company	<b>Transonic Systems, Inc.</b>
Location	Ithaca, NY, US
Contact	MARK S ALSBERGE
510(k) history	16 submissions · 16 cleared · 1988-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010253/> 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 4, 2026