

**K052004 VASCON NEUROPATH GUIDING CATHETER**Jan 19, 2006  
178 days to decisionK052004 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k052004/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jul 25, 2005
Decision date	Jan 19, 2006
Days to decision	178 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Vascon, LLC</b>
Location	Miami, FL, US
Contact	ROBERTA D GOODE
510(k) history	4 submissions · 4 cleared · 2004-2008

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052004/> 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 25, 2026