

**K022000 POURCHEZ RETRO TWIN LUMEN CHRONIC  
HEMODIALYSIS CATHETER WITH SEPARATED TIPS (WITH  
AND WITHOUT SIDE HOLES)**Jul 19, 2002  
30 days to decisionK022000 · Product code: **MSD** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k022000/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Hemodialysis, Implanted (MSD)
Date received	Jun 19, 2002
Decision date	Jul 19, 2002
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Spire Biomedical, Inc.</b>
Location	Bedford, MA, US
Contact	DONALD FICKETT
510(k) history	16 submissions · 12 cleared · 2002-2008

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022000/> 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 26, 2026