

**K011741 CRIT-LINE MONITOR III TQA (CLM TQA)**Jul 23, 2002  
413 days to decisionK011741 · Product code: **FIL** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k011741/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Dialysate Delivery, Single Pass (FIL)
Date received	Jun 5, 2001
Decision date	Jul 23, 2002
Days to decision	413 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Hemametrics</b>
Location	Kaysville, UT, US
Contact	MATTHEW L HAYNIE
510(k) history	1 submissions · 1 cleared · 2002-2002

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k011741/> 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