FDA 510(k) Premarket Notification

K011741 CRIT-LINE MONITOR III TQA (CLM TQA)

CLEARED

Jul 23, 2002
413 days to decision

K011741 · 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	**Hemametrics**
Location	Kaysville, UT, US
Contact	MATTHEW L HAYNIE
510(k) history	1 submissions · 1 cleared · 2002-2002