FDA 510(k) Premarket Notification

K982412 MODIFICATION OF CRIT-LINE MONITOR III (CLM III)

CLEARED

Oct 9, 1998
88 days to decision

K982412 · Product code: **MQS** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k982412/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Hemodialysis, Access Recirculation Monitoring (MQS)
Date received	Jul 13, 1998
Decision date	Oct 9, 1998
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**In-Line Diagnostics Corp.**
Location	Riverdale, UT, US
Contact	MATTHEW L HAYNIE
510(k) history	8 submissions · 8 cleared · 1995-2000