

K982412 MODIFICATION OF CRIT-LINE MONITOR III (CLM III)Oct 9, 1998
88 days to decisionK982412 · Product code: **MQS** · Gastroenterology & UrologySource: <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

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