

K982926 FRESENIUS BLOOD VOLUME MONITORSep 3, 1999
379 days to decisionK982926 · Product code: **KDI** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k982926/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Aug 20, 1998
Decision date	Sep 3, 1999
Days to decision	379 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Fresenius Medical Care North America
Location	Lexington, MA, US
Contact	DONALD A LINCOLN
510(k) history	43 submissions · 43 cleared · 1998-2025

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