

K011269 LIFEMATE MLA MULTI-LINE ADAPTORMar 4, 2002
312 days to decisionK011269 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k011269/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Apr 26, 2001
Decision date	Mar 4, 2002
Days to decision	312 days
Third-party review	No
Summary / Statement	Summary
Other names	LIFEMATE SFR STERILE FLUID RESERVOIR

APPLICANT

Company	Nxstage Medical, Inc.
Location	Tewksburt, MA, US
Contact	KAREN ST. ONGE
510(k) history	51 submissions · 51 cleared · 2001-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k011269/> Data sourced from FDA 510(k) public records (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. - Generated June 1, 2026