

**K843623 UNILYTE A**Nov 16, 1984  
87 days to decisionK843623 · Product code: **FKQ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k843623/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Dialysate Delivery, Central Multiple Patient (FKQ)
Date received	Aug 21, 1984
Decision date	Nov 16, 1984
Days to decision	87 days
Third-party review	No

**APPLICANT**

---

Company	<b>Medical Enterprises, Ltd.</b>
Location	Jackson, MS, US
Contact	HOLLAND
510(k) history	1 submissions · 1 cleared · 1984-1984

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k843623/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 3, 2026