FDA 510(k) Premarket Notification

K843623 UNILYTE A

CLEARED

Nov 16, 1984
87 days to decision

K843623 · Product code: **FKQ** · Gastroenterology & Urology
Source: 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	**Medical Enterprises, Ltd.**
Location	Jackson, MS, US
Contact	HOLLAND
510(k) history	1 submissions · 1 cleared · 1984-1984