FDA 510(k) Premarket Notification

K970799 ALEXANDER MANUFACTURING RECHARGEABLE BATTERY (M1/T)

CLEARED

May 14, 1997
71 days to decision

K970799 · Product code: **LIF** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k970799/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer Reprocessing System (LIF)
Date received	Mar 4, 1997
Decision date	May 14, 1997
Days to decision	71 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Alexander Mfg. Co.**
Location	Mason City, IA, US
Contact	STACEY HIPPEN
510(k) history	39 submissions · 37 cleared · 1997-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k970799/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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