FDA 510(k) Premarket Notification

K971865 ALEXANDER MANUFACTURING CO. RECHARGEABLE BATTERY PART NUMBER M12/500-3P

CLEARED

Jul 3, 1997
44 days to decision

K971865 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k971865/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	May 20, 1997
Decision date	Jul 3, 1997
Days to decision	44 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k971865/ 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 May 25, 2026