

K853843 IV MATEJan 9, 1986
115 days to decisionK853843 · Product code: **FLN** · General Hospital
Source: <https://www.510kdatabase.net/k853843/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Electric For Gravity Flow Infusion Systems (FLN)
Date received	Sep 16, 1985
Decision date	Jan 9, 1986
Days to decision	115 days
Third-party review	No

APPLICANT

Company	Medtech Intl., Inc.
Location	Mchenry, IL, US
Contact	JONATHAN ROSEN
510(k) history	6 submissions · 6 cleared · 1978-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k853843/>; 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 July 2, 2026