

K854058 PK-AC POWERED PATIENT MANUPULATING DEVICEFeb 25, 1986
145 days to decisionK854058 · Product code: **FNG** · General Hospital
Source: <https://www.510kdatabase.net/k854058/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lift, Patient, Ac-powered (FNG)
Date received	Oct 3, 1985
Decision date	Feb 25, 1986
Days to decision	145 days
Third-party review	No

APPLICANT

Company	Mannesmann Demag Corp.
Location	Solon, OH, US
Contact	DAVID R MARSHALL
510(k) history	1 submissions · 1 cleared · 1986-1986

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k854058/>; 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 9, 2026