FDA 510(k) Premarket Notification

K854058 PK-AC POWERED PATIENT MANUPULATING DEVICE

CLEARED

Feb 25, 1986
145 days to decision

K854058 · 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