FDA 510(k) Premarket Notification

K813356 DEVICE FOR REPOSITIONING A BET PATIENT

CLEARED

Feb 10, 1982
75 days to decision

K813356 · Product code: **FNJ** · General Hospital
Source: https://www.510kdatabase.net/k813356/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Manual (FNJ)
Date received	Nov 27, 1981
Decision date	Feb 10, 1982
Days to decision	75 days
Third-party review	No

APPLICANT

Company	**Ali-Oop Co., Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982