

K813356 DEVICE FOR REPOSITIONING A BET PATIENTFeb 10, 1982
75 days to decisionK813356 · Product code: **FNJ** · General HospitalSource: <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

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Device record: <https://www.510kdatabase.net/k813356/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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