FDA 510(k) Premarket Notification

K801517 BED ROCKING DEVICE

CLEARED

Oct 10, 1980
101 days to decision

K801517 · Product code: **FNJ** · General Hospital
Source: https://www.510kdatabase.net/k801517/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Manual (FNJ)
Date received	Jul 1, 1980
Decision date	Oct 10, 1980
Days to decision	101 days
Third-party review	No

APPLICANT

Company	**Wesler**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1980-1980