

K801517 BED ROCKING DEVICEOct 10, 1980
101 days to decisionK801517 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k801517/>; 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 10, 2026