

K823926 BED-CHECK: II-IFACApr 18, 1983
110 days to decisionK823926 · Product code: **KMI** · General Hospital
Source: <https://www.510kdatabase.net/k823926/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Bed Patient (KMI)
Date received	Dec 29, 1982
Decision date	Apr 18, 1983
Days to decision	110 days
Third-party review	No

APPLICANT

Company	Bed-Check Corp.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k823926/> 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