

K911079 AMB-ALERTSep 13, 1991
186 days to decisionK911079 · Product code: **KMI** · General HospitalSource: <https://www.510kdatabase.net/k911079/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Bed Patient (KMI)
Date received	Mar 11, 1991
Decision date	Sep 13, 1991
Days to decision	186 days
Third-party review	No

APPLICANT

Company	Medway
Location	Franklin, TN, US
Contact	JAMES J CALLAWAY
510(k) history	5 submissions · 5 cleared · 1991-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k911079/>, 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 May 26, 2026