

**K872358 STA-PUT**Aug 6, 1987  
49 days to decisionK872358 · Product code: **KMI** · General Hospital  
Source: <https://www.510kdatabase.net/k872358/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Bed Patient (KMI)
Date received	Jun 18, 1987
Decision date	Aug 6, 1987
Days to decision	49 days
Third-party review	No

**APPLICANT**

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Company	<b>Design Technologies, Inc.</b>
Location	Rahway, NJ, US
Contact	LOUIS G ACRI
510(k) history	4 submissions · 4 cleared · 1986-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872358/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 27, 2026