FDA 510(k) Premarket Notification

K943481 INFORMER PLUS

CLEARED

Nov 2, 1994
106 days to decision

K943481 · Product code: **KMI** · General Hospital
Source: https://www.510kdatabase.net/k943481/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Bed Patient (KMI)
Date received	Jul 19, 1994
Decision date	Nov 2, 1994
Days to decision	106 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Micro-Tech Medical, Inc.**
Location	East Hartford, CT, US
Contact	KEITH CHARLES
510(k) history	1 submissions · 1 cleared · 1994-1994

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k943481/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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