

K864190 E-Z HOLD - #4060Jan 14, 1987
79 days to decisionK864190 · Product code: **KMK** · General Hospital
Source: <https://www.510kdatabase.net/k864190/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Oct 27, 1986
Decision date	Jan 14, 1987
Days to decision	79 days
Third-party review	No

APPLICANT

Company	John F. Greer Co., Inc.
Location	Oakland, CA, US
Contact	LELAND H GREER
510(k) history	5 submissions · 5 cleared · 1986-1987

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k864190/>; 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 3, 2026