FDA 510(k) Premarket Notification

K864190 E-Z HOLD - #4060

CLEARED

Jan 14, 1987
79 days to decision

K864190 · 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