

K840683 FPP IV SECUREMar 6, 1984
20 days to decisionK840683 · Product code: **KMK** · General Hospital
Source: <https://www.510kdatabase.net/k840683/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Feb 15, 1984
Decision date	Mar 6, 1984
Days to decision	20 days
Third-party review	No

APPLICANT

Company	Fpp, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1984-1984

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Device record: <https://www.510kdatabase.net/k840683/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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