FDA 510(k) Premarket Notification

K840683 FPP IV SECURE

CLEARED

Mar 6, 1984
20 days to decision

K840683 · 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