FDA 510(k) Premarket Notification

K810721 PROCESS CHANGE

CLEARED

Apr 23, 1981
37 days to decision

K810721 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k810721/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 17, 1981
Decision date	Apr 23, 1981
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Medlon, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981