FDA 510(k) Premarket Notification

K781785 I.V. SET WITH IN-LINE FILTER

CLEARED

Dec 7, 1978
50 days to decision

K781785 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k781785/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Oct 18, 1978
Decision date	Dec 7, 1978
Days to decision	50 days
Third-party review	No

APPLICANT

Company	**Cutler Laboratories, Inc.**
Location	MD, US
510(k) history	2 submissions · 2 cleared · 1978-1978