FDA 510(k) Premarket Notification

K833659 IV EXTENSION SET

CLEARED

Nov 29, 1983
42 days to decision

K833659 · Product code: **FPK** · General Hospital
Source: https://www.510kdatabase.net/k833659/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubing, Fluid Delivery (FPK)
Date received	Oct 18, 1983
Decision date	Nov 29, 1983
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**Dna Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1982-1983