FDA 510(k) Premarket Notification

K812852 SET FOR ASEPTIC DECANTING OF A FLEX I.V

CLEARED

Nov 2, 1981
20 days to decision

K812852 · Product code: **KPE** · General Hospital
Source: https://www.510kdatabase.net/k812852/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Oct 13, 1981
Decision date	Nov 2, 1981
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Dip, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1981-1981