FDA 510(k) Premarket Notification

K821624 LOW VOLUME INFUSTION SET

CLEARED

Jun 22, 1982
20 days to decision

K821624 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k821624/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jun 2, 1982
Decision date	Jun 22, 1982
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Lifemed of California**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1982-2001