FDA 510(k) Premarket Notification

K820200 IV ALERT SYSTEM

CLEARED

Mar 18, 1982
51 days to decision

K820200 · Product code: **KPE** · General Hospital
Source: https://www.510kdatabase.net/k820200/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Jan 26, 1982
Decision date	Mar 18, 1982
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Allied Medical Technologies, Inc.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1982-1982