FDA 510(k) Premarket Notification

K802966 I.V. ADDITIVE CAP

CLEARED

Jan 7, 1981
47 days to decision

K802966 · Product code: **KPE** · General Hospital
Source: https://www.510kdatabase.net/k802966/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Nov 21, 1980
Decision date	Jan 7, 1981
Days to decision	47 days
Third-party review	No

APPLICANT

Company	**Baxa Corp., Sub. of Cook Group, Inc.**
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1978-1991