FDA 510(k) Premarket Notification

K830868 COMBISET OR INFUSION SET

CLEARED

Apr 14, 1983
27 days to decision

K830868 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k830868/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 18, 1983
Decision date	Apr 14, 1983
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Pan-Tek Intl., Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983