FDA 510(k) Premarket Notification

K832329 STERILE IV SET 4 SIZES

CLEARED

Aug 24, 1983
40 days to decision

K832329 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k832329/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 15, 1983
Decision date	Aug 24, 1983
Days to decision	40 days
Third-party review	No

APPLICANT

Company	**Abco Dealers, Inc.**
Location	Mchenry, IL, US
510(k) history	127 submissions · 127 cleared · 1976-1991