FDA 510(k) Premarket Notification

K900909 IN-LINE VIAL CONNECTION SITE

CLEARED

May 18, 1990
80 days to decision

K900909 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k900909/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 27, 1990
Decision date	May 18, 1990
Days to decision	80 days
Third-party review	No

APPLICANT

Company	**Baxter Healthcare Corp**
Location	Mchenry, IL, US
Contact	PATRICIA S BARSANTI
510(k) history	505 submissions · 496 cleared · 1977-2019