FDA 510(k) Premarket Notification

K933848 PRESPO 7, SELF-SUFFICIENT DRIP DEVICE

CLEARED

Mar 3, 1995
574 days to decision

K933848 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k933848/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 6, 1993
Decision date	Mar 3, 1995
Days to decision	574 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Bonding Corp.**
Location	Coraopolis, PA, US
Contact	JEAN-PIERRE LA RUDULIER
510(k) history	1 submissions · 1 cleared · 1995-1995