FDA 510(k) Premarket Notification

K013714 CODAN US I.V. ADMINISTRATION SET, MODEL B400 SP

CLEARED

Mar 28, 2002
140 days to decision

K013714 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k013714/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 8, 2001
Decision date	Mar 28, 2002
Days to decision	140 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Codan US Corp.**
Location	Irvine, CA, US
Contact	MARILYN R POURAZAR
510(k) history	7 submissions · 7 cleared · 1999-2012

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k013714/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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