FDA 510(k) Premarket Notification

K001872 I.V. SET-LUER LOCK WITH LATEX FREE INJECTION SITE, MODEL B411

CLEARED

Sep 7, 2000
79 days to decision

K001872 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k001872/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jun 20, 2000
Decision date	Sep 7, 2000
Days to decision	79 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Codan US Corp.**
Location	Irvine, CA, US
Contact	RICHARD A BASLER
510(k) history	7 submissions · 7 cleared · 1999-2012

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k001872/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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