FDA 510(k) Premarket Notification

CLEARED

K042800 LIFECARE PCA INFUSION SYSTEM WITH HOSPIRA MEDNET SOFTWARE

Oct 18, 2004
10 days to decision

K042800 · Product code: **MEA** · General Hospital
Source: https://www.510kdatabase.net/k042800/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion, Pca (MEA)
Date received	Oct 8, 2004
Decision date	Oct 18, 2004
Days to decision	10 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Hospira, Inc.**
Location	Lake Forest, IL, US
Contact	THOMAS KOZMA
Website	http://www.hospira.com
510(k) history	45 submissions · 44 cleared · 2004-2017

Hospira, Inc. was an American global pharmaceutical and medical device company headquartered in Lake Forest, Illinois. The company specialized in generic injectable pharmaceuticals and integrated infusion therapy systems for hospitals and alternate care settings. Hospira maintains an FDA 510(k) regulatory record of cleared devices from total submissions between 2004 and 2017. The company's primary focus was General Hospital devices, which comprised the majority of its submissions. Notable cleared products include the Plum 360 Infusion System, extension sets, administratio...