

**K143015 Hospira Primary Sets**Feb 25, 2015  
128 days to decisionK143015 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k143015/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Oct 20, 2014
Decision date	Feb 25, 2015
Days to decision	128 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hospira, Inc.</b>
Location	Lake Forest, IL, US
Contact	Anju Kurian
Website	<a href="http://www.hospira.com">http://www.hospira.com</a>
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...

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