

**K061585 OPTICATH CENTRAL VENOUS OXIMETRY CATHETER**Oct 5, 2006  
120 days to decisionK061585 · Product code: **DQE** · Cardiovascular  
Source: <https://www.510kdatabase.net/k061585/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Oximeter, Fiber-optic (DQE)
Date received	Jun 7, 2006
Decision date	Oct 5, 2006
Days to decision	120 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hospira, Inc.</b>
Location	Lake Forest, IL, US
Contact	NICOHL WILDING
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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