

**K131085 CENTRAL VENOUS PRESSURE SYSTEM - MESPERE VENUS 1000**

Aug 22, 2013  
126 days to decision

K131085 · Product code: **PFA** · Cardiovascular  
Source: <https://www.510kdatabase.net/k131085/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Non-invasive Central Venous Manometer (PFA)
Date received	Apr 18, 2013
Decision date	Aug 22, 2013
Days to decision	126 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Mespere Lifesciences, Inc.</b>
Location	Waterloo, On, CA
Contact	HELEN TAN
510(k) history	3 submissions · 3 cleared · 2013-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k131085/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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