

**K991886 PULSION PULSIOCATH, PULSION PCCO  
MONITORING KIT, PULSION IN-LINE INJECTATE SENSOR**May 11, 2000  
344 days to decisionK991886 · Product code: **KRB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k991886/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Thermodilution (KRB)
Date received	Jun 2, 1999
Decision date	May 11, 2000
Days to decision	344 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pulsion Medical Systems, Inc.</b>
Location	Cornelius, NC, US
Contact	ULF BORG
510(k) history	3 submissions · 3 cleared · 2000-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k991886/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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