

**K090671 PULSE OXIMETER, MODELS CMS50E, CMS50F, CMS60C, CMS60D**Jun 11, 2009  
90 days to decisionK090671 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k090671/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Mar 13, 2009
Decision date	Jun 11, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Contec Medical System Co., Ltd.</b>
Location	Zhong Shan, Shanghai, CN
Contact	Diana Hong
510(k) history	12 submissions · 12 cleared · 2008-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090671/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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