

**K192343 Patient Monitor**Dec 31, 2019  
125 days to decisionK192343 · Product code: **DQA** · Cardiovascular  
Source: <https://www.510kdatabase.net/k192343/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Aug 28, 2019
Decision date	Dec 31, 2019
Days to decision	125 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen Creative Industry Co., Ltd.</b>
Location	Flintville, TN, US
Contact	Jia Wang
510(k) history	11 submissions · 11 cleared · 2007-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k192343/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 27, 2026