

**K171241 Pulse Oximeter**Mar 27, 2018  
334 days to decisionK171241 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k171241/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Apr 27, 2017
Decision date	Mar 27, 2018
Days to decision	334 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hunan Accurate Bio-Medical Technology Co., Ltd.</b>
Location	Changsha, Hunan, CN
Contact	Zhou Le
510(k) history	5 submissions · 5 cleared · 2014-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k171241/> 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 26, 2026