

**K210305 Pulse Oximeter**Dec 22, 2021  
322 days to decisionK210305 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k210305/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Feb 3, 2021
Decision date	Dec 22, 2021
Days to decision	322 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dongguan Lingxin Technologies Co., Ltd.</b>
Location	Dongguan, CN
Contact	Cony Tang
510(k) history	1 submissions · 1 cleared · 2021-2021

**REGULATORY CONSULTANT**

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Consulting firm	<b>Shenzhen Reanny Medical Devices Mangement Consulting., Ltd.</b>
Contact	Reanny Wang

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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