

**K213984 Oximeter (Model: JZK-301,JZK-303,JZK-305,JZK-307)**Dec 16, 2022  
361 days to decisionK213984 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k213984/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Dec 20, 2021
Decision date	Dec 16, 2022
Days to decision	361 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen Zhengkang Technology Co., Ltd.</b>
Location	Shenzhen City, CN
Contact	Huayong Yang
510(k) history	3 submissions · 3 cleared · 2020-2022

**REGULATORY CONSULTANT**

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Consulting firm	<b>Feiyong Drug &amp; Medical Consulting Technical Service Group</b>
Contact	Becky Chen

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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