

**K100225 PULSE OXIMETER MODEL P10**Feb 9, 2011  
379 days to decisionK100225 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k100225/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jan 26, 2010
Decision date	Feb 9, 2011
Days to decision	379 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Mediana Co., Ltd.</b>
Location	Flintville, TN, US
Contact	CHARLIE MACK
510(k) history	10 submissions · 10 cleared · 2005-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k100225/> 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 June 18, 2026