

**K101577 MODIFICATION TO:FINGERTIP PULSE OXIMETER,
MODEL MD300C SERIES**Dec 2, 2010
178 days to decisionK101577 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k101577/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Oximeter (DQA)
Date received	Jun 7, 2010
Decision date	Dec 2, 2010
Days to decision	178 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Beijing Choice Electronic Technology Co., Ltd.
Location	Shanghai, CN
Contact	Diana Hong
510(k) history	29 submissions · 29 cleared · 2007-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k101577/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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