

**K100203 HANDHELD PULSE OXIMETER MODEL MD300A/
MD300K2/ MD300M**May 17, 2010
112 days to decisionK100203 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k100203/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jan 25, 2010
Decision date	May 17, 2010
Days to decision	112 days
Third-party review	No
Summary / Statement	Statement

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/k100203/> Data sourced from FDA 510(k) public records (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. - Generated May 17, 2026