

K160268 Fingertip Pulse Oximeter MD300CG11/MD300CG51Oct 14, 2016
255 days to decisionK160268 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k160268/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Feb 2, 2016
Decision date	Oct 14, 2016
Days to decision	255 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Beijing Choice Electronic Technology Co., Ltd.
Location	Beijing, CN
Contact	LEI CHEN
510(k) history	14 submissions · 14 cleared · 2016-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k160268/> 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 26, 2026