

K230172 Pulse OximeterAug 12, 2023
204 days to decisionK230172 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k230172/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jan 20, 2023
Decision date	Aug 12, 2023
Days to decision	204 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k230172/> 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 13, 2026