

K240092 Pulse Oximeter (YM101, YM102, YM103, YM104, YM201, YM301, YM202, YM302, YM111, YM112, YM113, YM114, YM211, YM212, YM314, YM601, YM602, YM401, YM402, YM403, YM501, YM502, YM503 and YM504)Oct 25, 2024
287 days to decisionK240092 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k240092/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jan 12, 2024
Decision date	Oct 25, 2024
Days to decision	287 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Shenzhen Yimi Life-Technology Co., Ltd.
Location	Shenzhen, CN
Contact	Peng Shande
510(k) history	2 submissions · 2 cleared · 2019-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k240092/> 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