

K220899 Oxehealth Vital SignsApr 29, 2022
32 days to decisionK220899 · Product code: **QME** · Cardiovascular
Source: <https://www.510kdatabase.net/k220899/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Software For Optical Camera-based Measurement Of Pulse Rate, Heart Rate, Breathing Rate, And/or Respiratory Rate (QME)
Date received	Mar 28, 2022
Decision date	Apr 29, 2022
Days to decision	32 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Oxehealth Limited
Location	Abingdon, GB
Contact	Iain Charlton
510(k) history	6 submissions · 5 cleared · 2021-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k220899/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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