

**K240890 PanopticAI Vital Signs**Dec 23, 2024  
266 days to decisionK240890 · Product code: **QME** · Cardiovascular  
Source: <https://www.510kdatabase.net/k240890/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Software For Optical Camera-based Measurement Of Pulse Rate, Heart Rate, Breathing Rate, And/or Respiratory Rate (QME)
Date received	Apr 1, 2024
Decision date	Dec 23, 2024
Days to decision	266 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>PanopticAI technologies Limited</b>
Location	Hong Kong, HK
Contact	Tsz Tai Chan
510(k) history	1 submissions · 1 cleared · 2024-2024

**REGULATORY CONSULTANT**

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Consulting firm	<b>Vee Care (Asia) Limited</b>
Contact	Jing Chen

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k240890/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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