

K231733 Neteera 130H-Plus Vital Sign Monitoring SensorFeb 9, 2024
241 days to decisionK231733 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k231733/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jun 13, 2023
Decision date	Feb 9, 2024
Days to decision	241 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Neteera Technologies , Ltd.
Location	Jerusalem, IL
Contact	Yossi Muncher
510(k) history	2 submissions · 2 cleared · 2022-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k231733/> 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