FDA 510(k) Premarket Notification

K231733 Neteera 130H-Plus Vital Sign Monitoring Sensor

CLEARED

Feb 9, 2024
241 days to decision

K231733 · Product code: **DRT** · Cardiovascular
Source: 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