

K211754 Vital Signs MonitorNov 15, 2021
161 days to decisionK211754 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k211754/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jun 7, 2021
Decision date	Nov 15, 2021
Days to decision	161 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Beijing Choice Electronic Technology Co., Ltd.
Location	Beijing, CN
Contact	Haiying Zhao
510(k) history	14 submissions · 14 cleared · 2016-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k211754/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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