

**K223170 Electronic Blood Pressure Monitor, model:
HTD6602US**Dec 5, 2023
420 days to decisionK223170 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k223170/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Oct 11, 2022
Decision date	Dec 5, 2023
Days to decision	420 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Hetaida Technology Co., Ltd.
Location	Dongguan, CN
Contact	Tom Chen
510(k) history	8 submissions · 8 cleared · 2018-2025

REGULATORY CONSULTANT

Consulting firm	Qimmiq Medical Consulting Service Co., Ltd.
Contact	You Yijie

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

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