

**K240729 Electronic Blood Pressure Monitor (Model:  
PG-800B30, PG-800B38, PG-800B39, PG-800B45, PG-800B46,  
PG-800B47, PG-800B48, PG-800B53, PG-800B55, PG-800B56,  
PG-800B57, PG-800B58, PG-800B61)**Jun 14, 2024  
88 days to decisionK240729 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k240729/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Mar 18, 2024
Decision date	Jun 14, 2024
Days to decision	88 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen Pango Medical Electronics Co., Ltd.</b>
Location	Shenzhen, CN
Contact	Yang Xiaoyun
510(k) history	3 submissions · 3 cleared · 2022-2024

**REGULATORY CONSULTANT**

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Consulting firm	<b>Guangzhou GLOMED Biological Technology Co., Ltd.</b>
Contact	Cassie Lee

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/k240729/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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