

**K193348 Blood Pressure Monitor**Jun 25, 2020  
205 days to decisionK193348 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k193348/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Dec 3, 2019
Decision date	Jun 25, 2020
Days to decision	205 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen Viatom Technology Co., Ltd.</b>
Location	Shen Zhen, CN
Contact	Zhou Saixin
510(k) history	5 submissions · 5 cleared · 2019-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k193348/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026