

K202934 Blood Pressure MonitorJan 26, 2021
119 days to decisionK202934 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k202934/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Sep 29, 2020
Decision date	Jan 26, 2021
Days to decision	119 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Bioland Technology, Ltd.
Location	North Point, HK
Contact	Yiqing Feng
510(k) history	8 submissions · 8 cleared · 2009-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k202934/> Data sourced from FDA 510(k) public records (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. - Generated May 27, 2026