FDA 510(k) Premarket Notification

K202934 Blood Pressure Monitor

CLEARED

Jan 26, 2021
119 days to decision

K202934 · Product code: **DXN** · Cardiovascular
Source: 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