

K191673 Upper Arm Blood Pressure MonitorNov 27, 2019
156 days to decisionK191673 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k191673/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jun 24, 2019
Decision date	Nov 27, 2019
Days to decision	156 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Famidoc Technology Company Limited
Location	Dongguan, CN
Contact	Leon Cao
510(k) history	3 submissions · 3 cleared · 2019-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k191673/> 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