

K090772 FULLY AUTOMATIC ELECTRONIC BLOOD PRESSURE MONITOR, MODEL KD-5903, KD-5909Jun 4, 2009
73 days to decisionK090772 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k090772/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Mar 23, 2009
Decision date	Jun 4, 2009
Days to decision	73 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Andon Health Co, Ltd.
Location	Tiajin, CN
Contact	LIU YI
510(k) history	92 submissions · 92 cleared · 2008-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k090772/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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