

**K091739 SEMI AUTOMATIC ELECTRONIC BLOOD PRESSURE MONITOR, MODELS KD-3917, KD 3909**Aug 18, 2009  
68 days to decisionK091739 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k091739/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jun 11, 2009
Decision date	Aug 18, 2009
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091739/> 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 June 22, 2026