

K123839 WA SERIES: WA100Jan 11, 2013
29 days to decisionK123839 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k123839/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Dec 13, 2012
Decision date	Jan 11, 2013
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary
Other names	WA200; WD100; WE100, WF100; WB200

APPLICANT

Company	Ningbo Diaier Electronic Co., Ltd.
Location	Yuyao City, Zhejiang, CN
Contact	LAO XIKUN
510(k) history	3 submissions · 3 cleared · 2009-2013

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k123839/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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