FDA 510(k) Premarket Notification

K123839 WA SERIES: WA100

CLEARED

Jan 11, 2013
29 days to decision

K123839 · 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