

K083395 SEMI-FULLY AUTOMATIC ELECTRONIC BLOOD PRESSURE MONITOR, MODELS: KD-322,KD-622,KD-575,KD-525E,KD-593,KD-595,KD-596,KD-598

Feb 27, 2009
102 days to decision

K083395 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k083395/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Nov 17, 2008
Decision date	Feb 27, 2009
Days to decision	102 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.net

Device record: <https://www.510kdatabase.net/k083395/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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