

**K122259 BLOOD PRESSURE MONITOR WITH STETHOSCOPE**Sep 24, 2012  
59 days to decisionK122259 · Product code: **DXQ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k122259/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	Jul 27, 2012
Decision date	Sep 24, 2012
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hangzhou Reli-On Co., Ltd.</b>
Location	Echo, OR, US
Contact	CHARILE MACK
510(k) history	1 submissions · 1 cleared · 2012-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122259/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 13, 2026