

**K131569 PG-800A SERIES**Jun 14, 2013  
15 days to decisionK131569 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k131569/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	May 30, 2013
Decision date	Jun 14, 2013
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen Pango Electronic Co., Ltd.</b>
Location	Shanghai, CN
Contact	Diana Hong
510(k) history	9 submissions · 9 cleared · 2010-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131569/> 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 May 26, 2026