

**K152770 ProBP 2400 Digital Blood Pressure Device**Oct 21, 2015  
26 days to decisionK152770 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k152770/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Sep 25, 2015
Decision date	Oct 21, 2015
Days to decision	26 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Microlife Intellectual Property GmbH</b>
Location	Great Neck, NY, US
Contact	GERHARD FRICK
510(k) history	54 submissions · 54 cleared · 2003-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k152770/> 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 25, 2026