

**K063814 BP-200 PLUS**Mar 7, 2007  
75 days to decisionK063814 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k063814/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Dec 22, 2006
Decision date	Mar 7, 2007
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Schiller AG</b>
Location	Baar, CH
Contact	RETO KUETTEL
510(k) history	16 submissions · 16 cleared · 1985-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063814/> 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 27, 2026