

**K972379 PRECISA N**Sep 24, 1997  
90 days to decisionK972379 · Product code: **DXQ** · CardiovascularSource: <https://www.510kdatabase.net/k972379/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	Jun 26, 1997
Decision date	Sep 24, 1997
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Rudolf Riester GmbH &amp; Co. KG</b>
Location	D-72417 Jungingen, DE
Contact	PATRICIA RIESTER-FREUDENMANN
510(k) history	30 submissions · 26 cleared · 1993-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k972379/> 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