

K051368 BR 102 PLUS BLOOD PRESSURE HOLTER SYSTEMAug 9, 2005
76 days to decisionK051368 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k051368/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	May 25, 2005
Decision date	Aug 9, 2005
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Schiller AG
Location	Baar, CH
Contact	MARKUS BUETLER
510(k) history	16 submissions · 16 cleared · 1985-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k051368/> Data sourced from FDA 510(k) public records (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. - Generated May 27, 2026