

**K221056 ARGUS PB-3000**Jul 6, 2023  
451 days to decisionK221056 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k221056/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Apr 11, 2022
Decision date	Jul 6, 2023
Days to decision	451 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Schiller AG</b>
Location	Baar, CH
Contact	Iru Rothfuchs
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/k221056/> 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 13, 2026