

**K053207 ARGUS, MODEL LCM**May 19, 2006  
184 days to decisionK053207 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k053207/>**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	Nov 16, 2005
Decision date	May 19, 2006
Days to decision	184 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Schiller AG</b>
Location	Baar, CH
Contact	MARKUS BUETLER
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/k053207/> 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