

K101619 WELCH ALLYN 1500 PATIENT MONITORNov 12, 2010
156 days to decisionK101619 · Product code: **MHX** · Cardiovascular
Source: <https://www.510kdatabase.net/k101619/>**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 | Jun 9, 2010 |
| Decision date | Nov 12, 2010 |
| Days to decision | 156 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k101619/> 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