

K950264 MAGLIFESep 27, 1996
613 days to decisionK950264 · Product code: **MHX** · CardiovascularSource: <https://www.510kdatabase.net/k950264/>**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 | Jan 23, 1995 |
| Decision date | Sep 27, 1996 |
| Days to decision | 613 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Odam |
| Location | Mchenry, IL, US |
| Contact | STEVEN CLARKE |
| 510(k) history | 6 submissions · 6 cleared · 1983-1998 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k950264/> 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 July 4, 2026