

K881772 SENSOCOR(TM) R-TEST 240 XLNov 9, 1988
197 days to decisionK881772 · Product code: **DXH** · CardiovascularSource: <https://www.510kdatabase.net/k881772/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Transmitters And Receivers, Electrocardiograph, Telephone (DXH) |
| Date received | Apr 26, 1988 |
| Decision date | Nov 9, 1988 |
| Days to decision | 197 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Sensormedix, Inc. |
| Location | Fort Lauderdale, FL, US |
| Contact | ROBERT P SILVER |
| 510(k) history | 2 submissions · 2 cleared · 1988-1989 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881772/>, 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 June 29, 2026