

K780432 ARRHYTHMIA PROGRAM MODULEMay 26, 1978
70 days to decisionK780432 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k780432/>**SUBMISSION DETAILS**

| | |
|-----------------------|--------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Detector And Alarm, Arrhythmia (DSI) |
| Date received | Mar 17, 1978 |
| Decision date | May 26, 1978 |
| Days to decision | 70 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Midwest Analog & Digital, Inc. |
| Location | Mchenry, IL, US |
| 510(k) history | 3 submissions · 3 cleared · 1978-1985 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k780432/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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