

K830013 EPI CARDIA SYS. - AMBULATORY MONITORFeb 28, 1983
56 days to decisionK830013 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k830013/>**SUBMISSION DETAILS**

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|-----------------------|--------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Detector And Alarm, Arrhythmia (DSI) |
| Date received | Jan 3, 1983 |
| Decision date | Feb 28, 1983 |
| Days to decision | 56 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Medicomp, Inc. |
| Location | Mchenry, IL, US |
| 510(k) history | 23 submissions · 23 cleared · 1983-2018 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k830013/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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