

**K902478 LIFEPAK 300 AUTOMATIC ADVISORY
DEFIBRILLATOR**Oct 31, 1990
149 days to decisionK902478 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k902478/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Dc-defibrillator, Low-energy, (including Paddles) (LDD) |
| Date received | Jun 4, 1990 |
| Decision date | Oct 31, 1990 |
| Days to decision | 149 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Physio-Control Corp. |
| Location | Mchenry, IL, US |
| Contact | SHERRI L POCOCK |
| 510(k) history | 80 submissions · 78 cleared · 1976-1999 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k902478/>; 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