

K900519 CLASSIC ECG/STIMULATING NO. 1700 SERIESMar 19, 1990
45 days to decisionK900519 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k900519/>**SUBMISSION DETAILS**

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|-----------------------|-------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrode, Electrocardiograph (DRX) |
| Date received | Feb 2, 1990 |
| Decision date | Mar 19, 1990 |
| Days to decision | 45 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Classic Medical Products, Inc. |
| Location | Muskego, WI, US |
| Contact | ROBERT A MACUR |
| 510(k) history | 2 submissions · 2 cleared · 1986-1990 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900519/>; 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