

K931794 MARINR SERIES EP DIAGNOSTIC CATHETERSJun 6, 1994
420 days to decisionK931794 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k931794/>**SUBMISSION DETAILS**

| | |
|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF) |
| Date received | Apr 12, 1993 |
| Decision date | Jun 6, 1994 |
| Days to decision | 420 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Medtronic Cardiorhythm |
| Location | San Jose, CA, US |
| Contact | CRAIG J COOMBS |
| 510(k) history | 5 submissions · 5 cleared · 1994-1998 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k931794/> 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 14, 2026