

**K033050 EN CIRCLR AL ADJUSTABLE LOOP MAPPING
CATHETER MODELS, 1045AL1, 1045AL2, 1060AL1, 1060AL2**Mar 26, 2004
179 days to decisionK033050 · Product code: DRF · Cardiovascular
Source: <https://www.510kdatabase.net/k033050/>**SUBMISSION DETAILS**

| | |
|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF) |
| Date received | Sep 29, 2003 |
| Decision date | Mar 26, 2004 |
| Days to decision | 179 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Medtronic Vascular |
| Location | Walker, MI, US |
| Contact | MAC MCKEEN |
| 510(k) history | 475 submissions · 453 cleared · 1977-2023 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k033050/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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