FDA 510(k) Premarket Notification

# K901555 MODEL 1730 NEOTRODE WITH ATTACHED LEADWIRE

**CLEARED**

Jul 18, 1990
106 days to decision

K901555 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k901555/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrode, Electrocardiograph (DRX) |
| Date received | Apr 3, 1990 |
| Decision date | Jul 18, 1990 |
| Days to decision | 106 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Medtronic Vascular** |
| Location | Walker, MI, US |
| Contact | JANICE M PEVIDE |
| 510(k) history | 475 submissions · 453 cleared · 1977-2023 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k901555/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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