FDA 510(k) Premarket Notification

# K833619 CONDUCTANCE METER

**CLEARED**

Jan 27, 1984
105 days to decision

K833619 · Product code: **KRC** · Cardiovascular
Source: https://www.510kdatabase.net/k833619/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Tester, Electrode, Surface, Electrocardiographic (KRC) |
| Date received | Oct 14, 1983 |
| Decision date | Jan 27, 1984 |
| Days to decision | 105 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Circadian, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 13 submissions · 13 cleared · 1980-1992 |