FDA 510(k) Premarket Notification

# K830655 QUIK-SILVER RESTING ECG ELECTRODE

**CLEARED**

Jun 15, 1983
105 days to decision

K830655 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k830655/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrode, Electrocardiograph (DRX) |
| Date received | Mar 2, 1983 |
| Decision date | Jun 15, 1983 |
| Days to decision | 105 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Quinton, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 164 submissions · 160 cleared · 1976-2003 |