

**K830655 QUIK-SILVER RESTING ECG ELECTRODE**Jun 15, 1983  
105 days to decisionK830655 · Product code: **DRX** · CardiovascularSource: <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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k830655/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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