

**K821757 QUIK - PREP2 ELECTRODE**Jul 2, 1982  
17 days to decisionK821757 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k821757/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jun 15, 1982
Decision date	Jul 2, 1982
Days to decision	17 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/k821757/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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