

K782079 ELECTRODE SYSTEM, QUIK-PREPFeb 1, 1979
50 days to decisionK782079 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k782079/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Dec 13, 1978
Decision date	Feb 1, 1979
Days to decision	50 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k782079/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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