

K810563 QUINTON LTE ELECTRODEMar 17, 1981
14 days to decisionK810563 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k810563/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Mar 3, 1981
Decision date	Mar 17, 1981
Days to decision	14 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.net

Device record: <https://www.510kdatabase.net/k810563/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026