

K840465 Q-MED ELECTRODEJun 19, 1984
137 days to decisionK840465 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k840465/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Feb 3, 1984
Decision date	Jun 19, 1984
Days to decision	137 days
Third-party review	No

APPLICANT

Company	Qmed, Inc.
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1984-1998

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k840465/> 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 June 29, 2026