

**K945836 1800 MONITORING ECG ELECTRODE AND 1870
DIAPHORETIC MONITORING ECG ELECTRODES**Jun 20, 1995
203 days to decisionK945836 · Product code: **DRX** · Cardiovascular
Source: <https://www.510kdatabase.net/k945836/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Nov 29, 1994
Decision date	Jun 20, 1995
Days to decision	203 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Conmed Andover Medical, Inc.
Location	Haverhill, MA, US
Contact	BRENMDA CULLINANCE
510(k) history	8 submissions · 8 cleared · 1994-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k945836/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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