

**K963787 EXACTRACE ADULT FOAM MONITORING ECG
ELECTRODE CONMED MODEL 1510 AND INSTATRACE ADULT
DIAPHORETIC FOAM ECG ELECTRODE CONM**Mar 11, 1997
172 days to decisionK963787 · Product code: **DRX** · Cardiovascular
Source: <https://www.510kdatabase.net/k963787/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Sep 20, 1996
Decision date	Mar 11, 1997
Days to decision	172 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Conmedcorp
Location	Dayton, OH, US
Contact	IRA D DUESLER, JR.
510(k) history	92 submissions · 92 cleared · 1981-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k963787/>; 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 27, 2026