

K982521 SKINTACT ECG ELECTRODES, S&W ECG ELECTRODESOct 26, 1999
463 days to decisionK982521 · Product code: **DRX** · Cardiovascular
Source: <https://www.510kdatabase.net/k982521/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jul 20, 1998
Decision date	Oct 26, 1999
Days to decision	463 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Leonhard Lang Co.
Location	Stillwater, MN, US
Contact	BURRHUS LANG
510(k) history	2 submissions · 2 cleared · 1999-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k982521/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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