

K030509 SKINTACT ECG TAB ELECTRODEMar 4, 2003
13 days to decisionK030509 · Product code: **DRX** · Cardiovascular
Source: <https://www.510kdatabase.net/k030509/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Feb 19, 2003
Decision date	Mar 4, 2003
Days to decision	13 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Leonhard Lang GmbH
Location	Stillwater, MN, US
Contact	ELAINE DUNCAN
510(k) history	17 submissions · 17 cleared · 2002-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030509/> 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 5, 2026