

K041883 SKINTACT MULTIFUNCTION ELECTRODES, MODEL DF 21Sep 7, 2004
57 days to decisionK041883 · Product code: **MKJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k041883/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jul 12, 2004
Decision date	Sep 7, 2004
Days to decision	57 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/k041883/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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