FDA 510(k) Premarket Notification

K041883 SKINTACT MULTIFUNCTION ELECTRODES, MODEL DF 21

CLEARED

Sep 7, 2004
57 days to decision

K041883 · 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