

K151329 SenTec Digital Monitor, OxiVent Sensor, V-Sign Sensor, Staysite Adhesive Pad, V-STATS PC Software including V-CareNet

Dec 17, 2015
213 days to decision

K151329 · Product code: **LKD** · Anesthesiology
Source: <https://www.510kdatabase.net/k151329/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Carbon-dioxide, Cutaneous (LKD)
Date received	May 18, 2015
Decision date	Dec 17, 2015
Days to decision	213 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sentec AG
Location	Egale, WI, US
Contact	ANKE WEISBRICH
510(k) history	5 submissions · 5 cleared · 2007-2025

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k151329/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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