K101690 SENTEC DIGITAL MONITOR, V-SIGN SENSOR AND V-SIGN SENSOR 2 WITH DIGITAL SENSOR ADAPER CABLE

CLEARED

Dec 3, 2010
170 days to decision

K101690 · Product code: **LKD** · Anesthesiology
Source: https://www.510kdatabase.net/k101690/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Carbon-dioxide, Cutaneous (LKD)
Date received	Jun 16, 2010
Decision date	Dec 3, 2010
Days to decision	170 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sentec AG**
Location	Egale, WI, US
Contact	STEPHEN H GORSKI
510(k) history	5 submissions · 5 cleared · 2007-2025

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k101690/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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