FDA 510(k) Premarket Notification

K121768 COSENSE

CLEARED

Oct 31, 2012
138 days to decision

K121768 · Product code: **CCJ** · Anesthesiology
Source: https://www.510kdatabase.net/k121768/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-monoxide, Gaseous-phase (CCJ)
Date received	Jun 15, 2012
Decision date	Oct 31, 2012
Days to decision	138 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Capnia, Inc.**
Location	Palo Alto, CA, US
Contact	Julie Blacklock
510(k) history	3 submissions · 3 cleared · 2012-2015

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k121768/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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