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Deviations from NIRS-derived optimal blood pressure are associated with worse outcomes after pediatric cardiac arrest

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Abstract

Aim: Evaluate cerebrovascular autoregulation (CAR) using near-infrared spectroscopy (NIRS) after pediatric cardiac arrest and determine if deviations from CAR-derived optimal mean arterial pressure (MAP_{opt}) are associated with outcomes.

Methods: CAR was quantified by a moving, linear correlation between time-synchronized mean arterial pressure (MAP) and regional cerebral oxygenation, called cerebral oximetry index (COx). MAP_{opt} was calculated using a multi-window weighted algorithm. We calculated burden

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Conflicts of interest

Dr. Morgan is supported by a grant from NIH NHLBI (K23HL148541). Dr. Sutton is supported by NIH grants to the institution. Dr. Topjian is the Chair of the AHA Emergency Cardiovascular Care Committee Pediatric Guidelines Writing Group. The remaining authors have disclosed that they do not have any conflicts of interest.

Appendix A. Supplementary material

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.resuscitation.2021.09.023>.

(magnitude and duration) of MAP less than 5 mmHg below MAP_{opt} ($MAP_{opt} - 5$), as the area between MAP and $MAP_{opt} - 5$ curves using numerical integration and normalized as percentage of monitoring duration. Unfavorable outcome was defined as death or pediatric cerebral performance category (PCPC) at hospital discharge ≥ 3 with ≥ 1 change from baseline. Univariate logistic regression tested association between burden of MAP less than $MAP_{opt} - 5$ and outcome.

Results: Thirty-four children (median age 2.9 [IQR 1.5,13.4] years) were evaluated. Median COx in the first 24 h post-cardiac arrest was 0.06 [0,0.20]; patients spent 27% [19,43] of monitored time with $COx \leq 0.3$. Patients with an unfavorable outcome ($n = 24$) had a greater difference between MAP and $MAP_{opt} - 5$ (13 [11,19] vs. 9 [8,10] mmHg, $p = 0.01$) and spent more time with MAP below $MAP_{opt} - 5$ (38% [26,61] vs. 24% [14,28], $p = 0.03$). Patients with unfavorable outcome had a higher burden of MAP less than $MAP_{opt} - 5$ than patients with favorable outcome in the first 24 h post-arrest (187 [107,316] vs. 62 [43,102] mmHg \times Min/Hr; OR 4.93 [95% CI 1.16–51.78]).

Conclusions: Greater burden of MAP below NIRS-derived $MAP_{opt} - 5$ during the first 24 h after cardiac arrest was associated with unfavorable outcomes.

Keywords

Cardiac arrest; Cerebral autoregulation; Cerebrovascular autoregulation; Hypoxic ischemic brain injury; NIRS; Pediatrics

Introduction

Among 20,000 children with cardiac arrests each year in the United States, most do not survive to hospital discharge and many survivors sustain new or worsened neurologic disability.^{1–6} The goal of post-cardiac arrest care is to reduce secondary brain injury and improve neurologic outcomes.^{7,8} Hypotension after cardiac arrest can cause cerebral hypoperfusion and is associated with worse outcomes.^{9–13} Thus, the American Heart Association (AHA) recommends maintaining a systolic blood pressure greater than the 5th percentile for age after cardiac arrest¹⁴; however, this approach does not account for variation in individual patients' cerebrovascular autoregulatory status that impacts cerebral perfusion.

Cerebrovascular autoregulation (CAR) maintains cerebral blood flow over a range of arterial blood pressures (ABP).^{15–17} After cardiac arrest, CAR is impaired, which can lead to reduced cerebral perfusion and secondary brain injury.^{7,8,18–20} CAR integrity can be quantified by the correlation between mean arterial pressure (MAP) and a surrogate of cerebral blood flow; a positive correlation reflects loss of CAR integrity. The cerebral oximetry index (COx) is the correlation of regional cerebral tissue oximetry (StO₂) derived from near infrared spectroscopy (NIRS) and MAP. The MAP at which a patient's CAR is most intact can be derived using COx, so-called optimal MAP (MAP_{opt}).^{21,22} To date, small adult and pediatric post-cardiac arrest studies have demonstrated associations between impaired CAR, CAR-derived MAP_{opt} , and patient outcomes.^{22–28} However, it is unknown if

deviations from CAR-derived MAP_{opt} are associated with outcomes after pediatric cardiac arrest and how CAR-derived MAP_{opt} compares to pediatric blood pressure targets.

The primary objective of this study was to determine the association of deviations from CAR-derived MAP_{opt} during the first 72 h after cardiac arrest using NIRS-derived StO_2 with outcomes. Secondary objectives were to determine the association of the severity of CAR impairment with outcomes and the age-based percentiles that correspond with CAR-derived MAP_{opt} . We hypothesized that patients with unfavorable outcomes would have a larger magnitude and duration of MAP deviation below CAR-derived MAP_{opt} than patients with favorable outcomes.

Methods

Study design

This was a retrospective analysis of prospectively collected data on patients ≥ 18 years old who received ≥ 1 min of cardiopulmonary resuscitation (CPR) for an in- or out-of-hospital cardiac arrest between November 2018 and March 2021 and received post-cardiac arrest care in the Children's Hospital of Philadelphia pediatric intensive care unit (PICU). Eligible patients had an invasive arterial catheter and either unilateral or bilateral cerebral NIRS connected to our integrative multimodality neuromonitoring device (Moberg Research, Ambler, PA, USA). Patients were excluded if they received extracorporeal support, had unrepaired cyanotic congenital heart disease, had concomitant severe acute brain injury due to traumatic brain injury or ruptured vascular malformation, or had formal limitations of care at time of eligibility. Additional exclusion criteria were time between ROSC and initiation of multimodal recording >24 h or recording duration less than 6 h.

Post-cardiac arrest care was determined by the clinical team, guided by an institutional pathway.²⁹ The clinical team was blinded to CAR and MAP_{opt} data. Prospectively defined clinical data were abstracted via medical record review and included patient demographics, cardiac arrest characteristics, and details regarding post-cardiac arrest care. Research coordinators abstracting data were blinded to the primary outcome. The study was approved by the Children's Hospital of Philadelphia Institutional Review Board with a waiver of consent.

Clinical outcomes

Clinical outcome was based on Pediatric Cerebral Performance Category (PCPC) scores, a 6-point scale of global neurologic function: (1) normal; (2) mild disability; (3) moderate disability; (4) severe disability; (5) coma or vegetative state; (6) death.^{30–32} It was assigned by trained nurse raters for preadmission baseline and at hospital discharge via medical record review and discussions with medical providers. Discrepancies were resolved by consensus of an independent internal review committee. Unfavorable outcome was defined as death or a change in PCPC ≥ 1 from pre-admission that resulted in PCPC score of 3, 4, or 5 at hospital discharge or 30 days post-cardiac arrest, whichever came first. Favorable outcome was defined as PCPC of 1 or 2 at hospital discharge or 30 days post-cardiac arrest,

whichever came first, or no change in PCPC from baseline to hospital discharge or 30 days post-cardiac arrest.

Neuromonitoring signal acquisition and processing

Decisions to place an arterial catheter or NIRS device (Nonin SenSmart (Nonin Medical, Inc., Plymouth, MN, USA) or Edwards Foresight (Edwards Lifesciences Corporation, Irvine, CA, USA)) were made by the clinical team.²⁹ An integrative bedside multimodality neuromonitoring device (Moberg Research, Ambler, PA, USA) was deployed by nurses per provider request and device availability. The device facilitated time synchronization of ABP and StO₂ signals.

Artifacts were computationally removed using a semi-automated sliding-window median filter with variable window length to account for differences in amount of artifact. Given the homogeneous nature of post-cardiac arrest hypoxic-ischemic brain injury, right and left cerebral StO₂ values were averaged together when both were available. Data obtained from the first 72 h after ROSC were divided into post-cardiac arrest Day 1 (0–24 h), Day 2 (24–48 h), and Day 3 (48–72 h).

Cerebrovascular autoregulation and MAP_{opt} determination

CAR was assessed using NIRS-based cerebral oximetry index (COx), calculated as a Pearson correlation coefficient between 10-s averaged values of MAP and corresponding NIRS-measured StO₂ over a 5-min window.^{33,34} COx values ranged from –1 to +1; negative or near-zero COx values, which result from MAP and StO₂ being either negatively correlated or not correlated, respectively, indicated intact CAR. In contrast, impaired CAR was indicated by positive COx values due to MAP and StO₂ being positively correlated. By moving the 5-min window by 1 min and repeating the correlation (80% overlap of data), COx can be analyzed as a continuous variable.³⁵ A COx = 0.3 was *a priori* defined as impaired CAR.³⁶

To determine each patient's MAP_{opt} over time, we used a multi-window approach.³⁷ COx was calculated using 3, 5, 10, 20, 30, 60, 90, and 120-min windows. Using windows of data from the prior 1–24 h (i.e., 1, 2, 4, 6, 8, 12, and 24 h), for each COx window, we plotted COx values versus corresponding MAP in 5 mmHg bins (e.g., 50–55 mmHg). We applied the Fisher transform to binned COx values to avoid ceiling effects.²¹ We then applied an automated curve fitting algorithm to fit a second-order polynomial representing a convex parabola.^{21,24,37–40} The nadir of the fitted curve (i.e., MAP where COx was most negative) represented MAP_{opt}. This process generated up to 51 COx versus MAP parabolic curves that were combined using a weighted average to determine MAP_{opt}. Curves with a better convex parabolic fit (i.e., greater adjusted r² value) and those with more negative COx values at the nadir were given greater weight to generate MAP_{opt}.^{37,38,41,42} This process was repeated to calculate an updated MAP_{opt} every minute.

We also determined lower and upper limits of autoregulation (LLA and ULA, respectively) over time for each patient. The LLA is the MAP at which CBF decreases with decreasing MAP. Similarly, ULA is the MAP at which CBF increases with increasing MAP. We defined LLA and ULA as MAPs where COx-MAP parabolic curves crossed a COx value of 0.3.³⁸

Data analyses

We compared each patient's MAP to their CAR-derived $MAP_{opt} \pm 5$ mmHg to account for anticipated clinical variations in MAP and to make the results more clinically applicable.^{24,26} For each subject and for each post-cardiac arrest day, we calculated mean magnitude of difference between MAP and $MAP_{opt} - 5$ mmHg during times when $MAP < MAP_{opt} - 5$ mmHg. Analogously, we calculated mean magnitude of the difference between MAP and $MAP_{opt} + 5$ mmHg during the times when $MAP > MAP_{opt} + 5$ mmHg. We further calculated the percent of time MAP was 1) below $MAP_{opt} - 5$ mmHg, 2) within $MAP_{opt} \pm 5$ mmHg, and 3) above $MAP_{opt} + 5$ mmHg, for Days 1, 2 and 3 post-cardiac arrest. Using numerical integration, the burden (combination of magnitude and duration; mmHg \times min/hour) of MAP below $MAP_{opt} - 5$ was defined as the normalized area between MAP and $MAP_{opt} - 5$ waveforms during the times when $MAP < MAP_{opt} - 5$ (Fig. 1). This burden was computed for each post-cardiac arrest day and normalized by the amount of time concomitant MAP and MAP_{opt} waveforms were available. Analogous burdens of MAP above $MAP_{opt} + 5$ (i.e., normalized area between MAP and $MAP_{opt} + 5$ waveforms during the times when $MAP > MAP_{opt} + 5$) were also computed for each post-cardiac arrest day.

In a secondary analysis, we calculated the burden of impaired CAR (i.e., the normalized area between the COx waveform and the COx threshold of 0.3 during the times when $COx > 0.3$ (Fig. 1)), for each post-cardiac arrest day.^{43,44} We additionally investigated brain oxygenation in relation to MAP by computing the mean StO₂ for each patient when MAP was 1) less than $MAP_{opt} - 5$, 2) within $MAP_{opt} \pm 5$, and 3) greater than $MAP_{opt} + 5$ for each post-cardiac arrest day. Finally, using published normative values from critically ill children,⁴⁵ we determined age-based MAP percentile equivalent for MAP, MAP_{opt} , LLA, and ULA, for each post-cardiac arrest day.

Statistical analysis

Descriptive statistics are reported as median and interquartile ranges (IQR) for continuous variables and frequencies with percentages for categorical variables. Chi-squared or Fisher exact tests were used to test associations between categorical variables and clinical outcome and Wilcoxon rank-sum was used to compare differences in continuous variables between outcome groups. All statistical tests were two-sided, and $p < 0.05$ was considered to indicate significance. The primary exposure was the burden of MAP less than $MAP_{opt} - 5$. Univariate logistic regression model tested the association between the normalized burden of MAP less than $MAP_{opt} - 5$ and outcome. Analyses were performed using GraphPad Prism (v5.03, GraphPad Software Inc., La Jolla, CA, USA), IBM SPSS Statistics (v26.0, IBM Corp., Armonk, NY, USA) or MATLAB (vR2018a, The Mathworks, Inc., Natick, MA, USA).

3D visualization of the burden of MAP below MAP_{opt}

A customized 3D visualization technique depicted the association between magnitude and duration of MAP below MAP_{opt} and outcome.⁴⁶ Using minute-by-minute MAP and MAP_{opt} data, for each pair of a given magnitude (MAP below MAP_{opt} by 0 to 30 mmHg) and for a given duration (0–40 min), the ratio between number of patients with an unfavorable outcome and total number of patients who experienced at least one such episode was

recorded. This ratio, indicating probability of an unfavorable outcome, was displayed on a 3D color-coded contour plot.

Results

Forty-six patients met inclusion criteria. Twelve patients were excluded due > 24 h from ROSC to data recording ($n = 3$), inadequate recording duration ($n = 4$), ECMO ($n = 2$), or concomitant non-hypoxic-ischemic severe acute brain injury ($n = 3$). Data from 34 patients were analyzed. The median age was 2.9 [IQR 1.5, 13.4] years and 71% ($n = 24$) were male. Thirty-eight percent had no past medical history and 59% had baseline PCPC of 1. Seventy-one percent ($n = 24$) of patients had unfavorable outcomes; 63% (15/24) did not survive to hospital discharge. Demographic and cardiac arrest characteristics are summarized in Table 1.

The median time between ROSC and initiation of data recording was 3 [2,7] h. A median of 52 [33, 63] h of concomitant MAP and StO₂ data were recorded in the first 72 h post-cardiac arrest.

Association between deviations from MAP_{opt} and outcome

We were able to calculate MAP_{opt} in 94% [86%, 97%] of recorded time. On post-cardiac arrest Day 1, the magnitude of difference between MAP and MAP_{opt} - 5 and percent of time MAP was less than MAP_{opt} - 5 were greater for patients with unfavorable compared to favorable outcomes (Table 2). Burden of MAP less than MAP_{opt} - 5 was greater for patients with unfavorable versus favorable outcomes (187 mmHg*Min/Hr [107, 316] vs. 62 mmHg*Min/Hr [43, 102], $p = 0.01$). Odds of unfavorable outcomes were 4.9 times higher for each standard deviation increase in burden of MAP less than MAP_{opt} - 5 (OR 4.93 [95% CI 1.16 to 51.78]). Fig. 2 demonstrates the impact of magnitude and duration of MAP less than MAP_{opt} - 5 on probability of unfavorable outcome.

There were no differences between patients with favorable and unfavorable outcomes for magnitude, duration, or burden of MAP less than MAP_{opt} - 5 on Days 2 or 3 post-cardiac arrest (Supplementary Table 1). Odds of unfavorable outcomes were not increased based on burden of MAP less than MAP_{opt} - 5 for Days 2 (OR 1.60 [0.72 to 4.59]) or 3 (OR 1.21 [0.58 to 3.08]) post-cardiac arrest. Magnitude, duration, and burden of MAP within MAP_{opt} ± 5 or greater than MAP_{opt} + 5 were not significantly different between patients with favorable or unfavorable outcomes on any post-cardiac arrest day (Supplementary Table 1).

There were no differences in StO₂ between patients with unfavorable or favorable outcomes on any day after arrest (Table 2 and Supplementary Table 1).

CAR impairment

COx was 0.3 28% [20%, 35%] of time. Median COx, percent time COx 0.3, and percent burden COx 0.3 on post-cardiac arrest Days 1, 2, or 3 did not differ between favorable and unfavorable outcome groups (Table 2 and Supplemental Table 1).

Comparing MAP_{opt}, LLA and ULA to age-based MAP percentiles

MAP_{opt} for post-cardiac arrest Days 1, 2, and 3 was equivalent to the 77th [62, 88] percentile of MAP for age. The LLA and ULA were equivalent to the 22nd [13,37] and 98th [94, 99] percentiles of MAP for age. The range between LLA and ULA was 38 mmHg [29, 44], and did not differ between favorable and unfavorable outcome groups (38 mmHg [27, 43] vs 39 mmHg [29, 44], $p = 0.696$). There were no differences in percentiles of MAP_{opt}, LLA and ULA for age between favorable and unfavorable outcome groups on any post-cardiac arrest day (Table 3).

Discussion

In this single center study of CAR following pediatric cardiac arrest, patients whose MAP was more than 5 mmHg below their CAR-derived MAP_{opt} in the first 24 h after cardiac arrest were more likely to have unfavorable outcomes. Both magnitude and duration of MAP deviation from MAP_{opt} were associated with unfavorable outcomes, however the severity of CAR impairment, as measured by COx, was not. Notably, CAR-derived MAP_{opt} was equivalent to the 77th percentile for age and the difference between the lower and upper limits of CAR was 38 mmHg, suggesting that after cardiac arrest the MAP range of intact CAR is substantially narrowed and that blood pressures higher than age-based means may be required to maintain adequate cerebral perfusion.

The association between magnitude and duration of MAP deviation below CAR-derived MAP_{opt} and outcomes implies that cerebral hypoperfusion occurs when MAP is below MAP_{opt} and this critical reduction in cerebral blood flow contributes to secondary brain injury and unfavorable outcomes. Interestingly, COx, an index of CAR, was not different between patients with favorable and unfavorable outcomes and the burden of impaired CAR was also not different. Thus, impaired CAR may predispose patients to brain injury, but both duration and magnitude of MAP deviation below each patient's individual MAP_{opt} is a more substantial contributor to secondary brain injury.

Our results build upon findings from adults^{23,24,26,47} and children²⁵ that have examined CAR using NIRS after cardiac arrest and found varying associations between impaired CAR, deviation of MAP from MAP_{opt}, and outcomes.²² Lee and colleagues found that a greater area under the MAP_{opt} curve, similar to the computed burden in the current study, on day 2 after cardiac arrest was associated with children receiving tracheostomy or gastrostomy tubes but did not find an association with changes in PCPC scores.²⁵ Ameloot et al. demonstrated percentage time spent below MAP_{opt} was negatively associated with survival.²³ A recent multicenter adult study using a similar multi-window approach to our study failed to find differences in CAR metrics and deviation from MAP_{opt} between outcome groups.²⁴

The AHA guidelines recommend maintaining a systolic blood pressure greater than the 5th percentile for age during post-cardiac arrest care.¹⁴ This threshold was based on observational studies that demonstrated worse outcomes, primarily survival, when systolic hypotension (below 5th percentile for age and sex) was present following cardiac arrest.^{7,14} In our study, CAR-derived MAP_{opt} was approximately the 75th percentile for age. In

adults, targeting a higher MAP (80–100 mmHg versus 65–75 mmHg) improved cerebral oxygenation, but not biomarkers of brain injury, hypoxic-ischemic injury on MRI, or neurologic outcomes.^{48,49} Our data suggest that active titration of blood pressure to cerebral hemodynamic parameters like MAP_{opt} or LLA is a potentially promising approach to improve outcomes rather than simply avoiding age-based 5th percentile blood pressures. This approach is the subject of an ongoing feasibility study in adults with traumatic brain injury.⁴¹

This study has several limitations. Although cerebral NIRS is attractive because of non-invasiveness and feasibility, the NIRS signal can be influenced by sensor placement, ambient light, and scalp blood flow, which would tend to skew our results to the null. Nevertheless, NIRS-derived CAR parameters were associated with outcomes. Due to small sample size, we were unable to control for cardiac arrest characteristics associated with outcomes (e.g., arrest location, arrest duration, initial rhythm). Due to small sample size, there was the potential for not finding significant differences where they may exist (type II errors). Similarly, with multiple comparisons in a small sample there was the potential for type I error. Future studies of larger cohorts of patients are necessary to adequately address these limitations. As in previous studies, duration of CPR was associated with outcomes.⁷ Because CAR impairment after cardiac arrest presumably reflects severity of brain injury, it is thus in the causal pathway from injury severity to outcome. Future larger studies are needed to evaluate the association of pre-, intra- and post-cardiac arrest features on CAR impairment. We used a COx threshold of 0.3 to define impaired CAR, consistent with adult post-cardiac arrest studies,^{24,26} although pediatric trials are needed to more clearly define thresholds of impaired CAR that are associated with outcomes. All patients had invasive arterial catheters, cerebral NIRS, and multimodality neuromonitoring. Therefore, these data may not be generalizable to less severely injured patients without such intensive post-cardiac arrest monitoring. While we attempted to enroll consecutive patients, data collection was limited by technical considerations or clinical circumstances. Some patients did not have data collection through Day 3 after cardiac arrest, mainly due to clinical decisions to remove the arterial catheter or stop NIRS monitoring.

Conclusions

A greater burden of MAP below NIRS-derived MAP_{opt} - 5 was associated with unfavorable outcomes in children within the first 24 h after cardiac arrest. Further research is needed to determine whether active titration of blood pressure to cerebral hemodynamic parameters like MAP_{opt} can limit secondary brain injury and improve outcomes after pediatric cardiac arrest.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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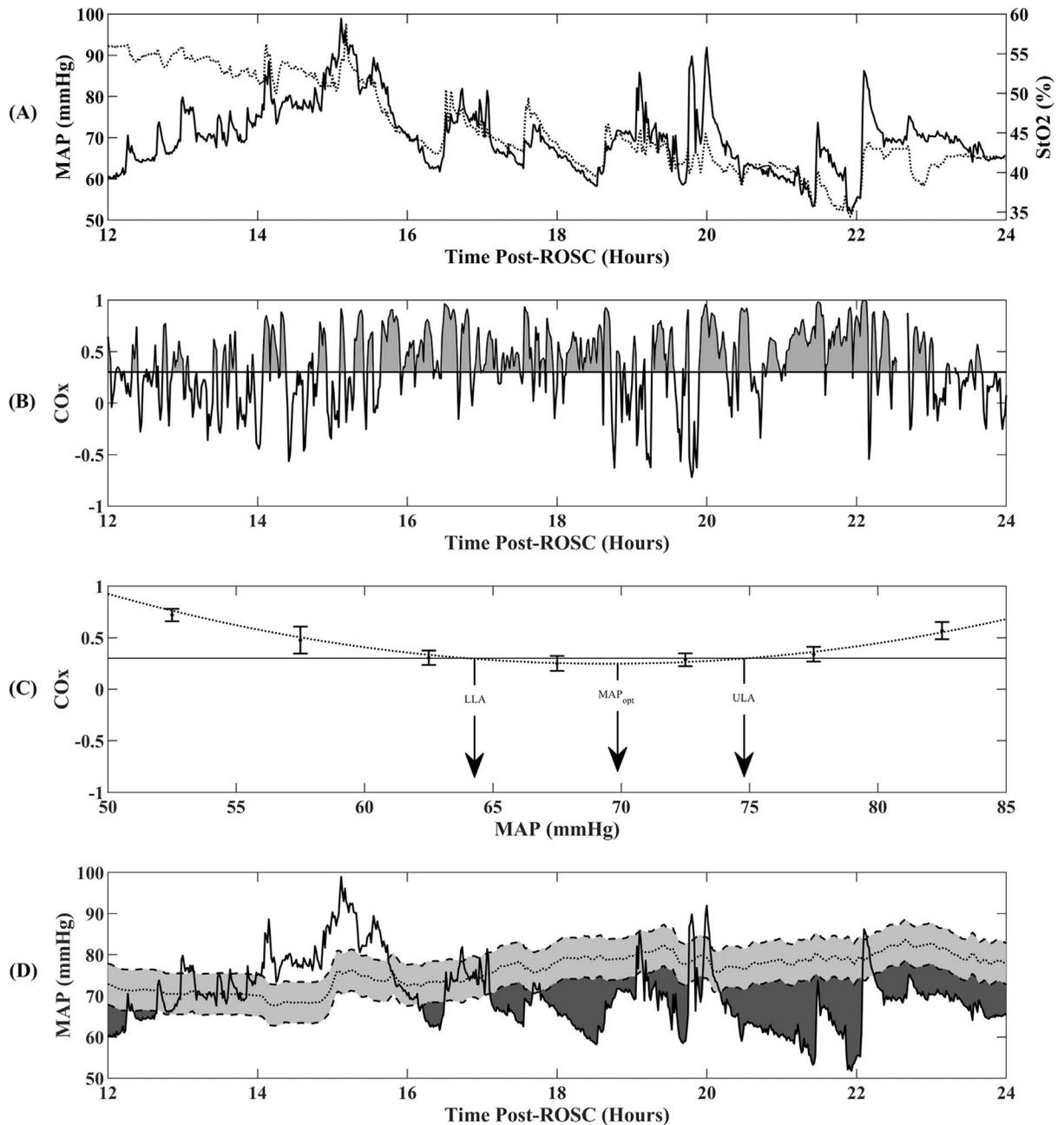


Fig. 1 –.

Physiologic data over 12 h of recording from a representative patient. The four rows depict (A) mean arterial pressure (MAP, solid line) and regional cerebral tissue oximetry (StO₂, dotted line) derived from NIRS; (B) cerebral oximetry index (CO_x), the gray shaded region on the CO_x curve represents the burden of impaired CAR which was defined as the area below the CO_x waveform and CO_x threshold of 0.3, and normalized as a percentage of the monitoring duration; (C) representative CO_x versus binned MAP parabolic curve demonstrating MAP_{opt} at the nadir of the curve and the lower and upper limits of autoregulation (LLA and ULA) where parabolic curve crosses 0.3; and (D) MAP (solid

line), MAP_{opt} (dotted line), gray shaded region is ± 5 mmHg of MAP_{opt} , black region is burden of MAP less than $MAP_{opt} - 5$.

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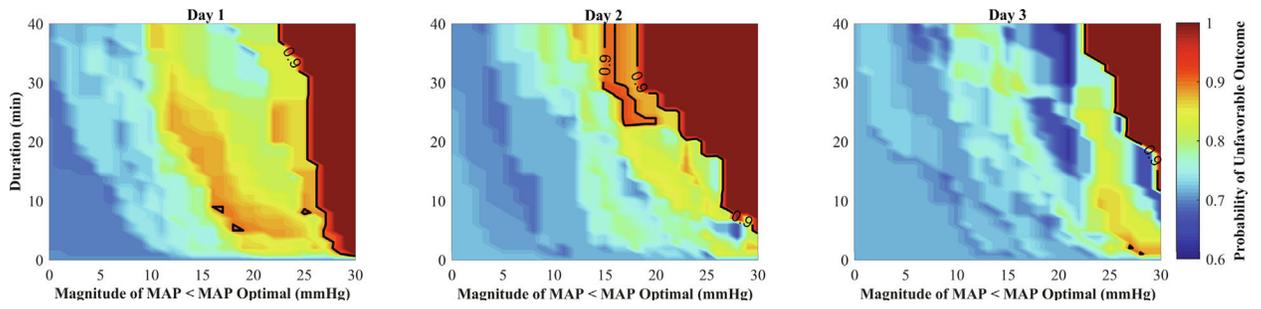


Fig. 2 –.

Association between magnitude and duration of MAP less than MAP_{opt} on the probability of an unfavorable outcome for Days 1, 2, and 3 after cardiac arrest. Color scale represents the probability of an unfavorable outcome. Dark blue indicates low probability of an unfavorable outcome and dark red indicates a high probability of an unfavorable outcome.

Table 1 –

Patient demographics, cardiac arrest characteristics by outcomes.

Patient demographics/Cardiac arrest characteristics	All Patients (n = 34)	Favorable Outcome (n = 10)	Unfavorable Outcome (n = 24)	P-value
Age (years), median' (IQR)	2.9 [1.5, 13.4]	11.7 [2.7, 14.6]	2.7 [0.7, 9.2]	0.11
Gender (male), N (%)	24 (71)	9 (90)	15 (63)	0.22
Preexisting conditions				
None	13 (38)	6 (60)	7 (29)	0.09
Premature infant	10 (29)	2 (20)	8 (33)	0.68
Congenital heart disease	6 (18)	1 (10)	5 (21)	0.64
Congenital malformation	4 (12)	0 (0)	4 (17)	0.30
Cancer/BMT	3 (9)	1 (10)	2 (8)	1.00
Respiratory disorder/CLD	10 (29)	1 (10)	9 (38)	0.22
Renal insufficiency	4 (12)	1 (10)	3 (13)	1.00
Neurological disorder	5 (15)	0 (0)	5 (21)	0.29
Solid organ transplant	1 (3)	0 (0)	1 (4)	1.00
Baseline PCPC				
PCPC 1	20 (59)	8 (80)	12 (50)	0.47
PCPC 2	4 (12)	0 (0)	4 (17)	
PCPC 3	6 (18)	1 (10)	5 (21)	
PCPC 4	4 (12)	1 (10)	3 (13)	
Arrest Etiology				0.19
Respiratory failure	12 (35)	3 (30)	9 (38)	
Arrhythmia	3 (9)	3 (30)	0 (0)	
Hypotension/shock/sepsis	3 (9)	1 (10)	2 (8)	
Ingestion/toxic	1 (3)	0 (0)	1 (4)	
Metabolic/electrolyte	2 (6)	1 (10)	1 (4)	
Seizures	1 (3)	0 (0)	1 (4)	
Drowning	3 (9)	0 (0)	3 (13)	
Unknown	9 (26)	2 (20)	7 (29)	
Out-of-hospital cardiac arrest	22 (65)	5 (50)	17 (71)	0.27
CPR duration (min), (n = 27)	12 [6,32]	6 [5,10]	28 [9, 36]	0.01

Patient demographics/Cardiac arrest characteristics	All Patients (n = 34)	Favorable Outcome (n = 10)	Unfavorable Outcome (n = 24)	P-value
Number of Epi doses				0.09
0–2	19 (56)	9 (90)	10 (42)	
3–5	4 (12)	0 (0)	4 (17)	
>5	7 (21)	1 (10)	6 (25)	
Unknown	4 (12)	0 (0)	4 (17)	
Initial Rhythm				0.06
PEA/Asystole	17 (50)	3 (30)	14 (58)	
Bradycardia	4 (12)	2 (20)	2 (8)	
V-tachycardia/fibrillation	3 (9)	3 (30)	0 (0)	
Not documented/Unknown	10 (29)	2 (20)	8 (33)	
Initial pH, median, (IQR)	6.98 [6.81, 7.19]	7.18 [7.05, 7.28]	6.87 [6.75, 7.09]	0.004
Initial Lactate, median* (IQR)	7.3 [4.8, 11.3]	6.0 [4.9, 7.3]	8.0 [4.7, 12.7]	0.14
Temperature target				0.07
33.0 °C	4 (12)	3 (30)	1 (4)	
36.0 °C	30 (88)	7 (70)	23 (96)	
PCPC (hospital discharge)				<0.001
PCPC 1	3 (9)	3 (30)	0 (0)	
PCPC 2	5 (15)	5 (50)	0 (0)	
PCPC 3	5 (15)	1 (10)	4 (17)	
PCPC 4	5 (15)	1 (10)	4 (17)	
PCPC 5	2 (6)	0 (0)	2 (8)	
PCPC 6	14 (41)	0 (0)	14 (58)	
PCPC Change (Discharge - Baseline)				<0.001
0	5 (15)	5 (50)	0 (0)	
1	7 (21)	5 (50)	2 (8)	
2	9 (27)	0 (0)	9 (38)	
3	4 (12)	0 (0)	4 (17)	
4	2 (6)	0 (0)	2 (8)	
5	7 (21)	0 (0)	7 (29)	

Table 2 –

Burden of MAP deviation from MAP_{opt}, index of impaired CAR, and StO₂ on Day 1 after cardiac arrest.

Cerebral physiology parameter Median [IQR]	Favorable (n = 10)	Unfavorable (n = 24)	p-value
MAP < MAP _{opt} -5			
Magnitude MAP below MAP _{opt} -5 (mmHg)	9 [8, 10]	13 [11, 17]	0.01
Percent time MAP < MAP _{opt} -5 (%)	24 [14, 28]	38 [26, 61]	0.03
Burden MAP < MAP _{opt} -5 (mmHg*Min/Hr)	62 [43, 102]	187 [107, 318]	0.01
MAP within ± 5 of MAP _{opt}			
Percent time MAP within ± 5 of MAP _{opt} (%)	36 [28, 40]	31 [17, 43]	0.30
MAP > MAP _{opt} + 5			
Magnitude MAP above MAP _{opt} + 5 (mmHg)	12 [10, 12]	12 [9, 15]	0.83
Percent time MAP > MAP _{opt} + 5 (%)	36 [27, 44]	21 [13, 38]	0.07
Burden MAP > MAP _{opt} + 5 (mmHg*Min/Hr)	272 [198, 306]	207 [92, 295]	0.14
COx			
COx	0.06 [-0.01, 0.20]	0.08 [-0.00, 0.20]	0.92
Percent time COx > 0.3 (%)	25 [19, 42]	32 [19, 44]	0.78
Percent COx burden > 0.3 (%)	8 [4, 16]	10 [5, 16]	0.66
StO ₂			
StO ₂ (%)	76 [70, 81]	70 [59, 78]	0.07
StO ₂ when MAP < MAP _{opt} -5 (%)	76 [73, 80]	70 [58, 78]	0.10
StO ₂ when MAP within ± 5 MAP _{opt} (%)	76 [69, 83]	71 [59, 79]	0.11
StO ₂ when MAP > MAP _{opt} + 5 (%)	77 [71, 81]	69 [60, 78]	0.07

Table 3 – Age-based blood pressure percentiles of MAP, MAP_{opt}, LLA and ULA by outcome.

Age-based percentile of MAP Median [IQR]	Favorable* (n = 10)	Unfavorable (n = 24)	p-value
Age-based percentile of MAP			
Day 1	77th [49, 85]	64th [48, 93]	0.84
Day 2	63rd [36, 91]	70th [51, 82]	0.80
Day 3	66th [36, 86]	77th [61, 85]	0.49
Age-based percentile of MAP _{opt}			
Day 1	68th [56, 80]	83rd [58, 94]	0.26
Day 2	66th [59, 77]	69th [59, 83]	0.56
Day 3	69th [43, 88]	82nd [65, 93]	0.23
Age-based percentile of LLA			
Day 1	24th [6, 34]	27th [19, 46]	0.20
Day 2	8th [3, 31]	23rd [16, 32]	0.15
Day 3	7th [3, 29]	23rd [8, 47]	0.13
Age-based percentile of ULA			
Day 1	94th [91, 98]	98th [90, 99]	0.28
Day 2	96th [93, 99]	95th [92, 99]	0.86
Day 3	98th [92, 99]	99th [94, 99]	0.32

* Day 2: n = 9 for favorable patients and 23 for unfavorable patients.

Day 3: n = 8 for favorable patients and 20 for unfavorable patients.