

research paper

Multimodal long term predictors of outcome in out of hospital cardiac arrest patients treated with targeted temperature management at 36°C

Erik Roman-Pognuz^{1*}, Johnathan Elmer², Frank X Guyette², Gabriele Poillucci³, Umberto Lucangelo¹, Giorgio Berlot¹, Paolo Manganotti⁴, Alberto Peratoner¹, Tommaso Pellis⁵, Fabio S. Taccone⁶ and Clifton W. Callaway²

¹ Affiliation 1; Department of Anesthesia and Intensive Care, †Azienda sanitaria universitaria Giuliano Isontina; romanpognuz.erik@gmail.com, umberto.lucangelo@asuits.sanita.fvg.it, a.peratoner@gmail.com, berlotg@virgilio.it

² Affiliation 2; Department of Emergency Medicine, University of Pittsburgh. elmerjp@upmc.edu, callawaycw@upmc.edu, guyefx@UPMC.edu

³ Affiliation 3; Department of radiology. Azienda sanitaria universitaria Giuliano Isontina. gabrielepoillucci@gmail.com

⁴ Affiliation 4; Department of Neurology, University of Trieste. paolo.manganotti@asuits.sanita.fvg.it

⁵ Affiliation 5; Department of Intensive Care, Azienda sanitaria Friuli Occidentale tommaso.pellis@asfo.sanita.fvg.it

⁶ Affiliation 6; Department of Intensive Care, Erasme Hospital, Université Libre de Bruxelles. fabio.Taccone@ulb.ac.be

*Correspondence: romanpognuz.erik@gmail.com; Tel.:+39 3394879119

Abstract:

Introduction: Early prediction of long term outcomes in patients resuscitated after cardiac arrest (CA) is still challenging. Guidelines suggested a multimodal approach combining multiple predictors. We evaluated whether the combination of the electroencephalography (EEG) reactivity, somatosensory evoked potentials (SSEPs) cortical complex and Gray to White matter ratio (GWR) on brain computed tomography (CT) at different temperatures could predict survival and good outcome at hospital discharge and after six months. **Methods:** We performed a retrospective cohort study including consecutive adult, non-traumatic patients resuscitated from out-of-hospital CA who remained comatose on admission to our intensive care unit from 2013 to 2017. We acquired SSEPs and EEGs during the treatment at 36°C and after rewarming at 37°C. Gray to white matter ratio (GWR) was calculated on the brain computed tomography scan performed within six hours of the hospital admission. We primarily hypothesized that SSEP was associated with favorable functional outcome at distance and secondarily that SSEP provides independent information from EEG and CT. Outcomes were evaluated using the Cerebral Performance Category (CPC) scale at six months from discharge. **Results:** Of 171 resuscitated patients, 75 were excluded due to missing of data or uninterpretable neurophysiological findings. EEG reactivity at 37 °C has been shown the best single predictor of good outcome (AUC 0.803) while N20P25 was the best single predictor for survival at each time point. (AUC 0.775 at discharge and AUC 0.747 at six months follow up) Predictive value of a model including EEG reactivity, average GWR, and SSEP N20P25 amplitude was superior (AUC 0.841 for survival and 0.920 for good outcome) to any combination of two tests or any single test. **Conclusion:** Our study, in which life-sustaining treatments were never suspended, suggests SSEP cortical complex N20P25, after normothermia ad off sedation, is a reliable predictor for survival at any time. When SSEP cortical complex N20P25 is added into a model with GWR average and EEG reactivity, the predictivity for good outcome and survival at distance is superior than each single test alone.



Keywords: 1. Cardiac arrest, 2. normothermia, 3. EEG, 4. SSEP, 5. GWR, 6. long term predictors

1. Introduction

After cardiac arrest (CA) most resuscitated patients are comatose as a result of hypoxic ischaemic brain injury [1, 2]. Differentiating patients who can awaken from coma from those with irrecoverable injury remains challenging. Bilateral absence of N20 cortical somatosensory evoked potentials (SSEP) at least 72 hours after return of spontaneous circulation (ROSC) is one reliable indicator of poor prognosis. [3][4][5] However, presence of cortical responses on SSEPs does not guarantee favorable outcome. [6] Consensus guidelines recommend that clinicians use SSEP as one part of a multimodal approach to prognostication. [7]

There are several gaps in knowledge about SSEP as a prognostic modality after cardiac arrest. First, most studies relate SSEP to short term outcomes like awakening from coma or survival to hospital discharge rather than more important long-term outcomes like function at 3 or 6 months. Second, many studies have used SSEP results in the decision to limit life support for subsets of patients, creating bias that would inflate the SSEP performance. Finally, few studies describe how much incremental information SSEP results provide over other clinical information [8]. To address some of these gaps Scarpino et colleagues investigated the role of a combination of measures, like SSEP, EEG and GWR, to predict CA patients's cerebral performance after 6 months follow up. [9]

Most studies simplify the complex SSEP waveform into the dichotomous presence or absence of the N20 cortical response. It is not clear whether quantifiable features like latency and amplitude additional also provide prognostic information (Figure.1)

To address some of these gaps, we collected data on the prognostic value of SSEP waveforms performed at regimented times during and after targeted temperature management for predicting outcome after 6 months in a cohort of CA patients with no limitations in life support. We hypothesized that the amplitude of SSEP cortical waveforms is associated with favorable functional recovery, and that SSEP provides independent information from EEG and CT scan.

2. Materials and Methods

2.1 Study design and population.

We performed a retrospective cohort study including consecutive non-traumatic patients resuscitated from out-of-hospital CA who remained comatose (Glasgow Coma Scale score ≤ 8) on admission to an intensive care unit (ICU) at a single center tertiary university hospital from January 2013 to May 2017. Our local ethics committee approved the study and waived the need for informed consent because of minimal risk.

We excluded patients under 18 years old, with traumatic cardiac arrest, with pregnancy, with previously diagnosed progressive neurodegenerative disease, and those deemed likely to meet the criteria for progress to brain death on admission. (Figure 2)

2.2 Study definition and data collection.

We defined cardiac arrest as the abrupt cessation of cardiac activity that required shocks and/or chest compression for the return of spontaneous circulation (ROSC). We dichotomized initial rhythm as shockable (ventricular fibrillation or pulseless ventricular tachycardia) or non-shockable (pulseless electrical activity or asystole). We estimated the time interval from collapse to ROSC from prehospital reports.

We performed non-contrast enhanced computerized tomography (CT) scan of the brain to exclude neurological causes of arrest or head trauma in unwitnessed collapse. We treated all patients with targeted temperature management (TTM) to 36°C for 24

hours using an intravenous cooling system (CoolGard 3000/Alsius Icy Heat Exchange Catheter, Zoll Medical). We sedated patients with propofol, although we substituted midazolam in cases of severe hypotension. We provide neuromuscular blockade with continuous infusion of cisatracurium to prevent shivering during the induction of TTM. We stopped sedation and paralysis at the beginning of the rewarming period.

We measured survival and functional recovery at discharge from the ICU and at 6 months after the event using medical records and cardiologic reports. Reports characterized functional neurological recovery using the Pittsburgh-Glasgow Cerebral Performance Categories scale (CPC), and we defined good outcome as CPC 1-2.[10] [11]

2.2 a EEGs and SSEPs

We acquired SSEPs and EEGs twice, first during the first 12 hours after reaching the target temperature of 36° C (approximately within 12 hours after ROSC) and second after withdrawal of sedation and rewarming to 37° C (approximately 72 hours after ROSC).

We recorded EEGs for at least 20 minutes using a portable machine (Galileo NT PMS version 3.90/00/17014 - SPD. EBN Neuro) using a 13 electrodes positioned according to the international 10-20 system. [12] A neurophysiologist determined EEG reactivity as present if there was a clear change in background frequency or amplitude after a pain or voice stimulation.[13] We considered EEGs where electroencephalographic seizures resulted from stimulation to be nonreactive. EEG background was classified as continuous (recognizable clear background activity), discontinuous (burst suppression of almost 10% of the recordings) or flat (isoelectric or suppression less than 10 μ V). [13] Epileptiform activity was identified as rhythmic spikes or waves and sharp waves and periodic epileptiform discharges (PEDs). [14][15]

A certified neurologist evaluated peripheral, spinal and cortical SSEPs in response to the stimulation of the median nerve at the wrist. The cortical N20 amplitude was defined as the highest amplitude of a reproducible potential of CP3/CP4 vs. Fz recordings and CP3/CP4 vs contralateral earlobe at least at 4.5 ms longer than the previous N13 peak (cervical spinal cord) and within 50 ms after stimulation.[16] We considered as P25 amplitude the first positive wave that follows N20 wave. (Figure 1) Consequently we examined the peak to peak N20-P25 absolute amplitude value in both sides. N20-P25 complex has been used previously as a prognostic indicator particularly in ischemic stroke patients. [17] [18] [19] In patients with background noise levels under 0.25 μ V for whom we could not discern any cortical waveform bilaterally, we considered the SSEPs to be absent. Clinical teams in our hospital do not use SSEP recordings as part of any decisions to withdraw or limit life support.

2.2 b GWR

We acquired non-contrast head CT scans on Aquilion 64 (Toshiba Medical Systems Europe B.V., Zoetermeer, The Netherlands) or Brilliance iCT 256 (Philips Healthcare, Best, The Netherlands) scanners, using 5 mm slices reconstruction in the axial plane. Two investigators blinded to clinical information examined CT scans for each patient using commercial image viewing software (suitEstensa Ris Pacs Software, Esaote Healthcare IT, Genova, Italy) with windowing adjusted to "brain," and identified comparable brain slices at the level of basal ganglia, and at two levels of superior cortex.

Investigators measured average attenuation in Hounsfield Units (HU) of circular regions of measurement (0.1-0.25 cm^2) using the method described by Torbey et al, with some modification according to more recent reports [20][21]. We recorded HU values bilaterally for gray matter (GM) in the caudate nucleus (CN), putamen (PU), and white matter (WM) in the corpus callosum (CC), and posterior limb of internal capsule (PIC). In particular we chose the anterior halves of posterior internal capsules in order to min-

imize the density variation of posterior internal capsules between the anterior and posterior halves due to the presence of focal low-attenuation lesions in the posterior half of the posterior internal capsule in 60% of normal brains, according to Choi et al.[22] We recorded values bilaterally for the medial cortex GM and medial WM at the level of the centrum semiovale (MC1 and MWM1, respectively) and high convexity area (MC2 and MWM2, respectively).

Increasing cerebral edema results in lower attenuation by gray matter and a lower GM to WM ratio (GWR). We calculated GWR basal ganglia = $(CN+PU)/(CC+PIC)$. We calculated GWR cerebrum = $(MC1+MC2)/(MWM1+MWM2)$. We calculated average GWR as the mean of the GWR basal ganglia and GWR cerebrum, and we used average GWR for analysis. We divided GWR results into 3 categories, normal (GWR>1.2), mild edema (GWR 1.1-1.2), or severe edema (GWR <1.1), based on prior studies. [23]

3 Statistical analysis

We describe data with mean (SD) for continuous variables, median (IWR) for non-normally distributed variables, and percentages for categorical variables. We compared variables that differed between patients with good and poor outcome using Chi square or Fisher's exact test for categorical variables and Wilcoxon rank-sum test for continuous variables.

We tested a correlation of SSEP N20P25 amplitude measurements between the sides and the inter rater reliability of GWR measurements using Pearson's correlation. We cross-tabulated GWR, EEG reactivity, and SSEP N20P25 amplitude to examine if any particular finding on one test perfectly predicted the results of another test.

We used logistic regression to test associations between average GWR, EEG reactivity, and SSEP N20P25 amplitude at 36°C and 37°C individually and in combination with survival and good outcome at discharge from ICU and after 6 months. Multiple models were created, and the AUC for each compared. With these 3 predictor variables, we created a total of 13 models: 1 model with GWR alone and 6 models at each temperature with SSEP N20P25 amplitude, EEG reactivity, SSEP N20P25+EEG, SSEP N20P25+GWR, EEG+GWR or SSEP N20P25+EEG+GWR.

We use a Hosmer-Lemeshow for goodness of fit for the logistic regression model.

We used a DeLong test to compare the area under the curve (AUC) as each predictor variable was added into subsequent logistic regression models. We used a DeLong comparing AUC both when models included SSEP N20P25 and when the same model was without SSEP N20P25.

Statistical analysis were performed using STATA version 15 (Stata corp, 4905 Lakeway Drive College Station, Texas, USA).

We considered p value≤ 0.05 statistically significant.

3. Results

3.1 Baseline characteristics

Of 171 consecutive patient admitted in our intensive care unit from January 2013 to May 2017, 75 patients were excluded due to missing data, uninterpretable neurophysiology data or because patients died during the treatment. (Picture 2) Table 1 summarizes baseline and resuscitation characteristics for all subjects and subgroups by ICU survival and outcome. Subjects who died were older. Subjects who died or had poor outcome, received higher total dose of adrenaline and had longer collapse-to-ROSC intervals.

Reliability of SSEPs N20-P25 amplitude measurement between the sides was high with a Pearson's correlation coefficient of 0.94. Inter rater reliability of GWR was similarly high with Pearson's correlation coefficient of 0.93.

3.2 Association between GWR, EEG and SSEP

Table 2 and 3 describes the association of GWR, EEG and SSEP features. Only one subject (1%) had severe edema on CT scan (GWR<1.1). This subject had no detectable N20-P25 waveform and no EEG reactivity both at 36 and 37 °C. Among all other subjects, an unfavorable finding in one of these modalities did not perfectly predict unfavorable findings in the others. We measured a strong Pearson correlation coefficient between GWR and SSEP (Pwcorr 0.608). The ROC analysis revealed an AUC for GWRav at discharge of 0.682 (IC 95% 0.551-0.811) for survival with a very high sensitivity and a very low specificity. The AUC was 0.674 (IC 95% 0.574-0.769) at 6 months with a decrease of sensitivity and an increase of specificity. The analysis at 6 months for good outcome demonstrated an AUC of 0.727 (IC 95% 0.629-0.815). Adding SSEP to GWRav to predict survival at discharge, we observed an increase of the AUC up to 0.838 (IC 95% 0.744-0.932) at 37°C with a slight decrease in sensitivity (91%) and an increase in specificity (56%). The results were similar at 36°C [AUC 0.781 (IC 95% 0.681-0.880)]. The combinations of SSEP and GWRav at both temperatures were superior to GWR alone. The predictability was also high for good outcome. [AUC 0.835 (IC 95% 0.755-0.914) at 37°C; AUC 0.823 (IC 95% 0.735-0.908) at 36°C]. (Table 5)

3.3 GWR and Outcome

GWRav ranged from 1.07 to 1.45. GWR was higher in subjects who survived relative to those who did not, and in subjects who had good outcome relative to those who had poor outcome (Table 3). The lowest GWRav value among subjects with a good outcome was 1.16, and among subjects who survived was 1.15. The median amplitude for cortical complex SSEP N20P25 with GWR \geq 1.2 at 36 °C was 0.63 and 0.66 at 37°C. (Table 2)

3.4 EEG and Outcome

EEG reactivity to stimulus at 37°C after rewarming and withdrawal of sedation was present in 41 patients, of whom 31 (69%) survived and 29 (83%) had good outcome at 6 months. There were 6 patients without EEG reactivity at 37°C who subsequently survived with a good outcome at 6 months. (Tab 3)

3.5 SSEP and Outcome

Amplitudes of the N20P25 waveform at 36°C and at 37°C were larger in subjects who survived relative to subjects who did not survive. Amplitudes were also larger in subjects with good outcome relative to subjects with poor outcome (Table 3).

3.6 Prognostic value of combined modality

Table 4 shows the AUC for EEG reactivity, average GWR, and SSEP N20P25 amplitude alone and in combination for predicting good outcome and survival at discharge at 6 months and the survival at discharge. EEG reactivity was the best single predictor of good outcome while N20P25 was the best single predictor for survival at each time point.

Predictive value of each combination of two tests was superior to any test alone. Predictive value of a model including EEG reactivity, average GWR, and SSEP N20P25 amplitude was superior (AUC 0.841 for survival and 0.920 for good outcome) to any combination of two tests or any single test.

3.7. Figures and Tables

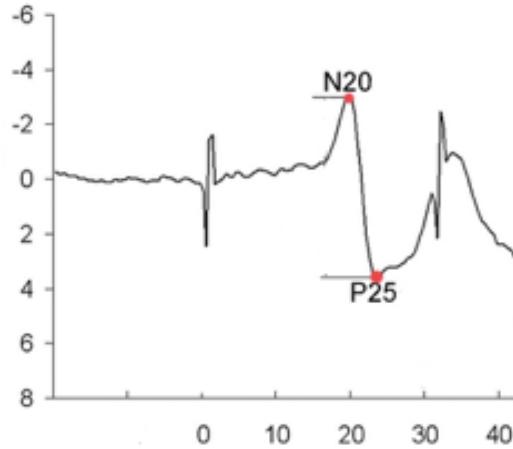


Figure 1. N20P25 cortical complex

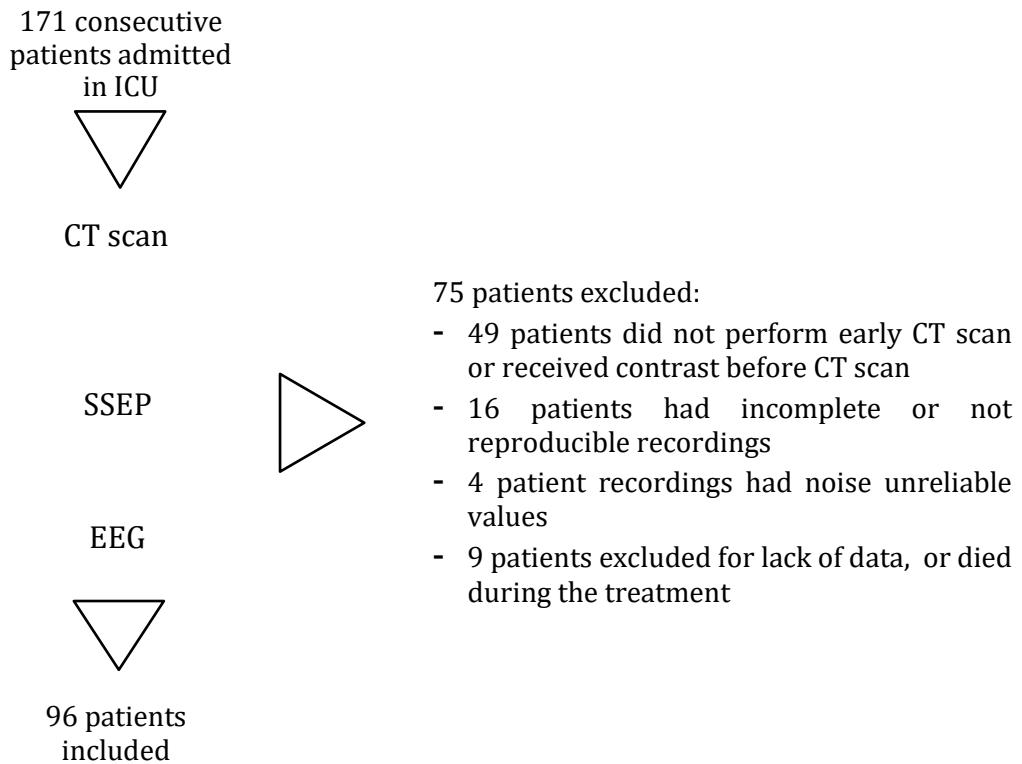


Figure 2 Study design

at discharge**at 6 months**

	<i>tot</i>	<i>survival</i>	<i>non survi- val</i>	<i>p</i>	<i>survival</i>	<i>non survi- val</i>	<i>p</i>	<i>good outcome</i>	<i>bad out- come</i>	<i>p</i>
<hr/>										
number (n, %)	96 (100)	69 (72)	27 (28)		50 (52)	46 (48)		40 (42)	56 (58)	
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age (median, IQR)	62 (53- 71)	61 (50-67)	72 (60-76)	<0.001	60 (48-66)	67 (60-74)	<0.001	61 (50-67)	72 (60-76)	0.01
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Baseline characteristics (n, %)										
Male (%)	69 (72)	49 (71)	20 (74)	0.8	37 (74)	32 (70)	0.6	28 (70)	41 (73)	0.7
Hypertension	23 (25)	14 (22)	9 (33)	0.2	10 (21)	13 (29)	0.4	7 (19)	16 (30)	0.2
Coronary artery disease	13 (14)	7 (11)	6 (22)	0.2	5 (11)	8 (18)	0.3	3 (8)	10 (18)	0.2
Chronic heart failure	8 (9)	5 (8)	3 (11)	0.6	4 (8)	4 (9)	0.9	2 (5)	6 (11)	0.3
Diabetes	21 (23)	12 (19)	9 (33)	0.1	9 (19)	12 (27)	0.4	5 (13)	16 (30)	0.07
Chronic kidney injury	7 (8)	5 (8)	2 (7)	0.2	5 (11)	2 (4)	0.3	4 (11)	3 (6)	0.3
Smoke history	21 (23)	13 (20)	8 (30)	0.3	10 (21)	11 (25)	0.7	8 (22)	13 (24)	0.8
COPD	9 (10)	5 (8)	4 (15)	0.3	3 (6)	6 (14)	0.2	3 (8)	6 (11)	0.6
<hr/>										
Resuscitation details										
Adrenaline mg (mean, IQR)	3 (1-5)	2 (0-4)	4 (2-6)	<0.001	2 (0-3)	4 (2-6)	<0.001	1 (0-3)	4 (2-6)	<0.0001
Shockable rhythm (n, %)	58 (68)	46 (74)	12 (52)	0.05	34 (74)	24 (54)	0.2	28 (76)	30 (62)	0.2

	<i>at discharge</i>				<i>at 6 months</i>							
	<i>tot</i>	<i>survival</i>	<i>non survi- val</i>	<i>p</i>		<i>survival</i>	<i>non survi- val</i>	<i>p</i>		<i>good outcome</i>	<i>bad out- come</i>	<i>p</i>
Time to Rosc minutes (mean, IQR)	17 (12- 26)	15 (10-24)	23 (16-29)	0.01		14 (10-20)	23 (16-30)	0.001		14 (10-17)	22 (16-30)	0.0001
nº of Shocks during CPR (mean, IQR)	2 (0-5)	2 (1-5)	1 (0-5)	0.1		2 (1-5)	2 (0-5)	0.3		2 (1-4)	2 (0-5)	0.8

Table 1. baseline characteristics and resuscitation details

GWR average			
	Severe Edema <1.1	Mild Edema 1.1-1.2	No Edema ≥1.2
<i>n</i> (%)	1(1)	17 (18)	78 (81)
EEG Reactivity at 36°C	0(0)	4 (27)	23 (31)
EEG Reactivity at 37°C	0(0)	4 (25)	37 (51)
N20-P25 amplitude at 36°C, (median, IQR)	0 (0-0)	0 (0-1.13)	0.63 (0-1.73)
If N20-P25 ampl ≠ 0 (median, IQR)	-	1.1 (0.4-1.9)	1.4 (0.7-2.8)
N20-P25 amplitude at 37°C (median, IQR)	0 (0-0)	0 (0-1.98)	0.66 (0-1.88)
If N20-P25 ampl ≠ 0 (median, IQR)	-	2.3 (0.6-2.7)	1.5 (0.7-2.3)

Table 2 Association between GW ratio and SSEP amplitudes and EEG reactivity at different temperature. Ø: no data, -: no observation, GWR average : Gray to white matter ratio average

	at discharge				at 6 months				p	
	tot	survival	non survi- val	p	survival	non survi- val	p	good out- come	bad outcome	
Basal Ganglia	1.25 (1.2- 1.3)	1.27 (1.23- 1.3)	1.22 (1.17- 1.29)	0.03	1.27 (1.23- 1.31)	1.24 (1.19- 1.27)	<0.005	1.28 (1.25- 1.34)	1.23 (1.19- 1.27)	<0.001
Average	1.25 (1.2-1.31)	1.26 (1.22- 1.3)	1.21 (1.17-1.29)	<0.01	1.27 (1.22- 1.32)	1.23 (1.19-1.29)	<0.005	1.29 (1.23- 1.33)	1.23 (1.19- 1.27)	<0.001
SSPEs (median, IQR)										
n20p25 amplitude at 36°C	0.48 (0- 1.47)	0.83 (0-2.16)	0 (0.043)	<0.0005	1.17 (0-2.59)	0 (0-0.73)	<0.0005	1.41 (0.12-2.83)	0.04 (0-0.94)	0.0001
if n20p25 ampl ≠ 0	1.38 (0.7- 2.2)	1.47 (0.8-2.8)	0.6 (0.4-1.4)	<0.01	1.9 (0.8-3.5)	0.93 (0.4-1.39)	0.001	2.2 (1.2-3.9)	0.94 (0.4-1.4)	<0.0005
n20p25 amplitude at 37°C	0.52 (0-1.88)	1.06 (0-2.18)	0 (0-0.16)	< 0.0001	1.49 (0.23-2.33)	0 (0-0.62)	<0.0001	1.66 (0.6-2.36)	0 (0-0.68)	<0.0001
if n20p25 ampl ≠ 0	1.62 (0.7-2.3)	1.8 (0.9-2.4)	0.48 (0.2-0.7)	<0.005	1.93 (1.1-2.6)	0.7 (0.35-1.9)	<0.005	1.98 (1.3-2.6)	0.73 (0.5-1.89)	<0.005

Table 3 Association of GWR, EEG and SSEP features at 36 °C and 37 °C

Discharge from ICU **At 6 months**

	survival		survival		good outcome	
	AUC	(IC 95%)	AUC	(IC 95%)	AUC	(IC 95%)
EEG Reactivity 36°C	0.647	(0.566-0.729)	0.631	(0.540-0.722)	0.658	(0.562-0.754)
EEG Reactivity 37°C	0.764	(0.683-0.845)	0.731	(0.626-0.819)	0.803	(0.699-0.875)
GWRav	0.682	(0.551-0.811)	0.674	(0.574-0.769)	0.727	(0.629-0.815)
N20p25 amplitude 36°C	0.731	(0.629-0.815)	0.707	(0.607-0.797)	0.730	(0.629-0.815)
N20p25 amplitude 37°C	0.775	(0.674-0.850)	0.747	(0.651-0.833)	0.759	(0.662-0.842)
EEG reactivity 36°C GWRav	0.783	(0.674-0.893)	0.745	(0.642-0.848)	0.804	(0.709-0.899)

	Discharge from ICU		At 6 months		
	survival	survival	survival	good outcome	
EEG reactivity 37°C GWRav	0.819 (0.721-0.917)	0.790 (0.697-0.882)	0.872 (0.797-0.947)		
EEG reactivity 36°C N20p25 amplitude 36°C	0.821 (0.730-0.913)	0.781 (0.687-0.874)	0.831 (0.746-0.917)		
EEG reactivity 37 °C N20p25 amplitude 37°C	0.859 (0.778-0.940)	0.812 (0.722-0.902)	0.887 (0.816-0.958)		
GWRav N20p25 amplitude 36°C	0.781 (0.681-0.880)	0.733 (0.680-0.865)	0.823 (0.735-0.908)		
GWRav N20p25 amplitude 37°C	0.838 (0.744-0.932)	0.798 (0.706-0.890)	0.835 (0.755-0.914)		
EEG reactivity 36°C GWRav N20p25 amplitude 36°C	0.845 (0.755-0.936)	0.818 (0.731-0.906)	0.882 (0.812-0.953)		
EEG reactivity 37°C GWRav N20p25 amplitude 37°C	0.882 (0.801-0.963)	0.841 (0.760-0.922)	0.920 (0.864-0.977)		

Table 4. AUC for EEG reactivity, average GWR, and SSEP N20P25 amplitude for predicting good outcome and survival

4. Discussion

We demonstrate that when SSEP cortical complex N20P25 is added into a model with GWR average and EEG reactivity, the predictive value for good outcome and survival at distance is superior than when using each single test alone. Furthermore, the predictive value is high also for survival at discharge from ICU. Few prior studies analysed EEG and SSEP recordings in the same patients and, to the extent of our knowledge, none of them added GWR in the model to predict good outcome and survival at distance.[14][20-23]

Few isolated cases of N20 false positive prediction have been reported. Often a retrospective analysis of the data confirmed that was due to background noise levels or misinterpretation. [24-25] Furthermore, the level of tolerance in noise levels remains still unclear. [7]. In order to overcome the risk that noise misinterpretation could lead to erroneous conclusions of absent N20, our findings suggest using the total amplitude of the SSEPs wave from the lowest value of N20 to the highest of P25. As in a recent paper, we considered the relationship between cortical amplitudes and outcome overcoming the presence absence dichotomy.[26]

SSEPs cortical complex has been previously utilized in other clinical situation for predicting outcome, but no previous studies specifically considered cortical complex N20P25 amplitude as an indicator for good outcome and survival at distance from CA. [27][28] Many papers considered the value of SSEPs only in patients who remained in a comatose state after rewarming, whereas we measured SSEPs in all patients. Although SSEPs are relatively unaffected by sedation, we found, as in previous studies, a difference in amplitude between 36 and 37 °C.[16].

EEG reactivity after withdrawal of sedation at 37 °C was the best solo indicator for good outcome at distance. SSEP complex at 37°C was the best solo predictor for survival at any time. Normally sedation affects more EEG reactivity than SSEP. [29] As shown in Table 4, the three tests combined during hypothermia showed a predictivity for outcomes slightly inferior than after normothermia and off sedation, these results suggest caution using prognostic information in the early stages of treatment.

The minimum amplitude value compatible with a good outcome for the N20-P25 complex was 0.23 μ V. Unfortunately there are few patients with amplitudes very close to this values, making our estimate potentially imprecise. Prospective validation is needed to achieve a higher certainty in this threshold. A retrospective study has previously investigated the relationship between N20-P25 complex and outcome after CA, and the results showed the amplitude reduction of N20-P25 complex were associated with poor outcome. [19]

Although the requisite for amplitude interpretation were well described previously, these data could not consider the difference between cortical and subcortical potentials. Amplitude could be influenced by the intensity and the number of the stimulus, even if we considered a minimum delay after recording. Adding GWR average to SSEPs complex considerably augmented the predictivity for long term outcome [30]. Patients with a hypoxic ischemic encephalopathy manifest a loss of distinction in gray and white matter. Previous studies focused on the presence of cerebral edema in the very early acute period, GWR values are higher when CT scan is performed within the first 6 hours. [31-32] We found only one patient with a GWR value compatible with severe edema (GWR<1.1). In this patient, the corresponding N20-P25 complex amplitude value was zero and there was no reactivity at each temperature we tested. The majority of patients were gathered into the no edema group (GWR \geq 1.2). Early prognostication could lead to a more rapid treatment and eventually referral to rehabilitation programs. Concerning the model for predictivity at 6 months, many factors could influence the survival and the neurological outcome in a so long amount of time. As suggested in other studies, a more extensive evaluation than just CPC score must be considered for recovery evaluation.[33]

There are several limitations to this study. First of all, it is a single center retrospective study, with all the implication that this carries with it. Second, studies on prognostication after CA can suffer from self-fulfilling prophecy. We usually do not consider SSEP in order to withdraw any therapy. Despite this, we can't exclude that single clinicians could be influenced by neurophysiological results. Third, reactivity was determined as a change in EEG background after a pain or voice stimulation, which are not standardized interventions. Although EEG reactivity is considered a strong predictor of awakening, we need to develop standardized methods for testing and interpretations. [34]

5. Conclusions

Our study suggest a strong association between SSEPs N20P25 complex and very early GWR measurement and EEG reactivity to predict good outcome at six months and survival at any time. We predicted high probability of survival and good outcome with GWR>1.15, EEG reactivity present, and N20P25 amplitude >0.83.

The predictability of long terms outcomes could be influenced by several factors. Despite this, the early detection of all these indicators seems to have value for clinical practice.

6. Patents

This section is not mandatory but may be added if there are patents resulting from the work reported in this manuscript.

Supplementary Materials: The following are available online at www.mdpi.com/xxx/s1, Figure S1: title, Table S1: title, Video S1: title.

Author Contributions: Conceptualization, Erik Roman-Pognuz. and Clifton W Callaway; methodology, Johnatan Elmer.; software, Erik Roman-Pognuz and Johnathan Elmer.; validation, Clifton W Callaway , Frank X Guyette, and Erik Roman-Pognuz.; formal analysis, Clifton W Callaway.; investigation, Erik Roman-Pognuz, Alberto Peratoner, Umberto Lucangelo, Gabriele Poillucci and Giorgio Berlot; resources, Erik Roman-Pognuz.; data curation, Erik Roman-Pognuz.; writing—original draft preparation, Erik Roman-Pognuz.; writing—review and editing, Clifton W Callaway, Fabio S Taccone, Tommaso Pellis.; visualization, Erik Roman-Pognuz.; supervision, Erik Roman-Pognuz, Johnatan Elmer, Clifton W Callaway ; project administration, Erik Roman-Pognuz; All authors have read and agreed to the published version of the manuscript.

Funding This research received no external funding

Institutional Review Board Statement: The study was conducted according to the guidelines of the Declaration of Helsinki, and approved by the Ethics Committee Comitato Etico Unico regionale EGAS of Friuli Venezia Giulia region Italy, protocol code 9637 - study number 88/2017 (date of approval March 23 2018).

Informed Consent Statement: Please add “Informed consent was obtained from all subjects involved in the study.” OR “Patient consent was waived due to REASON (please provide a detailed justification).” OR “Not applicable” for studies not involving humans.

Data Availability Statement: Please refer to suggested Data Availability Statements in section “MDPI Research Data Policies” at <https://www.mdpi.com/ethics>.

Acknowledgments: In this section, you can acknowledge any support given which is not covered by the author contribution or funding sections. This may include administrative and technical support, or donations in kind (e.g., materials used for experiments).

Conflicts of Interest: The authors declare no conflict of interest.

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