

STUDY PROTOCOL

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Electroencephalography (EEG) for neurological prognostication after cardiac arrest and targeted temperature management; rationale and study design

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Abstract

Background: Electroencephalography (EEG) is widely used to assess neurological prognosis in patients who are comatose after cardiac arrest, but its value is limited by varying definitions of pathological patterns and by inter-rater variability. The American Clinical Neurophysiology Society (ACNS) has recently proposed a standardized EEG-terminology for critical care to address these limitations.

In the Target Temperature Management (TTM) trial, a large international trial on temperature management after cardiac arrest, EEG-examinations were part of the prospective study design. The main objective of this study is to evaluate EEG-data from the TTM-trial and to identify malignant EEG-patterns reliably predicting a poor neurological outcome.

Methods/Design: In the TTM-trial, 399 post cardiac arrest patients who remained comatose after rewarming underwent a routine EEG. The presence of clinical seizures, use of sedatives and antiepileptic drugs during the EEG-registration were prospectively documented.

After the end of the trial, the EEGs were retrieved to form a central EEG-database.

The EEG-data will be analysed using the ACNS EEG terminology. We designed an electronic case record form (eCRF). Four EEG-specialists from different countries, blinded to patient outcome, will independently classify the EEGs and report through the eCRF. We will describe the prognostic values of pre-specified EEG patterns to predict poor as well as good outcome. We hypothesise three patterns to always be associated with a poor outcome (suppressed background without discharges, suppressed background with continuous periodic discharges and burst-suppression). Inter- and intra-rater variability and whether sedation or level of temperature affects the prognostic values will also be analyzed.

Discussion: A well-defined terminology for interpreting post cardiac arrest EEGs is critical for the use of EEG as a prognostic tool.

The results of this study may help to validate the ACNS terminology for assessing post cardiac arrest EEGs and identify patterns that could reliably predict outcome.

Trial registration: The TTM-trial is registered at ClinicalTrials.gov (NCT01020916).

Keywords: EEG, Prognosis, Interrater variability, Brain injury, Out-of-hospital cardiac arrest, Hypothermia, Resuscitation

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Background

Out of hospital cardiac arrest is a common cause of sudden death. It has a yearly incidence of 40–54 per 100 000 inhabitants [1,2], and an overall survival rate of 3%-10% [3-5]. Comatose survivors are admitted to an intensive care unit (ICU) and approximately half will die during hospital stay. While extensive hypoxic-ischemic brain injury is the main cause of death, severe neurological disability is uncommon among survivors [6].

For patients who remain in coma when sedative drugs have been cleared after rewarming, a neurological examination is the foundation for decisions whether or not to continue intensive care. Prediction of outcome is based on repeated clinical neurological examinations, most often combined with other methods of prognostication [7,8]. Accurate prediction requires test methods with a very high specificity, i.e. a very low false positive rate, for a poor outcome. A recent survey showed that, next to the clinical neurological examination, electroencephalography (EEG) is the most commonly used tool to assess prognosis after cardiac arrest (Friberg et al. Unpublished observation).

EEG is also important for detecting subclinical seizure activity in the sedated and sometimes paralyzed post cardiac arrest patient. Electrographic seizures and postanoxic status epilepticus are associated with a poor prognosis [9-15], but the electrographic definitions vary and there are several reports describing survivors with good neurological outcome [10,12,16,17]. Currently, there is no high-level evidence regarding whether treatment of electrographic seizures or even status epilepticus alters outcome after cardiac arrest. Guidelines on post-arrest care recommend an EEG to rule out seizure activity and for prognostication [18,19].

However, EEG has some important limitations when used in this setting. Firstly, it may be affected by sedation, which is routinely used during temperature management. A decreased body temperature prolongs the metabolism of sedative drugs and it may be difficult to know whether residual sedation still affects the patient or not [20].

Secondly, grading systems and definitions of malignant and benign EEG-patterns vary between studies, which complicate comparisons and meta-analysis. Most grading systems were developed decades ago, before the era of target temperature management [21-25]. In most recent studies, only a limited number of EEG-features were evaluated and it is reasonable to assume that the interpreting EEG-specialists could have been biased by other EEG features. Thus, a standardized terminology evaluating all important features would be preferable to assess which individual features of the EEG are the most predictive.

Thirdly, EEG-specialists usually are confident regarding their own interpretations, but considerable inter-rater variability has been described [26-28]. We are not aware

of any previous studies on the prognostic value of post cardiac arrest EEG including an evaluation of inter- and intra-rater variabilities. Furthermore, most previous studies were performed at single centres by experienced researchers and the generalizability of the results is questionable.

The American Clinical Neurophysiology Society (ACNS) recently proposed a revised version of a standardized critical care EEG terminology [29], to characterize both rhythmic and periodic patterns as well as background activity. The intentions were to standardize terminology of the EEG patterns seen in encephalopathic patients to allow collaborative, multicentre studies and maximize inter-rater reliability.

The Target Temperature Management trial (TTM-trial) randomized 950 comatose cardiac arrest patients to a targeted body temperature of 33°C and 36°C post-arrest and found no significant differences in mortality or neurological function between the two groups [30].

In the TTM-trial, a routine EEG was performed in patients who were still comatose 12 hours after rewarming. Overall, 399 EEGs were performed. This protocol describes the rationale and methods of the EEG-analysis of the TTM-trial.

Aims

We will investigate the prognostic value, as well as intra- and inter-rater variability of specific EEG patterns in a routine EEG performed after rewarming, according to the ACNS standardized EEG terminology.

Our main objective is to identify highly malignant patterns that predict a poor neurological outcome with a zero false positive ratio (FPR) and acceptable inter-rater variability.

Hypotheses

1. For patients who have suffered a cardiac arrest and remain in coma after rewarming, the following EEG features (defined according to ACNS terminology) are:
 - a. always associated with a poor outcome (“highly malignant EEG” with a FPR = 0%):
 - Suppressed background (<10 μ V) without Discharges.
 - Suppressed background (<10 μ V) with Continuous Periodic Discharges.
 - Burst-suppression background (with or without Discharges) with suppression periods (<10 μ V) constituting >50% of the recording.
 - b. nearly always associated with a poor outcome (“malignant EEG” with a FPR < 5%):

Periodic or rhythmic patterns:

 - Abundant Periodic Discharges (>50% of recording).

- Abundant Rhythmic SW (=polyspike-/spike-/sharp-and-wave) (>50% of recording).
 - Unequivocal electrographic seizure (at least one).
- Background EEG:
- Discontinuous background with suppression periods (<10 μ V) constituting >10% of the recording.
 - Low voltage background (most activity <20 μ V).
 - Reversed anterior-posterior gradient.
- Reactivity:
- Absence of background reactivity or only stimulus-induced discharges (SIRPIDs) for both sound and pain stimuli.
- c. associated with a good outcome in the majority of cases ("benign EEG"):
 - Absence of all malignant features stated above.
 2. The prognostic ability of a malignant EEG improves if more than one of the following malignant features (stated in hypothesis 1b) are present in the same EEG:
 - Malignant periodic or rhythmic pattern.
 - Malignant background EEG.
 - Unreactive background.
 3. The prognostic ability of a benign EEG improves if combined with presence of background reactivity for either sound or pain stimuli.
 4. The prevalence and prognostic ability of highly malignant, malignant and benign EEGs after rewarming are similar between patients treated with 33°C or 36°C.
 5. The prognostic ability of highly malignant and malignant patterns is similar between patients that were affected by sedation and those that were not. Influence of sedation is defined as ongoing sedation during the EEG-registration or if the treating physician considered the patient affected by residual sedation on the day of EEG-registration.
 6. There is an acceptable inter- and intra-rater variability (Kappa >0,6) for grouping EEG patterns into the highly malignant, malignant and benign categories stated above and for all the individual features of the ACNS EEG terminology.

Methods/Design

Ethical approval and trial registration

The TTM-trial protocol included a routine EEG and was approved by the ethics committees in all participating countries and institutions. The TTM-trial is registered at ClinicalTrials.gov (NCT01020916) and the trial protocol has been published elsewhere [31].

Recording of EEGs in the TTM-trial

The recording of EEG-data in the TTM-trial was pre-specified [31] and approved by the TTM steering group.

The TTM-trial recruited patients from November 2010 to January 2013. A routine EEG was performed according to the trial protocol in patients that remained unconscious 12 hours after rewarming.

Routine EEGs were performed at all 36 sites (intensive care units in Europe and Australia) in the trial. Sites were given recommendations on how to perform the EEG-registrations including the use of ≥ 16 EEG-channels and testing of background reactivity for both sound and pain stimuli. The result (EEG-report) of the routine EEG was used at each centre according to local routine. The presence of seizures, sedatives, antiepileptic medication and the level of consciousness were prospectively documented in the TTM-database.

A physician blinded for target temperature level performed a neurological evaluation 72 hours after rewarming and issued a recommendation for the continuation or withdrawal of life-sustaining therapy. In the TTM-trial, persisting coma (Glasgow Coma Scale-Motor 1–2) 72 hours after rewarming in combination with a treatment refractory postanoxic status epilepticus defined by EEG was one of the findings that allowed discontinuation of active intensive care [31].

Study population

Adult patients resuscitated after out-of-hospital cardiac arrest of presumed cardiac cause that were still unconscious on admission to the hospital. The patients were randomized to target temperature management at 33°C or 36°C. Sedation was mandated in both groups until after rewarming, 36 hours after randomisation.

Study design for the EEG analysis

For the EEG analysis, four EEG-specialists (Sweden: *EW*, Denmark: *TWK*, The Netherlands: *AFvR*, and Switzerland: *AOR*) were invited together with members of the steering group, with special interest in neurophysiology, to form the EEG group of the TTM-trial (May 2013).

EEG database

On behalf of the TTM-trial steering group, a central EEG database was created at the department of Clinical Neurophysiology, Skane University Hospital, Lund, Sweden. All 36 site investigators were contacted and the local EEG laboratories were encouraged to provide anonymised routine EEGs in electronic format (EDF+).

All notations in the EEGs were translated to English. Notations that were important for the interpretation of the EEGs were kept, for instance notations concerning artefacts, electrode adjustments, movements of the patient, clinical seizure descriptions and if significant changes in level of sedation occurred during the registration. Information that could obviously bias the EEG-specialists for instance notations about the level of consciousness,

including voluntary movements and clinical reaction to stimuli, and results of cranial nerve examination were removed. All data on gender and age were also removed.

Inclusion criteria

A total of 399 patients had a routine EEG according to the TTM-database, all will be considered for inclusion in this study.

Exclusion criteria

1. EEGs performed before reaching normothermic body temperature.
2. EEGs not yet received from the sites before start of analysing phase three (autumn 2014).
3. EEGs that do not meet the following quality criteria:
 - electronic format compatible with the EEG-specialists EEG-readers.
 - ≥ 8 electrodes + reference electrode + ground electrode.
 - registration time of ≥ 10 minutes.

EEG analysis

The EEGs will be analysed using the American Clinical Neurophysiology Society (ACNS) standardized critical care EEG terminology [29]. The four EEG-specialists will independently perform the EEG-analysis, after studying the terminology and ACNS web based training module (www.acns.org).

First phase (pilot study)

Twenty EEGs, chosen to cover important aspects of the ACNS EEG terminology, were analysed during the autumn 2013 by the four EEG-specialists. The purpose was to evaluate inter-rater variability for all categories of the ACNS EEG terminology. For this pilot study paper case report forms were used to report the EEG-findings. There was moderate agreement on the presence and type of periodic and rhythmic patterns [32].

Second phase

A web based eCRF was created (Experlytics AB) for the EEG analysis. The eCRF contains a limited number of categories from the ACNS EEG terminology. The categories were chosen after evaluating inter-rater variability in the pilot study and clinical relevance (literature study, see Table 1). Ten categories were excluded (typical duration, longest duration, phases, relative amplitude, polarity, onset, symmetry, breach effect, posterior dominant rhythm, variability) due to low kappa values (< 0.4). In addition, these categories were considered less relevant for evaluation of post cardiac arrest EEGs.

In the second phase, 103 EEGs will be analysed independently and in a randomized order by the four EEG-

specialists during 2014. These EEGs represent all EEGs from 8 selected sites, which were chosen as they routinely performed testing of reactivity. The 20 EEGs analysed in the first phase will also be included in the 103 EEGs of the second phase for the purpose of evaluating intra-rater variability. When the EEG analysis of the second phase is completed, we will analyse inter- and intra-rater variability (hypothesis 6).

Third phase

We estimate that an additional 200 EEGs will be included in the third phase. In some of these EEGs, testing of background reactivity will not have been performed.

Outcome measures

All patients were followed-up until 180 days after cardiac arrest. The outcome measures were neurological function and mortality. The hypotheses will be correlated to outcome using the cerebral performance category scale (CPC) [52]. Good outcome is defined as CPC 1–2 at 180 days after cardiac arrest or a best achieved CPC of 1–2 at any time. Poor outcome is defined as best achieved CPC of 3–5 during the first 180 days after cardiac arrest.

Statistical analysis

Descriptive statistics

Prevalence will be presented for all individual features in hypotheses 1a-c as well as grouping the features into “highly malignant EEG”, “malignant EEG”, “benign EEG” as well as subgroups of “malignant EEG”: “malignant periodic or rhythmic patterns”, “malignant background EEG” and “unreactive EEG”. The prevalence will be presented as range of the findings reported by the four interpreters as well as a “majority decision”. The “majority decision” for the findings of an individual EEG means that if two or more interpreters have reported a feature, this EEG will be judged to have this feature. If two interpreters have reported one feature and the two others interpreters have reported another feature, the most malignant feature will be reported if the “malignancy” of the features can be graded (i.e. “suppressed background” is more malignant than “burst-suppression background”). If the features cannot be graded (i.e. “low voltage background” versus “reversed anterior-posterior gradient”) both features will be presented.

Prognostic ability

The evaluation of the prognostic ability of EEG will be presented by specificity, sensitivity and false positive rate (FPR), defined as 1-specificity. Similarly as the prevalence, these values will be given both as a range of findings reported by the four interpreters as well as a “majority decision”. Prognostic measures will be compared between

Table 1 Previous publications describing malignant EEG patterns are presented

Described malignant EEG patterns in previous publications	Features of the ACNS EEG terminology included in the hypotheses of the present study	Number of previous publications:					
		Treated with TTM			Not treated with TTM		
		FPR = 0 PPV = 100	FPR < 5 PPV > 95	FPR > 5 PPV < 95	FPR = 0 PPV = 100	FPR < 5 PPV > 95	FPR > 5 PPV < 95
"Burst-suppr"	Burst-suppr (suppr >50%)	3 [10,33,34]	1 [35]	0	5 [36-40]	1 [41]	0
"Discontinuous background with suppr >1 sec"	Discontinuous background (suppr >10%)	0	0	0	3 [22,39,42]	0	0
"Low voltage background (<20 µV)"	Low voltage background (most activity <20 µV)	1 [33]	1 [10]	0	8 [22,36-42]	0	0
"Alfa coma pattern"	Reversed anterior-posterior gradient	1 [33]	0	0	5 [22,36,37,43,44]	1 [45]	3 [39,46,47]
"PDs on a flat background"	Continuous PDs and Voltage = suppressed (<10 µV)	0	0	0	2 [22,36]	0	0
"Continuous Epileptiform Activity or Electrographic Status Epilepticus (broad definition)"	Abundant PDs or Abundant Polyspike-/Spike-/Sharp-and-Waves	3 [14,33,48]	1 [10]	0	1 [39]	0	1 [38]
"Electrographic seizure activity"	Unequivocal seizures	1 [33]	0	0	0	0	0
"Unreactive background"	Reactivity absent or SIRPIDs only	3 [33,49,50]	0	0	1 [39]	0	1 [51]

Malignant EEG patterns after cardiac arrest described in previously published studies [10,14,22,33-51] are presented according to their predicting values (PPV) and false positive ratio (FPR) and whether target temperature management (TTM) were applied or not. We have listed the features (or combinations of features) of the American Clinical Neurophysiology Society's critical care EEG terminology, that we consider correspond to the previously published malignant patterns, which are chosen to be tested in the hypotheses of this study. They were chosen since at least one previous study reported FPR < 5% or PPV > 95% for corresponding EEG patterns. The patterns were thus considered to have a potential to become markers for poor prognosis. For some studies the described malignant patterns were part of an EEG grading system and therefore the prevalence of these patterns could not be decided. PDs = Periodic Discharges, SIRPIDs = Stimulus-induced rhythmic, periodic, or ictal discharges.

groups using a generalized linear mixed model including values from all four interpreters.

Inter-rater variability

Inter-rater variability (Kappa and percent agreement) will be presented for all individual features of the ACNS EEG terminology as well as grouping the features according to hypotheses 1a-c ("highly malignant EEG"; "malignant EEG"; "benign EEG", "malignant periodic or rhythmic patterns"; "malignant background EEG"; "unreactive EEG").

Sample size

Following the primary hypothesis (1a) we assume a FPR of 0 and the study is planned to include approximately 400 patients among whom 125 had a good neurological outcome. Based on these assumptions a 95%-confidence interval (using Wilson's method) of FPR will be 0-3%.

Discussion

Earlier studies of hypothermia-treated patients found that approximately one third of patients are still comatose 3 days after rewarming [53] and we found the proportion of patients that had an EEG in the TTM-trial (n = 399, 42%) to be close to the expected. Thereby, the 399 EEGs are likely to compose a representative sample of EEGs performed for the purpose of prognostication.

EEG has been used for decades for prognostication after cardiac arrest, but its accuracy as a prognostic method is

limited by lack of consensus on definitions of malignant patterns [54-56] and by inter-rater variability [26-28].

The recently revised standardized critical care EEG terminology by ACNS [29] also includes characterization of the background activity, which was lacking in the earlier version [57]. Thereby, it has become possible to use the terminology to investigate the prognostic significance of different EEG patterns in comatose cardiac arrest patients.

Our main objective is to identify highly malignant patterns with an acceptable level of inter-rater variability that can reliably be used to aid in prognostication and in decisions on whether or not to withdraw life sustaining treatment or continue active care. The terminology was designed to maximize inter-rater reliability, which will also be tested. Previously described malignant EEG patterns were recently reviewed [54-56] and we used the evidence on malignant patterns from these three reviews to design the hypotheses of our study (Table 1).

The much used American Academy of Neurology's (AAN) practice parameters did not consider patients treated with temperature management and listed three EEG-patterns as highly malignant; generalized suppression, burst-suppression with generalized epileptiform activity, and generalized periodic complexes on a flat background; these correspond to the highly malignant patterns in the hypotheses of our study. Since the definition of burst suppression in the ACNS terminology is rather conservative (suppression periods constituting more than 50% of the recording), we hypothesised that burst-

suppression is a highly malignant pattern per se, even without discharges.

It is likely that our restriction of highly malignant patterns to those with a false positive ratio (FPR) of zero will lead to a low sensitivity for detecting poor outcome. In order to increase the sensitivity of EEG to detect poor outcome, we will also test other features of the ACNS terminology that we consider potentially malignant (an expected FPR <5%). We therefore selected patterns with a FPR <5% or a PPV >95% in at least one of the previously published studies (Table 1). The corresponding categories of the ACNS terminology were considered to have a potential to become markers of a poor prognosis and were therefore termed “malignant” in the hypotheses of this study.

We used the CPC categories 3–5 to define poor neurological outcome as was done in the TTM-trial and in accordance with the majority of EEG-studies in recent years [55,56].

EEG is affected by sedation [58]. The TTM-trial registered information about ongoing sedation during the routine EEG and whether the treating physician judged that sedation was still affecting the level of consciousness of the patient. Since the routine EEG was performed after rewarming, sedation is likely to have been tapered for most patients. If some EEG-patterns will prove predictive of outcome despite residual sedation, this would be clinically relevant information. We therefore chose not to exclude EEGs judged to be affected by sedation from our main hypotheses, but to analyse the effect of sedation (considering it a dichotomous variable) separately (hypothesis 5).

Postanoxic status epilepticus was previously described as a malignant pattern [9-13]. Based on our own experience and previously published studies using routine or continuous EEG monitoring, we expect the majority of status epilepticus to have started and thus be detected at the time point of the routine EEG in the TTM-trial [10]. Since a treatment refractory status epilepticus in combination with persisting coma 72 hours after rewarming was one of the findings that allowed discontinuation of active intensive care, the validity of the prognostic value may have been affected by early withdrawal of care, a phenomenon known as self-fulfilling prophecy.

Background reactivity has been described to have important prognostic implications [33,49,50,59] and all sites were recommended to perform reactivity testing for sound and pain stimuli. However, this was not performed in all sites and in addition some EEG software systems could not export notations about stimulations. Therefore, we will only be able to evaluate reactivity in approximately 150 EEGs.

It is of paramount importance to establish a well-defined and comprehensible terminology for interpretation of EEG, especially when EEG is used as a prognostic tool and as a

support for decisions on whether active care should be continued or withdrawn. The ACNS EEG terminology is standardized and available for all EEG-specialists. The results of our study should help establish this terminology for assessment of prognosis in comatose post cardiac arrest patients.

Abbreviations

AAN: American academy of neurology; ACNS: American clinical neurophysiology society; CA: Cardiac arrest; CPC: Cerebral performance category scale; eCRF: Electronic case report form; EDF+: European data format; EEG: Electroencephalography; FPR: False positive ratio; GCS: Glasgow coma scale; NPV: Negative predictive value; PPV: Positive predictive value; SIRPIDs: Stimulus-induced rhythmic, periodic, or ictal discharges; SW: Polyspike-/spike/sharp-and-wave; TTM: Target temperature management.

Competing interests

The authors have no competing interests to report.

Authors' contributions

EW, IR, AOR, AFvR, TWK, JH, NN and TC are members of the EEG group of the TTM-trial. EW, IR, AOR, AFvR, TWK and TC designed the EEG-analyses. SU is the statistician of the EEG-analyses. EW, AOR, AFvR, TWK will perform the retrospective EEG-interpretations. In the TTM-trial: NN is principal investigator; HF is senior investigator; NN, HF, TC, and JH are members of the steering group. EW and TC drafted the manuscript that was critically revised and finally approved by all co-authors.

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References

- Atwood C, Eisenberg MS, Herlitz J, Rea TD: **Incidence of EMS-treated out-of-hospital cardiac arrest in Europe.** *Resuscitation* 2005, **67**(1):75–80.
- Rea TD, Eisenberg MS, Sinibaldi G, White RD: **Incidence of EMS-treated out-of-hospital cardiac arrest in the United States.** *Resuscitation* 2004, **63**(1):17–24.
- Adielsson A, Hollenberg J, Karlsson T, Lindqvist J, Lundin S, Silfverstolpe J, Svensson L, Herlitz J: **Increase in survival and bystander CPR in out-of-hospital shockable arrhythmia: bystander CPR and female gender are predictors of improved outcome. Experiences from Sweden in an 18-year perspective.** *Heart* 2011, **97**(17):1391–1396.
- Kitamura T, Iwami T, Kawamura T, Nitta M, Nagao K, Nonogi H, Yonemoto N, Kimura T, Japanese Circulation Society Resuscitation Science Study G: **Nationwide improvements in survival from out-of-hospital cardiac arrest in Japan.** *Circulation* 2012, **126**(24):2834–2843.
- Stiell IG, Wells GA, Field B, Spaitte DW, Nesbitt LP, De Maio VJ, Nichol G, Cousineau D, Blackburn J, Munkley D, Luinstra-Toohey L, Campeau T, Dagnone E, Lyver M: **Advanced cardiac life support in out-of-hospital cardiac arrest.** *N Engl J Med* 2004, **351**(7):647–656.
- Dragancea I, Rundgren M, Englund E, Friberg H, Cronberg T: **The influence of induced hypothermia and delayed prognostication on the mode of death after cardiac arrest.** *Resuscitation* 2013, **84**(3):337–342.
- Samaniego EA, Persoon S, Wijman CA: **Prognosis after cardiac arrest and hypothermia: a new paradigm.** *Curr Neurol Neurosci Rep* 2011, **11**(1):111–119.
- Oddo M, Rossetti AO: **Predicting neurological outcome after cardiac arrest.** *Curr Opin Crit Care* 2011, **17**(3):254–259.
- Rundgren M, Rosen I, Friberg H: **Amplitude-integrated EEG (aEEG) predicts outcome after cardiac arrest and induced hypothermia.** *Intensive Care Med* 2006, **32**(6):836–842.
- Rundgren M, Westhall E, Cronberg T, Rosen I, Friberg H: **Continuous amplitude-integrated electroencephalogram predicts outcome in hypothermia-treated cardiac arrest patients.** *Crit Care Med* 2010, **38**(9):1838–1844.
- Rossetti AO, Logroscino G, Liaudet L, Ruffieux C, Ribordy V, Schaller MD, Despland PA, Oddo M: **Status epilepticus: an independent outcome predictor after cerebral anoxia.** *Neurology* 2007, **69**(3):255–260.
- Rossetti AO, Oddo M, Liaudet L, Kaplan PW: **Predictors of awakening from postanoxic status epilepticus after therapeutic hypothermia.** *Neurology* 2009, **72**(8):744–749.
- Legriel S, Bruneel F, Sediri H, Hilly J, Abbosh N, Lagarrigue MH, Troche G, Guezennec P, Pico F, Bedos JP: **Early EEG monitoring for detecting postanoxic status epilepticus during therapeutic hypothermia: a pilot study.** *Neurocrit Care* 2009, **11**(3):338–344.
- Wennervirta JE, Ermes MJ, Tiainen SM, Salmi TK, Hynninen MS, Sarkela MO, Hynninen MJ, Stenman UH, Viertio-Oja HE, Saastamoinen KP, Pettilä VV, Vakkuri AP: **Hypothermia-treated cardiac arrest patients with good neurological outcome differ early in quantitative variables of EEG suppression and epileptiform activity.** *Crit Care Med* 2009, **37**(8):2427–2435.
- Bouwes A, van Poppelen D, Koelman JH, Kuiper MA, Zandstra DF, Weinstein HC, Tromp SC, Zandbergen EG, Tijssen MA, Horn J: **Acute posthypoxic myoclonus after cardiopulmonary resuscitation.** *BMC Neurol* 2012, **12**:63.
- Hovland A, Nielsen EW, Kløver J, Salvesen R: **EEG should be performed during induced hypothermia.** *Resuscitation* 2006, **68**(1):143–146.
- Westhall Erik RM, Lilja G, Friberg H, Cronberg T: **Postanoxic status epilepticus can be identified and treatment guided successfully by continuous electroencephalography: a case report.** *Ther Hypothermia Temp Manag* 2013, In Press.
- Deakin CD, Nolan JP, Soar J, Sunde K, Koster RW, Smith GB, Perkins GD: **European Resuscitation Council Guidelines for Resuscitation 2010 Section 4. Adult advanced life support.** *Resuscitation* 2010, **81**(10):1305–1352.
- Claassen J, Taccone FS, Horn P, Holtkamp M, Stocchetti N, Oddo M, Neurointensive Care Section of the European Society of Intensive Care M: **Recommendations on the use of EEG monitoring in critically ill patients: consensus statement from the neurointensive care section of the ESICM.** *Intensive Care Med* 2013, **39**(8):1337–1351.
- Tortorici MA, Kochanek PM, Poloyac SM: **Effects of hypothermia on drug disposition, metabolism, and response: a focus of hypothermia-mediated alterations on the cytochrome P450 enzyme system.** *Crit Care Med* 2007, **35**(9):2196–2204.
- Prior PF, Scott DF: **Outcome after severe brain damage.** *Lancet* 1973, **1**(7806):770.
- Scollo-Lavizzari G, Bassetti C: **Prognostic value of EEG in post-anoxic coma after cardiac arrest.** *Eur Neurol* 1987, **26**(3):161–170.
- Synek VM: **EEG abnormality grades and subdivisions of prognostic importance in traumatic and anoxic coma in adults.** *Clinical EEG* 1988, **19**(3):160–166.
- Bassetti C, Karbowski K: **Prognostic value of electroencephalography in non-traumatic comas.** *Schweiz Med Wochenschr* 1990, **120**(39):1425–1434.
- Young GB, McLachlan RS, Kreeft JH, Demelo JD: **An electroencephalographic classification for coma.** *Can J Neurol Sci* 1997, **24**(4):320–325.
- Grant AC, Abdel-Baki SG, Weedon J, Arnedo V, Chari G, Koziorynska E, Lushbough C, Maus D, McSween T, Mortati KA, Reznikov A, Omurtaq A: **EEG interpretation reliability and interpreter confidence: a large single-center study.** *Epilepsy Behav* 2014, **32**:102–107.
- Mani R, Arif H, Hirsch LJ, Gerard EE, LaRoche SM: **Interrater reliability of ICU EEG research terminology.** *J Clin Neurophysiol* 2012, **29**(3):203–212.
- Abend NS, Gutierrez-Colina A, Zhao H, Guo R, Marsh E, Clancy RR, Dlugos DJ: **Interobserver reproducibility of electroencephalogram interpretation in critically ill children.** *J Clin Neurophysiol* 2011, **28**(1):15–19.
- Hirsch LJ, LaRoche SM, Gaspard N, Gerard E, Svoronos A, Herman ST, Mani R, Arif H, Jette N, Minzad Y, Kerrigan JF, Vespa P, Hantus S, Claassen J, Young GB, So E, Kaplan PW, Nuwer MR, Fountain NB, Drislane FW: **American clinical neurophysiology society's standardized critical care EEG terminology: 2012 version.** *J Clin Neurophysiol* 2013, **30**(1):1–27.
- Nielsen N, Wetterslev J, Cronberg T, Erlinge D, Gasche Y, Hassager C, Horn J, Hovdenes J, Kjaergaard J, Kuiper M, Pellis T, Stammen P, Wanscher M, Wise MP, Aneman A, Al-Subaie N, Boesgaard S, Bro-Jeppesen J, Brunetti I, Bugge JF, Hingston CD, Juffermans NP, Koopmans M, Køber L, Langørgen J, Lilja G, Møller JE, Rundgren M, Rylander C, Smeid O, et al: **Targeted temperature management at 33 degrees C versus 36 degrees C after cardiac arrest.** *N Engl J Med* 2013, **369**(23):2197–2206.
- Nielsen N, Wetterslev J, Al-Subaie N, Andersson B, Bro-Jeppesen J, Bishop G, Brunetti I, Cranshaw J, Cronberg T, Edqvist K, Erlinge D, Gasche Y, Glover G, Hassager C, Horn J, Hovdenes J, Johnsson J, Kjaergaard J, Kuiper M, Langørgen J, Macken L, Martinell L, Martner P, Pellis T, Pelosi P, Petersen P, Persson S, Rundgren M, Saxena M, Svensson R, et al: **Target temperature management after out-of-hospital cardiac arrest—a randomized, parallel-group, assessor-blinded clinical trial—rationale and design.** *Am Heart J* 2012, **163**(4):541–548.
- Westhall E, Rosén I, Rossetti AO, van Rootselaar AF, Wesenberg Kjaer T, Horn J, Ullén S, Cronberg T: **Interrater agreement of EEG interpretation in comatose post cardiac arrest patients.** In *Abstract, 30th International Congress of Clinical Neurophysiology, Berlin, 2014.* 2014. Accepted (march 2014) for publication in *Clinical Neurophysiology*, Elsevier.
- Crepeau AZ, Rabinstein AA, Fugate JE, Mandrekar J, Wijidicks EF, White RD, Britton JW: **Continuous EEG in therapeutic hypothermia after cardiac arrest: prognostic and clinical value.** *Neurology* 2013, **80**(4):339–344.
- Kawai M, Thapalia U, Verma A: **Outcome from therapeutic hypothermia and EEG.** *J Clin Neurophysiol* 2011, **28**(5):483–488.
- Oh SH, Park KN, Kim YM, Kim HJ, Youn CS, Kim SH, Choi SP, Kim SC, Shon YM: **The prognostic value of continuous amplitude-integrated electroencephalogram applied immediately after return of spontaneous circulation in therapeutic hypothermia-treated cardiac arrest patients.** *Resuscitation* 2013, **84**(2):200–205.

36. Bassetti C, Bomio F, Mathis J, Hess CW: **Early prognosis in coma after cardiac arrest: a prospective clinical, electrophysiological, and biochemical study of 60 patients.** *J Neurol Neurosurg Psychiatry* 1996, **61**(6):610–615.
37. Rothstein TL, Thomas EM, Sumi SM: **Predicting outcome in hypoxic-ischemic coma. A prospective clinical and electrophysiologic study.** *Electroencephalogr Clin Neurophysiol* 1991, **79**(2):101–107.
38. Zandbergen EG, Hijdra A, Koelman JH, Hart AA, Vos PE, Verbeek MM, de Haan RJ: **Prediction of poor outcome within the first 3 days of postanoxic coma.** *Neurology* 2006, **66**(1):62–68.
39. Edgren E, Hedstrand U, Nordin M, Rydin E, Ronquist G: **Prediction of outcome after cardiac arrest.** *Crit Care Med* 1987, **15**(9):820–825.
40. Lemmi H, Hubbert CH, Faris AA: **The electroencephalogram after resuscitation of cardiocirculatory arrest.** *J Neurol Neurosurg Psychiatry* 1973, **36**(6):997–1002.
41. Young GB, Doig G, Ragazzoni A: **Anoxic-ischemic encephalopathy: clinical and electrophysiological associations with outcome.** *Neurocrit Care* 2005, **2**(2):159–164.
42. Hockaday JM, Potts F, Epstein E, Bonazzi A, Schwab RS: **Electroencephalographic changes in acute cerebral anoxia from cardiac or respiratory arrest.** *Electroencephalogr Clin Neurophysiol* 1965, **18**:575–586.
43. Berkhoff M, Donati F, Bassetti C: **Postanoxic alpha (theta) coma: a reappraisal of its prognostic significance.** *Clin Neurophysiol* 2000, **111**(2):297–304.
44. Vignaendra V, Wilkus RJ, Copass MK, Chatrian GE: **Electroencephalographic rhythms of alpha frequency in comatose patients after cardiopulmonary arrest.** *Neurology* 1974, **24**(6):582–588.
45. Kaplan PW, Genoud D, Ho TW, Jallon P: **Etiology, neurologic correlations, and prognosis in alpha coma.** *Clin Neurophysiol* 1999, **110**(2):205–213.
46. Chokroverty S: **"Alpha-like" rhythms in electroencephalograms in coma after cardiac arrest.** *Neurology* 1975, **25**(7):655–663.
47. Grindal AB, Suter C, Martinez AJ: **Alpha-pattern coma: 24 cases with 9 survivors.** *Ann Neurol* 1977, **1**(4):371–377.
48. Rittenberger JC, Popescu A, Brenner RP, Guyette FX, Callaway CW: **Frequency and timing of nonconvulsive status epilepticus in comatose post-cardiac arrest subjects treated with hypothermia.** *Neurocrit Care* 2012, **16**(1):114–122.
49. Rossetti AO, Oddo M, Logroscino G, Kaplan PW: **Prognostication after cardiac arrest and hypothermia: a prospective study.** *Ann Neurol* 2010, **67**(3):301–307.
50. Rossetti AO, Carrera E, Oddo M: **Early EEG correlates of neuronal injury after brain anoxia.** *Neurology* 2012, **78**(11):796–802.
51. Zhang Y, Su YY, Haupt WF, Zhao JW, Xiao SY, Li HL, Pang Y, Yang QL: **Application of electrophysiologic techniques in poor outcome prediction among patients with severe focal and diffuse ischemic brain injury.** *J Clin Neurophysiol* 2011, **28**(5):497–503.
52. Jennett B, Bond M: **Assessment of outcome after severe brain damage.** *Lancet* 1975, **1**(7905):480–484.
53. Cronberg T, Rundgren M, Westhall E, Englund E, Siemund R, Rosen I, Widner H, Friberg H: **Neuron-specific enolase correlates with other prognostic markers after cardiac arrest.** *Neurology* 2011, **77**(7):623–630.
54. Wijdicks EF, Hijdra A, Young GB, Bassetti CL, Wiebe S: **Practice parameter: prediction of outcome in comatose survivors after cardiopulmonary resuscitation (an evidence-based review): report of the Quality Standards Subcommittee of the American Academy of Neurology.** *Neurology* 2006, **67**(2):203–210.
55. Sandroni C, Cavallaro F, Callaway CW, D'Arrigo S, Sanna T, Kuiper MA, Biancone M, Della Marca G, Farcomeni A, Nolan JP: **Predictors of poor neurological outcome in adult comatose survivors of cardiac arrest: a systematic review and meta-analysis. Part 2: patients treated with therapeutic hypothermia.** *Resuscitation* 2013, **84**(10):1324–1338.
56. Sandroni C, Cavallaro F, Callaway CW, Sanna T, D'Arrigo S, Kuiper M, Della Marca G, Nolan JP: **Predictors of poor neurological outcome in adult comatose survivors of cardiac arrest: a systematic review and meta-analysis. Part 1: patients not treated with therapeutic hypothermia.** *Resuscitation* 2013, **84**(10):1310–1323.
57. Hirsch LJ, Brenner RP, Drislane FW, So E, Kaplan PW, Jordan KG, Herman ST, LaRoche SM, Young B, Bleck TP, Scheuer ML, Emerson RG: **The ACNS subcommittee on research terminology for continuous EEG monitoring:**

proposed standardized terminology for rhythmic and periodic EEG patterns encountered in critically ill patients. *J Clin Neurophysiol* 2005, **22**(2):128–135.

58. Sessler CN, Grap MJ, Ramsay MA: **Evaluating and monitoring analgesia and sedation in the intensive care unit.** *Crit Care* 2008, **12** Suppl 3:S2.
59. Oddo M, Rossetti AO: **Early multimodal outcome prediction after cardiac arrest in patients treated with hypothermia.** *Crit Care Med* 2014, doi:10.1097/CCM.0000000000000211.

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