Prognostic Value of Resting-State EEG Structure in Disentangling Vegetative and Minimally Conscious States: A Preliminary Study

Andrew A. Fingelkurts, PhD 1,*, Alexander A. Fingelkurts, PhD 1, Sergio Bagnato, MD, PhD, 2,3 Cristina Boccagni, MD 2,3, and Giuseppe Galardi, MD 2,3

1 BM-Science – Brain and Mind Technologies Research Centre, Espoo, Finland
2 Neurorehabilitation Unit, Rehabilitation Department, Fondazione Istituto “San Raffaele - G. Giglio”, Cefalu (PA), Italy
3 Neurophysiology Unit, Rehabilitation Department, Fondazione Istituto “San Raffaele - G. Giglio”, Cefalu (PA), Italy

Abstract

Background: Patients in a vegetative state pose problems in diagnosis, prognosis and treatment. Currently, no prognostic markers predict the chance of recovery, which has serious consequences, especially in end-of-life decision-making.

Objective: We aimed to assess an objective measurement of prognosis using advanced electroencephalography (EEG).

Methods: EEG data (19 channels) were collected in 14 patients who were diagnosed to be persistently vegetative based on repeated clinical evaluations at 3 months following brain damage. EEG structure parameters (amplitude, duration and variability within quasi-stationary segments, as well as the spatial synchrony between such segments and the strength of this synchrony) were used to predict recovery of consciousness 3 months later.

Results: The number and strength of cortical functional connections between EEG segments were higher in patients who recovered consciousness (P < .05 – P < .001) compared with those who did not recover. Linear regression analysis confirms that EEG structure parameters are capable of predicting (P = .0025) recovery of consciousness 6 months post-injury, whereas the same analysis failed to significantly predict patient outcome based on aspects of their clinical history alone (P = .629) or conventional EEG spectrum power (P = .473).

Conclusions: The result of this preliminary study demonstrates that structural strategy of EEG analysis is better suited for providing prognosis of consciousness recovery than existing methods of clinical assessment and of conventional EEG. Our results may be a starting point for developing reliable prognosticators in patients who are in vegetative state, with the potential to improve their day-to-day management, quality of life, and access to early interventions.

Key words
Vegetative state; minimally conscious state; EEG; brain damage; brain synchrony; functional connectivity.

Abbreviations:
EEG = electroencephalogram; fMRI = functional Magnetic Resonance Imaging; ISS = Index of Structural Synchrony; LCF = Levels of Cognitive Functioning; LTG = lamotrigine; MCS = minimally conscious state; PB = phenobarbital; TBI = traumatic brain injury; VS = vegetative state.
Introduction

The vegetative state (VS) or unresponsive wakefulness syndrome (UWS) is a complex neurological condition of “unawareness of self and environment in which the patient breathes spontaneously, has a stable circulation, and shows cycles of eye closure and opening which may simulate sleep and waking”.¹ Thus, VS/UWS is commonly agreed to be a state of “wakeful unconsciousness”;² whereas the type of consciousness researchers have in mind is phenomenal consciousness – the sort of awareness that there is something that it is like to enjoy, from the subject’s point of view.³ According to the Royal College of Physicians,¹ a VS/UWS is classified as persistent when it lasts longer than a month with recovery rates approaching zero after 12 months for patients older than 40 years.⁴ Non-traumatic brain injuries are considered to have a poorer prognosis.⁵

Once VS/UWS is diagnosed, the chance of recovery is considered to be low-to-moderate and almost always involves some level of disability;² sometimes VS/UWS lasts the whole life.⁵ In any case, giving a precise estimate of the likelihood of further recovery remains difficult.

The first stage of recovery is characterised by a minimally conscious state (MCS), which is “a condition of severely altered consciousness in which minimal but definite behavioural evidence of self or environmental awareness is demonstrated. In MCS, cognitively mediated behaviour occurs inconsistently, but is reproducible or sustained long enough to be differentiated from reflexive behaviour”.⁶ Like VS/UWS, the MCS may be transitory and precede further recovery of consciousness or last the whole life.²

An accurate and reliable judgment of VS/UWS patients’ awareness is of paramount importance for their diagnosis and prognosis.⁷ Despite an increasing number of published scientific research in recent years, at present, we still do not have objective and validated prognostic markers that allow clinicians to predict the chances of recovery in VS/UWS patients.⁸ Wrong estimation of the chances of recovery can lead to serious consequences, especially when end-of-life decision-making is concerned.⁹ Therefore, objective measurement tools through which an individual’s level of retained awareness can be assessed (without explicit reports) are needed in order to achieve more accurate estimates of prognosis.

Potential Role of Electroencephalogram
An electroencephalogram (EEG), which permits bedside assessment, could be particularly helpful since, in contrast to fMRI, it is routinely available in most clinics/laboratories, affordable, and allows direct and objective recording of spontaneous brain activity without the need of any behavioural response from the patient. More specifically, EEG measures a highly organized macro-level electrophysiological phenomena in the brain, which capture the operations of large-scale cortical networks (neuronal assemblies) and which are remarkably correlated with behaviour, cognition and consciousness.

Despite compelling evidence demonstrating the usefulness of standard resting-state EEG in predicting recovery from coma, there are only scarce reports that show some promise of predicting VS recovery or overall survival. Furthermore, the predictive value of individual resting-state EEG classifications has not been adequately addressed. As a result, conventional resting-state EEG (based on spectral analysis) is typically used only for gross and qualitative analysis and is not practical for long-term patient monitoring nor as a sophisticated prognostic tool.

An alternative strategy to examining EEG is offered by micro-structural analysis of the signal. Using this strategy, recent studies have established that each EEG signal (channel) within the multichannel EEG recording could be represented as a sequence of quasi-stationary (nearly stable) segments. Throughout the duration of each segment, the neuronal assembly that generates the oscillation is supposed to be in steady, quasi-stationary state. The transition of one EEG segment to another reflects change in neuronal assembly microstate. Temporal synchronisation of quasi-stationary EEG segments among different EEG channels could thus represent synchronised activity of different local and transient neuronal assemblies that play an important role in cognition and consciousness. This strategy of EEG analysis has been validated in a number of electrophysiological, cognitive, and clinical studies and is proven to be robust, consistent and statistically reliable.

In applying microstructural strategy of resting-state EEG analysis to VS/UWS and MCS patients, it has been suggested that the incapacity of VS/UWS patients to generate consciousness is most likely linked to disruptions in local and large-scale EEG structures. In particular, it has been found that the absence of consciousness in VS/UWS patients is paralleled by impairment in (a) EEG segmental characteristics (small amplitude, short duration and high amplitude variability) and (b) temporal synchronicity among EEG segments obtained from different EEG channels (decreased number and strength of functional connections). At the same time, fluctuating (minimal) awareness in MCS patients
was paralleled by partial restoration of EEG segmental characteristics (increased amplitude and duration as well as decreased amplitude variability) and their temporal synchrony parameters (increased number and strength of functional connections), approaching those found in healthy fully-conscious subjects. These findings are summarised schematically in Figure 1.

**Aim of the Study**

Here, we investigated whether the aforementioned characteristics of resting-state EEG segments (amplitude, duration and variability), as well as the spatial synchrony of EEG segments and the strength of such synchrony could predict the recovery of conscious awareness after 6 months post-injury in a sample of fourteen patients who were diagnosed to be persistently vegetative on the basis of repeated and thorough clinical evaluations within 3 months following brain damage.
Methods

Patients Cohort

We recruited patients with traumatic and non-traumatic brain injury admitted in Neurorehabilitation Unit at the Fondazione Istituto “San Raffaele – G. Giglio” who met the currently accepted international definition of persistent VS/UWS.1,5,30 Additionally, the Levels of Cognitive Functioning (LCF) scale score31 was assessed on the day of admission (about 3 months post-injury) and three days later when an EEG was registered, to estimate the stable expression of clinical consciousness.21,22 The LCF has a linearly graded scale ranging from 1 to 8 (1 – patient is unconscious; 8 – patient is self-oriented and conscious of the environment) and is well correlated with resting-state EEG abnormalities in patients with brain damage.21,22 At the time of EEG scanning, all patients had a LCF score of 1 or 2 (1.6 ± 0.5).

Inclusion criteria for the patients included (a) confirmation of diagnosis of VS/UWS according to the diagnostic criteria;1,5,30 (b) within 3 months after acute brain event onset; (c) first-ever acute brain event; and (d) stable LCF score during 3 days. Exclusion criteria comprised (a) any acute comorbidity or unstable vital signs; (b) obvious communicating or obstructive hydrocephalus; (c) a history of neurological disease before admission; and (d) severe spasticity (causing constant EMG artefacts). Demographic information is summarised in the Table 1.

Based on clinical evaluations at 3-months follow-up after the EEG registration (thus 6 months post-injury), patients were retrospectively divided into two groups: 10 unrecovered – continued to be vegetative (VS-Pers patients; age 44.7 ± 22 years; follow-up LCF score: 1.7 ± 0.7) and 4 recovered – classified as minimally conscious (VS-MCS patients; age 38 ± 17.3 years; follow-up LCF score: 5 ± 2.4). Though patients with severe brain damage have recovery chances within a year after injury2,4 and therefore some patients in the unrecovered group at 6 months post-injury might in fact recover some level of consciousness one year post-injury; as it is evident from the Table 1, none of the VS-Pers patients recovered consciousness after one year and three patients died. Hence, we used for our analysis the 6 month post-injury assessment as a cut-off point in time for prediction of recovery of consciousness.

The study was approved by the local institutional Ethics Committee and complies with Good Medical Practice. Informed and overt consent of patients’ legal representatives, in line
with the Code of Ethics of the World Medical Association (Declaration of Helsinki) and standards established by the Fondazione Istituto “San Raffaele – G. Giglio” Review Board were acquired. Data use was authorized by means of written informed consent of the VS/UWS patients’ caregivers.

EEG Registration

Waking resting EEG was recorded (0.5–70 Hz bandpass; 200 Hz sampling rate; ~30 min) with a Neuropack (Nihon Kohden, Japan) from 19 electrodes positioned according to the International 10–20 system. The impedance was below 5 kΩ. An electrooculogram (0.5–70 Hz bandpass) was also collected.

EEG recordings were started if patients had their eyes open spontaneously, the eyelids were then closed by hand and kept closed until the end of registration. At the end of the recordings all patients opened their eyes spontaneously, suggesting an unchanged vigilance level throughout EEG registration. The presence of an adequate EEG-signal was determined by visual inspection of the raw signal.

### Table 1. Basic demographic and clinical characteristics of patients

<table>
<thead>
<tr>
<th>Patient ID</th>
<th>Age</th>
<th>Gender</th>
<th>Type of condition</th>
<th>Disorder</th>
<th>CT/MRI findings (in the acute phase)</th>
<th>Time (in days)</th>
<th>LCF recording</th>
<th>LCF after EEG recording</th>
<th>LCF &amp; awakening</th>
<th>LCF after awakening</th>
<th>LCF after awakening</th>
<th>LCF in acute</th>
<th>LCF at 6 months post-injury</th>
<th>LCF at 12 months post-injury</th>
<th>LCF at 6 months post-injury</th>
<th>LCF at 12 months post-injury</th>
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</thead>
<tbody>
<tr>
<td>001</td>
<td>36</td>
<td>M</td>
<td>VS/UWS Trauma</td>
<td></td>
<td>left parieto-temporal intraparenchymal hemorrhage; several intraparenchymal micro-hemorrhages; subdural and epidural hematoma in the right hemisphere; widespread intraparenchymal microhemorrhages</td>
<td>36</td>
<td>PB 100</td>
<td>Delta, Theta1</td>
<td>2</td>
<td>2</td>
<td>Not recov.</td>
<td>Not recov.</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>002</td>
<td>35</td>
<td>M</td>
<td>VS/UWS Trauma</td>
<td></td>
<td>cortical contusions in the frontal lobes and in the right temporal lobes; subdural hematoma; diffuse axonal injury</td>
<td>46</td>
<td>None</td>
<td>Delta</td>
<td>1</td>
<td>1</td>
<td>Not recov.</td>
<td>Not recov.</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>003</td>
<td>28</td>
<td>M</td>
<td>VS/UWS Trauma</td>
<td></td>
<td>subdural hematoma in the left hemisphere; widespread intraparenchymal microhemorrhages in the right</td>
<td>63</td>
<td>None</td>
<td>Delta, Theta1</td>
<td>2</td>
<td>3</td>
<td>Recovered</td>
<td>Recovered</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
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<td>55</td>
<td>M</td>
<td>VS/UWS Trauma</td>
<td></td>
<td>subdural hematoma in the left hemisphere; widespread intraparenchymal microhemorrhages in the right</td>
<td>48</td>
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<td>Delta, Theta1</td>
<td>2</td>
<td>3</td>
<td>Recovered</td>
<td>Recovered</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>005</td>
<td>14</td>
<td>M</td>
<td>VS/UWS Trauma</td>
<td></td>
<td>intraparenchymal microhemorrhages in the right frontal, temporal and parietal lobes; diffuse axonal injury</td>
<td>90</td>
<td>PB 100</td>
<td>Delta, Theta1</td>
<td>2</td>
<td>2</td>
<td>Not recov.</td>
<td>Died</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>006</td>
<td>19</td>
<td>M</td>
<td>VS/UWS Trauma</td>
<td></td>
<td>fronto-temporo-parietal intraparenchymal hemorrhage in the left hemisphere</td>
<td>30</td>
<td>None</td>
<td>Delta, Theta1</td>
<td>1</td>
<td>8</td>
<td>Recovered</td>
<td>Recovered</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>007</td>
<td>35</td>
<td>M</td>
<td>VS/UWS Vascular</td>
<td></td>
<td>left subarachnoid hemorrhage and left tempo-parieto-occipital ischemia (due to vasospasm)</td>
<td>36</td>
<td>None</td>
<td>Delta, Theta1</td>
<td>2</td>
<td>2</td>
<td>Not recov.</td>
<td>Not recov.</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>008</td>
<td>41</td>
<td>M</td>
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<td></td>
<td>fronto-temporo-parietal intraparenchymal hemorrhage in the left hemisphere</td>
<td>60</td>
<td>None</td>
<td>Delta, Theta1</td>
<td>1</td>
<td>1</td>
<td>Not recov.</td>
<td>Died</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>009</td>
<td>79</td>
<td>F</td>
<td>VS/UWS Vascular</td>
<td></td>
<td>intraparenchymal hemorrhage in left parieto-occipital region</td>
<td>92</td>
<td>LTG 200, PB 100</td>
<td>Delta, Theta1</td>
<td>2</td>
<td>2</td>
<td>Not recov.</td>
<td>Not recov.</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>010</td>
<td>50</td>
<td>M</td>
<td>VS/UWS Vascular</td>
<td></td>
<td>hemorrhage in the right putamen</td>
<td>65</td>
<td>None</td>
<td>Delta, Theta1</td>
<td>2</td>
<td>6</td>
<td>Recovered</td>
<td>Recovered</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>011</td>
<td>66</td>
<td>M</td>
<td>VS/UWS Vascular</td>
<td></td>
<td>right fronto-temporo-parietal intraparenchymal and subarachnoid hemorrhage</td>
<td>60</td>
<td>PB 100</td>
<td>Delta, Theta1, Theta2</td>
<td>1</td>
<td>2</td>
<td>Not recov.</td>
<td>Not recov.</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>012</td>
<td>57</td>
<td>M</td>
<td>VS/UWS Vascular</td>
<td></td>
<td>brainstem hemorrhage; right microhemorrhages</td>
<td>90</td>
<td>PB 100</td>
<td>Delta</td>
<td>2</td>
<td>2</td>
<td>Not recov.</td>
<td>Not recov.</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>013</td>
<td>16</td>
<td>M</td>
<td>VS/UWS Anoxia</td>
<td></td>
<td></td>
<td>92</td>
<td>None</td>
<td>Delta, Theta1</td>
<td>2</td>
<td>2</td>
<td>Not recov.</td>
<td>Not recov.</td>
<td></td>
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<tr>
<td>014</td>
<td>68</td>
<td>M</td>
<td>VS/UWS Anoxia</td>
<td></td>
<td></td>
<td>63</td>
<td>None</td>
<td>Delta, Theta1</td>
<td>1</td>
<td>1</td>
<td>Not recov.</td>
<td>Died</td>
<td></td>
<td></td>
<td></td>
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</tr>
</tbody>
</table>

M - male; F - female; VS/UWS - vegetative state/unresponsive wakefulness syndrome; LCF - level of cognitive functioning scale; PB - phenobarbital; LTG - lamotrigine; Delta - 1.5–3 Hz, Theta1 - 3.5–4.5 Hz, Theta2 - 5–6.5 Hz
Prior to analysis, each EEG-signal was bandpass-filtered (Butterworth filter of the sixth order) in the alpha (7–13 Hz), beta-1 (15–25 Hz) and beta-2 (25–30 Hz) frequency bands. Phase shifts were eliminated by forward and backward filtering. The mentioned frequency bands were chosen based on our previous study; only these frequency oscillations have shown dynamics consistent with the analytical consciousness model.

**EEG Segmentation**

In short, the adaptive segmentation algorithm can be described in two main stages (see for details Ref 28 and Fig. 2): (1) preliminary identification of the boundaries using automated algorithm that moves a double window screening along each separate EEG channel; (2) selection of actual (real) boundaries based on the steepness of previously detected EEG amplitude changes and Student criteria. Three EEG segment attributes were further estimated: (1) average amplitude within each segment (microvolts); (2) average length of segments (milliseconds); (3) coefficient of amplitude variability within segments (%). These attributes inform about different features of neuronal assemblies: size, life-span and stability.

**Synchronisation of EEG Segments Among EEG Channels**

Index of Structural Synchrony (ISS; for details see Ref 28) estimates synchronization of EEG quasi-stationary segments obtained from different brain locations. In brief, each boundary in one EEG channel (from any pair of EEG channels) was surrounded by a short (ms) “window”. Any boundary from another channel was considered to coincide if it fell within this window (Fig. 2). To arrive at a direct estimate (5 % level) of statistical significance ($P < .05$) of the ISS, computer simulation of boundaries coupling was undertaken based on random shuffling of segments (500 independent trials) for each pair of EEG channels. As a result of this procedure, the stochastic levels of RTP coupling ($\text{ISS}_{\text{stoch}}$), together with the upper and lower thresholds of $\text{ISS}_{\text{stoch}}$ significance (5%) were calculated. Where there is no synchronization, the ISS tends toward zero, whereas positive (higher than upper stochastic level) or negative (lower than low stochastic level) values are indicative of synchronization (coupling of EEG segments is observed significantly more often than expected by chance as a result of random shuffling during a computer simulation) or de-synchronization (coupling of EEG segments is observed significantly less than expected by chance as a result of random shuffling during a computer simulation) respectively. The strength of EEG structural
synchrony is proportional to the actual value of ISS in each pair of EEG channels: the higher this value, the greater the strength of functional connection. The number of connections corresponds to a number of pairs of EEG channels with a statistically valid ISS.

![Figure 2. Schematic presentation of microstructural strategy of EEG analysis. Explanation in the text.](image)

**Conventional Power Spectral Analysis**

For conventional power spectrum estimates artefact-free EEG signals were filtered in the 1–30 Hz frequency range. Individual power spectra were calculated in the range of 1–30 Hz with 0.5-Hz resolution, using a Fast Fourier Transform with a 2-sec Hanning window shifted by 50 data-samples (0.39-sec) for each channel of one-minute EEG. After calculation of EEG short-term power spectra they were averaged within each EEG channel, then across all one-minute EEGs for each patient. Spectral power was integrated within following frequency bands: delta (1.5–3.0 Hz), theta (3.5–6.5 Hz), alpha (7–13 Hz), beta 1 (15–25 Hz), and beta 2 (25.5–30.0 Hz).

**Results**

*Demographic and Clinical Information*
Six of the patients had sustained a traumatic brain injury (TBI), whereas the remaining eight had sustained a non-traumatic brain injury (non-TBI). There were no significant differences between the two groups (VS-Pers – unrecovered and VS-MCS – recovered) at the time of EEG registration in terms of length of time since brain injury (Student $t$-test, $P = .262$), patients’ age (Student $t$-test, $P = .599$), LCF scale score (Student $t$-test, $P = .733$), or hemispheric localisation of brain damage (Chi-square, $P = 1.000$). However, 50% of patients from VS-Pers group were administered medication, while no one from the VS-MCS group was medicated; this difference was statistically significant (Chi-square, $P = .0000001$).

All patients in both groups underwent the same rehabilitation treatment during hospitalization, consisting of daily verticalization in the standing position (30 min) and kinesitherapy (2 hrs), as well as regular bed mobilization and chair-transfers.

**Prognostic Value of EEG Segments’ Attributes**

Consistent with expectations based on our previous study,\textsuperscript{17} we found that average amplitude within EEG quasi-stationary segments and the length of these segments were larger, while the amplitude variability within EEG segments was lower in VS/UWS patients who recovered some level of consciousness 3 months later (thus 6 months post-injury), when compared with unrecovered VS/UWS patients (Fig. 3). This observation was similar for all three frequency bands (alpha, beta-1 and beta-2).

To determine whether the two groups of patients differed significantly, group-EEG segment-attribute averages and respective standard deviations were calculated for the whole pull of correspondent 1-min EEGs. A comparison of the same segment attributes between VS-Pers and VS-MCS groups was performed using Wilcoxon’s $t$-test. We found no evidence of statistically significant differences between the two groups of patients ($\text{Alpha}_{amplitude}, P = .297$; $\text{Alpha}_{length}, P = .364$; $\text{Alpha}_{instability}, P = .179$; $\text{Beta-1}_{amplitude}, P = .811$; $\text{Beta-1}_{length}, P = .189$; $\text{Beta-1}_{instability}, P = .33$; $\text{Beta-2}_{amplitude}, P = .909$; $\text{Beta-2}_{length}, P = .959$; $\text{Beta-2}_{instability}, P = .388$), though such differences have consistent pattern for all studied frequency bands (Fig. 3). Therefore, these results suggest that the studied parameters of EEG segments may indicate a potential predictive trend and do not carry a predictive value.
Figure 3. EEG segment attributes. Data averaged across all 1-min EEG epochs and all EEG channels for each subject within each group: unrecovered (VS-Pers; \(n = 10\)) and recovered (VS-MCS; \(n = 4\)). The mean values of segment attributes indicated by the Y-axis: amplitude within each segment (microvolt); length of segments (milliseconds); coefficient of amplitude variability within segments (%). Lines in the graphs represent the tendency, which is presented by the linear equation and \(R^2\). Alpha – EEG rhythm within 7-13 Hz, Beta-1 – EEG rhythm within 15-25 Hz, Beta-2 – EEG rhythm within 25-30 Hz.
Prognostic Value of Functional Connections and Their Strength

The number and strength of EEG segmental synchrony was assessed using an ISS index (see Methods section and Ref 28 for details). The differences in number and strength of ISS patterns between the two groups (VS-Pers and VS-MCS) were assessed using Wilcoxon’s rank t-test. At first, all statistically valid EEG functional connections were averaged for each group and the whole pull of correspondent 1-min EEGs within nine categories of functional connectivity (shortleft/right, shortanterior/posterior, longleft/right, longanterior/posterior and longinterhemispheric). The same procedure was done separately for the number and strength of these functional connections. Since the absolute number of possible functional connections within each category was different, their per-category percentage was calculated. During the final stage, an average of all nine categories was calculated. Thus, average values of functional connections for the whole cortex were used for further analysis.

Statistically higher values for the number and strength of functional connections were found during the first assessment of VS/UWS patients (3 months post-injury) who showed good outcome (recovered) at 6 months post-injury, in comparison with unrecovered VS/UWS patients (Fig. 4, Alpha number, P = .01; Alpha strength, P = .028; Beta-1 number, P = .007; Beta-1 strength, P = .038; Beta-2 number, P = .038; Beta-2 strength, P = .021). Similar differences were observed in all three (alpha, beta-1 and beta-2) frequency bands (Fig. 4). These results suggest that the number and strength of cortical functional connections recorded at third month following brain damage provide potentially useful information on the outcome of persistent VS/UWS patients 3 months later (thus 6 months post-injury).

To determine whether an increase in the number and strength of cortical functional connections has a true predictive value of patients’ recovery, compared with clinical history parameters, we used a stepwise linear regression analysis. For large-scale parameters of resting-state EEG structure two factors were used for each frequency band (alpha, beta-1 and beta-2): (a) number of functional connections and (b) strength of functional connections. For the clinical history parameters four factors were used: (a) age at time of injury (years), (b) time since injury (days), (c) LCF score and (d) hemispheric localisation of brain damage (left/right). For conventional EEG spectrum power analysis five factors were used: (a) delta frequency band, (b) theta frequency band, (c) alpha frequency band, (d) beta1 frequency band and (e) beta2 frequency band.

Linear regression analysis confirms that large-scale parameters of resting-state EEG structure (number and strength of functional connections) could successfully discriminate
between recovered and unrecovered VS/UWS patient groups ($F = 12.61$, $P = .0025$), while the same analysis failed to significantly predict VS/UWS patient outcome based on aspects of their clinical history alone ($F = 0.245$, $P = .629$) or conventional EEG spectrum power ($F = 0.754$, $P = .473$). These results indicate that in contrast to the parameters of clinical history and conventional EEG analysis, the large-scale parameters of resting-state EEG structure measured 3 months post-injury can predict the recovery outcome of persistent VS/UWS patients 6 months post-injury. In other words, the higher the number and the strength of cortical functional connections in persistent VS/UWS patients, the higher their chance to recover some level of consciousness in future.

![Graph 1: Number of functional connections](image1)

**Figure 4. EEG structural synchrony.** Data averaged across nine connectivity categories for all pairs of EEG channels within each category and all subjects within each group. The Y-axis presents mean values of either number or strength of functional connections ($n = 9$ categories). Alpha – EEG rhythm within 7-13 Hz, Beta-1 – EEG rhythm within 15-25 Hz, Beta-2 – EEG rhythm within 25-30 Hz. Bars represent means ± s.e.m. * $P < .05$ and ** $P < .01$. 
However, the distribution of patients who had sustained a traumatic brain injury (TBI) was not equal in the VS-Pers (30 %, $n = 3/10$) and VS-MCS (75 %, $n = 3/4$) groups (Chi-square, $P = .000001$). This difference may have influenced the results since it is considered that patients with TBI have better prognosis. To check whether the larger ratio of TBI patients in the VS-MCS group was influencing the main result of this study, we repeated the EEG segmental synchrony analysis for TBI patients only ($n = 3$ for VS-Pers and $n = 3$ for VS-MCS). This analysis resulted in the same significant differences between VS-Pers and VS-MCS groups for all studied frequency bands (Alpha$_{number}$, $P = .008$; Alpha$_{strength}$, $P = .045$; Beta-1$_{number}$, $P = .045$; Beta-1$_{strength}$, $P = .045$; Beta-2$_{number}$, $P = .045$; Beta-2$_{strength}$, $P = .045$) as in the full sample groups. Therefore, we could conclude that the increased ratio of TBI patients in VS-MCS group was not affecting our results.

There was another potential confounding factor: 50 % of patients from the unrecovered group (VS-Pers) were treated with phenobarbital (PB) and lamotrigine (LTG), whereas none of the patients from the recovered group (VS-MCS) were administered medications. These treatments may have influenced the EEG structure, giving rise to the observed differences between VS-Pers and VS-MCS groups. However, both drugs either have no effect or may cause increased coupling among EEG channels. In the present study increased EEG synchrony was observed in the non-medicated group, therefore it is unlikely that the PB and LTG treatments affected the result of the study; in fact they may have diminished the discovered statistically significant difference between the groups (such in the analysis of EEG segmental characteristics).

**Discussion**

Prognostic accuracy for patients in persistent VS/UWS poses serious medical and ethical concerns because treatment decisions typically include the possibility of life-support being withdrawn. Currently, prognosis of the outcome is determined primarily through diagnosis (VS/UWS or MCS) and also by aetiology of brain injury (traumatic, vascular, or anoxic) as well as the age of the patient. However, misdiagnoses of VS/UWS are very common and have been shown to be as high as 37–45 % if not using an appropriate behavioural scales.

Here we have demonstrated that certain parameters of large-scale resting-state EEG structure could predict future recovery of some level of consciousness in persistent VS/UWS
patients. Indeed, 29% (4/14) of patients who appeared to be persistently vegetative on the basis of repeated clinical assessment at the time of admission to rehabilitation unit, but recovered consciousness 3 months later (thus 6 months post-injury), had a similar number and strength of cortical functional connections to those found in MCS patients from our previous study. At the same time, 71% (10/14) of persistent VS/UWS patients who did not recover demonstrated an impaired large-scale structure of EEG as predicted based on our previous observations. These results emphasise the modern understanding that intact communication between brain areas, namely, the coherent dynamic binding of operations performed by multiple neuronal assemblies, which are organized within a nested hierarchical brain architecture, is a basic requirement for consciousness. Furthermore, our results extend several lines of evidence on the strong implication of cortical alpha and beta rhythms on human higher functions and consciousness.

We cannot draw any certain conclusions about the inner subjective experiences in this cohort of patients at the time of admission to the rehabilitation unit based solely on EEG structure analysis. However, our results may indicate that at least 4 VS/UWS patients had a resting-state EEG architecture compatible with partial preservation of awareness despite fulfilling the clinical criteria for a diagnosis of persistent vegetative state. To determine whether these patients may be exhibiting partially preserved conscious processing that is though not expressed behaviourally more research is needed and this is an objective for future studies. Currently, we can only state that prognosis for future recovery of consciousness in persistent VS/UWS patients can be determined accurately on the basis of large-scale resting-state EEG structure analysis alone at the time when patients meet all clinical criteria for the persistent VS (they show no signs of awareness). To fully appreciate the value of this result, it is necessary to consider that a clear-cut difference in resting-state EEG segment synchrony between the two groups (recovered and unrecovered) occurred at the early stage when reliable communication with patients could not yet be established and before spontaneous EEG showed significant modifications in the conventional (based on spectral analysis) parameters (see Table 1). Thus, the result of this study demonstrates that microstructural strategy of resting-state EEG analysis is better suited for providing prognosis of consciousness recovery than existing methods of clinical assessment and of conventional EEG.

Even though the findings of this preliminary study are quite promising, the limitation is that data for the analysis are based upon 14 patients only with varied medical conditions,
which is typical for this kind of population. Therefore one should take with caution the above stated conclusion and to confirm the results presented in this paper, future studies that include a larger group of patients are warranted. Additionally, the results of this study are restricted by the fact that predictive value of the structural EEG measures were not compared with other than conventional (based on spectral analysis) measures, e.g. event-related EEG measures, and that such a comparison should be an object of future studies.

**Conclusion**

As EEG (a) is inexpensive, portable, and available in most hospitals, (b) can be performed at the bedside and (c) can be used in patients who have metal implants, the results of this preliminary work could have a direct clinical significance following further validation: they may establish a starting point in the development of reliable early markers for VS/UWS patients prognosis. This could contribute in improving day-to-day management, access to early interventions, and quality of life in such patients.

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**Declaration of Conflicting Interests**

The authors declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

**Author Contributions**

Conceived and designed the study: AnAF, AI AF, SB, GG. Performed the experiments and collected clinical data: SB, CB. Analysed the data: AnAF, AI AF. Interpreted the results: AnAF, AI AF, SB, CB, GG. Wrote the first draft of the manuscript: AnAF, AI AF. Contributed to the writing of the manuscript: SB, CB, GG. Criteria for authorship read and met: AnAF, AI AF, SB, CB, GG. Agree with manuscript results and conclusions: AnAF, AI AF, SB, CB, GG.
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