Impaired effective cortical connectivity in vegetative state:

Preliminary investigation using PET

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ABSTRACT

Vegetative state (VS) is a condition of abolished awareness with persistence of arousal. Awareness is part of consciousness, which itself is thought to represent an emergent property of cerebral neural networks. Our hypothesis was that part of the neural correlate underlying VS is an altered connectivity, especially between the associative cortices. We assessed regional cerebral glucose metabolism (rCMRGlou) and effective cortical connectivity in four patients in VS by means of statistical parametric mapping (SPM) and [18F]fluorodeoxyglucose (FDG) - positron emission tomography (PET). Our data showed a common pattern of impaired rCMRGlou in the prefrontal, premotor and parieto-temporal association areas and posterior cingulate cortex / precuneus in VS. In a next step, we demonstrated that in VS patients various prefrontal and premotor areas have in common that they are less tightly connected with the posterior cingulate cortex than in normal controls. These results provide a strong argument for an alteration of cortical connectivity in VS patients.

INTRODUCTION

Consciousness is thought to represent an emergent property of cortical and subcortical neural networks and their reciprocal projections. Its multifaceted aspects can be seen as expressions of various specialised areas of the cortex that are responsible for processing external and internal stimuli, short- and long-term storage, language comprehension and production, information integration and problem solving, and attention (Posner, 1994). Vegetative state (VS) is, by definition, a disorder of consciousness: arousal, sleep-wake cycles, ventilation and autonomic control persist but external awareness (including all cognitive function and emotion) is abolished. The clinical diagnosis of VS can be made soon after a brain injury. It may be partially or totally reversible or it may progress to a persistent
VS or death. *Persistent* VS is defined as a VS that persists for at least one month after the acute brain injury (ANA Comittee on Ethical Affairs, 1993; The Multi- Society Task Force on PVS, 1994).

The underlying neuroanatomical basis of VS remains poorly understood. Neuropathological studies of patients in persistent VS of nontraumatic origin show widespread cortical damage with involvement of the association cortices in conjunction with the primary and secondary sensory cortices (Kinney and Samuels, 1994b). Previous positron emission tomography (PET) studies, in patients in VS of diverse origin and duration, showed a reduction in metabolism of overall cortex (Levy et al., 1987). On a regional basis, the most profound reduction in metabolic activity was found in the parieto-occipital and mesiofrontal cortices in persistent VS of anoxic origin (De Volder et al., 1990). Neither study did perform statistical parametric mapping analysis.

We tested the hypothesis that patients in VS, independent of the cause or duration of their condition, suffer from an altered connectivity between the associative cortices. Regional cerebral glucose metabolism distribution (rCMRGlu) was assessed in four cases of VS by means of statistical parametric mapping (SPM) and $^{[18]}$Ffluorodeoxyglucose (FDG) - PET. We first localised the common pattern of cortical metabolic impairment in our VS patients and then studied effective cortical connectivity (Friston et al., 1993) in these patients as compared to normal subjects. Given the small number of observations, this paper should be viewed as a preliminary record of an ongoing research.

**MATERIALS AND METHODS**

The study was approved by the Ethics Committee of the University of Liège. Informed consent was obtained for all control subjects. We retrospectively analysed patients scanned on clinical demand.
All patients fulfilled the international criteria for VS: (1) spontaneous eye opening without evidence of awareness of the environment; (2) no evidence of reproducible voluntary behavioural responses to any stimuli; (3) no evidence of language comprehension or expression; (4) intermittent wakefulness and behaviourally assessed sleep-wake cycles; (5) normal cardiorespiratory function and blood pressure control (ANA Committee on Ethical Affairs, 1993; The Multi-Society Task Force on PVS, 1994).

Patient 1

A 40-year old right-handed woman evolved to a vegetative state after a CO intoxication. She showed spontaneous eye opening and preserved pupillary, oculocephalic, corneal and vestibulo-ocular reflexes. Brain magnetic resonance imaging (MRI) performed 14 days after the insult was normal. Electroencephalography (EEG) showed a 6 Hz basal activity with more pronounced slowing on the left parietal regions. Auditory evoked potentials (AEP) were normal. Somesthetic evoked potentials (SEP) of the median nerve showed normal latency and amplitude of P14 and N20 potentials without any late cortical components. FDG-PET was performed 15 days after the insult. After nineteen days the patient regained consciousness and full autonomy. One year after the accident she showed a spastic gait with altered fine motor function, most prominent one the right, a slurred speech and minor short term memory disturbances.

Patient 2

A 40-year old right-handed woman presented a cardiorespiratory arrest due to a severe bronchial asthma and was resuscitated. She remained in a VS with spontaneous eye opening and preserved pupillary, oculocephalic, corneal and vestibulo-ocular reflexes. Brain MRI performed 22 days after the insult showed bilateral hemorrhagic ischemic lesions in the basal
ganglia. EEG showed a nearly isoelectric basal activity with diffuse low amplitude 3 Hz activity. Visual evoked potentials were absent and AEP was normal. FDG-PET was performed 22 days after the insult. Six months after the accident the patient remained in a VS.

Patient 3

A 26-year-old woman was in VS since 5 years after a traumatic head injury. At examination, eyes were spontaneously open and corneal as well as oculo-cephalic reflexes were present. Spontaneous limb movements with left predominance were observed. Myotatic reflexes were increased and plantar responses were extensor. The following investigations were performed contemporary to FDG-PET: cranial MRI, showing severe cortico-subcortical atrophy and a focal lesion in the pons; EEG, showing diffuse 5 Hz slow dysrhythmia; AEP, showing absence of wave V after left stimulation (suggesting a ponto-mesencephalic lesion); and SEP, showing abnormal responses after stimulation of median nerves (N20 potentials absent on the left side and delayed and broadened on the right side).

Patient 4

A 28-year-old woman was admitted in coma after a suicide attempt using insulin and benzodiazepines. A CT scan performed 4 days after admission showed signs of diffuse cerebral oedema. Patient's condition evolved to VS and FDG-PET was performed 1 month after admission. Other investigations included EEG, showing diffuse delta dysrhythmia, and evoked potentials, showing delayed and broadened cortical responses after stimulation of median nerves and optic nerves. The patient regained consciousness 2 weeks after the PET study. Six months after the cerebral insult she showed a spasticity of the left leg, echolalia alternated with periods of mutism and spatiotemporal disorientation. CT scan now showed diffuse cortico-subcortical atrophy and bilateral pallidal lesions.
FDG – PET data of the four patients were retrospectively analysed. Patient 1 and 2 were scanned on a Siemens CTI 951 scanner at the Cyclotron Research Centre (CRC) of the university of Liege, patient 3 and 4 were scanned on a Siemens 953 scanner at the PET/Biomedical Cyclotron Unit of the Erasmus hospital of the 'Université Libre de Bruxelles'. In both centres data were obtained approximately 15-20 mm above the canthomeatal line and in two dimensional mode. A transmission scan was performed to allow a measured attenuation correction. Five to 10 mCi of FDG was injected intravenously.

PET data were analysed using the statistical parametric mapping software (SPM96 version; Welcome Department of Cognitive Neurology, Institute of Neurology, London, UK) implemented in MATLAB (Mathworks Inc., Sherborn, MA). The use of SPM to assess inter-subject (rather than intra-subject) variability is unlikely to alter the relevance of our results given their high degree of significance. Data from each subject were normalised to a standard stereotactic space and then smoothed with a 16 mm full width half maximum isotropic kernel (Friston et al., 1995a).

The control population was age-matched and consisted of drug-free, healthy volunteers of both sexes (n=53, mean age 42 ± 21 years for patient 1 and 2; and n=25, mean age 27 ± 9 years for patient 3 and 4).

The design matrix included the 4 patient’s scans and the 53 and 25 control subject’s scans of both PET Centres. Global flow normalisation was performed by proportional scaling. The analysis identified brain regions where glucose metabolism was significantly lower in the patient than in the respective centre defined control group. The resulting set of voxel values for each contrast, constituting a map of the t statistics [SPM(t)], was transformed to the unit normal distribution SPM(Z) and thresholded at p<0.001 (Z=3.09). The resulting foci were characterised in terms of peak height over the entire volume analysed at a threshold of voxel-
level corrected p < 0.05 (Friston et al., 1995b). A conjunction analysis provided the common pattern in rCMRGlu decreases characterising our four VS patients.

In a second step, we used a psychophysiological interaction analysis to test the hypothesis on altered effective cortical connectivity in VS as compared to normals (Friston et al., 1993). The design matrix included the same scans as described above and took into account group differences in mean levels of glucose consumption. Now the analysis looked for brain regions that experienced a significant difference in reciprocal modulation with/from the cortical regions most attained in VS. It included the most significant peaks of the 16 different Brodmann area's (voxel level corrected p < 0.05) obtained from the first analysis, and assessed the difference in modulation of every of these 16 voxels depending on the condition patient or normal control. Results were significant at voxel level corrected p<0.05 (Z ≥ 3.96).

RESULTS

A common pattern of significantly impaired rCMRGlu in the four VS patients was observed in the left and right middle and superior frontal gyri, the left inferior frontal gyrus, the left inferior parietal lobule, the left middle temporal gyrus, the right superior temporal gyrus, the posterior cingulate cortex / precuneus, and the left pre- and post-central gyri (Figure 1 and Table 1).

We observed a significant difference in effective connectivity between the most significant voxels of the left prefrontal and premotor cortex (BA 6, 8, 9, 10; marked with an asterix in Table 1) and the posterior cingulate cortex (BA 23/31; x=2, y=-32, z=28; Z score 4.66; p corrected = 0.01) in patients in VS as compared to controls (Figure 2). No such result was obtained with right-sided anterior areas.
DISCUSSION

At the time of PET scanning all patients were in a VS according to the international criteria (ANA Comittee on Ethical Affairs, 1993; The Multi- Society Task Force on PVS, 1994). It is important to distinguish between VS and persistent VS, the latter being defined as a VS that has continued or endured for at least 1 month. In the present study, we were interested in the impairments of effective cortical connectivity in VS independent of its duration, outcome, or cause. Patient 1 and 4 recovered from their VS, patient 2 and 3 failed to recover. The aetiology of their condition was different for each case (toxic, anoxic, posttraumatic and metabolic).

The presented data show a significant dysfunction of the prefrontal, premotor and parieto-temporal association areas and posterior cingulate cortex / precuneus as the common neural correlate in our VS patients. The observed pattern of metabolic impairment is in agreement with post-mortem findings in persistent VS where involvement of the association cortices is reported as critical neuroanatomic substrate (Kinney and Samuels, 1994b).

Furthermore, we observed that the left sided anterior regions differently modulate the posterior cingulate cortex in VS patients as compared to controls. Numerous neuropsychological studies in nonhuman primates suggest the role of the posterior cingulate cortex in spatial orientation and memory (Vogt et al., 1992). In addition, clinical studies have implicated the posterior cingulate cortex in anterograde and retrograde amnesia in humans (Valenstein et al., 1987) (Rudge and Warrington, 1993). Functional imaging studies have shown activation of the retrosplenial region (encompassing the posterior cingulate gyrus and the precuneus) in episodic verbal memory retrieval (Shallice et al., 1994), modulation of visual perception by attention (Dolan et al., 1997) and mental imagery (Fletcher et al., 1995). Interestingly, the posterior cingulate cortex is one of the most active cerebral regions in conscious waking (Andreasen et al., 1995; Maquet et al., 1997) and is systematically one of
the least active regions in unconscious or minimally conscious states such as slow wave sleep and rapid eye movement sleep (Maquet et al., 1997; Maquet et al., 1996). These arguments suggest that the posterior cingulate cortex might represent part of the neural network subserving conscious experience.

The choice of the anterior regions included in the psychophysiological interaction analysis was dictated by the current knowledge about cortical afferents to the posterior cingular cortex in mammals. The posterior cingulate cortex of rats, cats, and monkeys is linked by direct neural pathways both to areas with known sensory and motor functions and to areas affiliated to the limbic system. Neuroanatomical tracer studies have shown projections to the posterior cingulate cortex from the prefrontal cortex (areas 9 and 10) and from the premotor areas (area 6 and 8) (Bates and Goldman-Rakic, 1993; Olson and Musil, 1992; Pandya et al., 1981).

As our statistical model included influences from not one but all 16 regions that showed decreased CMRGlut in VS, the observed inferences about the psychophysiological interactions are valid in relation to effective connectivity as defined by Friston et al. (Friston et al., 1997). It seems that the sum of activity in the left prefrontal and premotor regions differentially predicts activity in the posterior cingulate cortex between the four examined patients in VS and their control population. This is in accordance with experiments in nonhuman primates where the functional integrity of the prefrontal cortex and its interactions with modality specific posterior brain regions is considered critically dependant for working memory (Goldman-Rakic, 1988).

No significant impairment in effective connectivity was observed for the right frontal cortex nor for the posterior cortices (BA 39, 40). Neither was any significant loss of effective connectivity observed for the overall cortex from subcortical nuclei (e.g. the thalamus) (Kinney et al., 1994a). This negative result might be due to the small number of patients
studied and to the heterogeneity of the lesions. Further observations are needed to investigate
the respective role of other neural circuitry involving more cortical and subcortical (thalamic)
structures underlying VS.

This report provides the first and preliminary argument suggesting a disturbed
connectivity between distal cerebral areas in VS patients.

REFERENCES

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Figure 1. The common pattern of altered cerebral metabolism characterising vegetative state patients. SPM(Z) thresholded at voxel and cluster level corrected p<0.05 (Z > 3.963), normalised to the stereotaxic space of Talairach (Friston et al., 1995a) and projected on a normalised MRI template.
Figure 2. Localisation of voxels that showed impaired connectivity with the left prefrontal / premotor cortex (Brodmann area 6, 8, 9 and 10; marked with an asterix in Table 1.) in patients in vegetative state as compared to controls. SPM\{Z\} thresholded at voxel and cluster level p<0.05 (Z > 4.25) projected on a normalised MRI template.
Table: Statistical results and localisation of the most significant voxels where rCMRGlux is decreased in patients in vegetative state as compared to their respective control population (voxel level corrected p<0.05). All the voxels were included in the statistical model assessing effective cortical connectivity. Voxels were the sum of activity differentially predicts the activity in posterior cingulate cortex / precuneus are marked with an asterix. Coordinates are defined in the stereotaxic space of Talairach (Friston et al., 1995a).

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<th>Area (Brodman’s area)</th>
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<th>Y</th>
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