

Impaired Desynchronization of Beta Activity Underlies Memory Deficits in People with Parkinson's Disease

Hayley J. MacDonald^{#1,3}, John-Stuart Brittain^{2,3}, Bernhard Spitzer⁴, Simon Hanslmayr^{2,3}, Ned Jenkinson^{1,3}

¹School of Sport, Exercise and Rehabilitation Sciences

²School of Psychology

³Centre for Human Brain Health

University of Birmingham, Birmingham, United Kingdom

⁴Center for Adaptive Rationality, Max Planck Institute for Human Development, Berlin, Germany

#Corresponding author:

Dr Hayley MacDonald

School of Sport, Exercise and Rehabilitation Sciences

The University of Birmingham, Edgbaston, Birmingham B15 2TT, United Kingdom

Phone: +441214147242

Email: h.j.macdonald@bham.ac.uk

Keywords: Parkinson's disease, episodic memory, beta desynchronization, semantic encoding

Running title: Beta desynchronization and memory deficits in Parkinson's

Abstract

There is a pressing need to better understand the mechanisms underpinning the increasingly recognised non-motor deficits in Parkinson's disease. Brain activity during Parkinson's disease is excessively synchronized within the beta range (12–30Hz). However, relatively little is known about how the abnormal beta rhythms impact on non-motor symptoms. In healthy adults, beta desynchronization is necessary for successful episodic memory formation. We investigated whether there was a direct relationship between decreased beta modulation and memory formation in Parkinson's disease. Electroencephalography recordings were made during an established memory-encoding paradigm. Parkinson's participants showed impaired memory strength ($P = 0.023$) and reduced beta desynchronization ($P = 0.014$) relative to controls. Longer disease duration was correlated with a larger reduction in beta desynchronization, and a concomitant reduction in memory performance. These novel results extend the notion that pathological beta activity is causally implicated in the motor and (lesser appreciated) non-motor deficits inherent to Parkinson's disease.

38 Introduction

39 Parkinson's disease (PD) is classified as a movement disorder. However, there is growing
40 recognition that non-motor burdens also significantly impact those suffering with the
41 condition. Non-demented PD patients can experience cognitive difficulties, including long-
42 term memory deficits (for a review see (Raskin, Borod, & Tweedy, 1990; Zgaljardic, Borod,
43 Foldi, & Mattis, 2003) and specifically the ability to recall verbal memory (Cohn,
44 Moscovitch, & Davidson, 2010; Dujardin et al., 2015; Edelstyn et al., 2015).

45 One striking feature of PD demonstrated repeatedly over the last 20 years is that the
46 electrical activity recorded from basal ganglia (BG) networks in people with PD is
47 excessively synchronized within the beta frequency range (12–30Hz) compared to healthy
48 controls. Under normal circumstances beta activity is modulated with voluntary movement,
49 where the amplitude of oscillations (power) in the beta range drops at the onset of movement
50 and rises again at the end. It is suggested that elevated beta is associated with tonic motor
51 state and event-related desynchronization (ERD) within BG networks “allows” movement to
52 take place (Brittain & Brown, 2014; Joundi et al., 2013), and as such the hyper-synchronized
53 beta state seen in PD prevents desynchronization and thus interferes with voluntary
54 movement (Jenkinson & Brown, 2011). Indeed, therapies that reduce the hyper-synchronized
55 activity, such as dopamine replacement therapy (Ray et al., 2008) or deep brain stimulation
56 (Eusebio et al., 2011), also proportionately improve bradykinesia and rigidity (Ray et al.,
57 2008). Interestingly, beta desynchronization can also occur in the absence of motor output
58 during imagined voluntary movements (McFarland, Miner, Vaughan, & Wolpaw, 2000;
59 Miller et al., 2010). However, to date the link between exaggerated beta activity and motor
60 symptoms in PD remains circumstantial and correlative. It therefore remains an unresolved
61 question as to whether pathological beta activity is causal or an epiphenomenon.

Given that beta activity has shown elevated coupling throughout the BG–thalamocortical circuit in PD, and that this coupling has been observed over broad areas of frontal cortex (Litvak et al., 2011), we postulated that the excessive beta seen in PD should interfere with other neural mechanisms that normally operate within these spatial and temporal domains. Identifying such beta dependent processes and demonstrating a deficit of function in PD would provide further evidence that increased beta is responsible for the motor and non-motor symptoms of the disease. Recent experimental evidence suggests a role for beta oscillations in the encoding of explicit long-term memory. Specifically, a greater amount of beta ERD occurs in the left inferior frontal cortex (IFC) during memory formation of words that are subsequently remembered compared with those that are not (Hanslmayr, Spitzer, & Bauml, 2009; Hanslmayr et al., 2011; Meconi et al., 2016; Meeuwissen, Takashima, Fernandez, & Jensen, 2011; Sederberg, Kahana, Howard, Donner, & Madsen, 2003). This relationship is especially strong if the explicit memory strategy requires semantic processing (Hanslmayr et al., 2009). Memory strategies utilizing semantic processing are examples of deep encoding; when people engage with the meaning of the words e.g. put them into the context of a sentence or make a judgment about whether they relate to living/nonliving entities. Conversely, in shallow encoding an individual only engages with the presented items on a superficial and more perceptual level, as opposed to a cognitive level (Craik & Lockhart, 1972). Examples are detecting whether a presented word contains a specific letter, or whether the first and last letters of the word are in alphabetical order (Otten, Henson, & Rugg, 2001). Unlike in deep encoding, beta ERD during shallow encoding is not predictive of memory performance (Hanslmayr et al., 2009). Furthermore, beta ERD is not seen when similar words are deeply encoded but using non-semantic strategies (Fellner, Bauml, & Hanslmayr, 2013). Therefore, it appears that beta desynchronization is specifically driven by the semantic nature of the encoding task. If the explicit motor deficits in PD are a result of increased beta

synchrony in motor areas of the brain, it stands-to-reason that the memory deficits may well be the result of the elevated levels of beta synchrony which prevent the encoding driven ERD required for semantic processing, and memory formation as a result thereof.

Employing a semantic-encoding memory task to investigate the role of pathological beta in PD has several advantages. Firstly, it removes the confound of movement during the beta desynchronization window. Therefore, if a relationship exists between behavior and beta ERD this would argue against impaired beta desynchronization seen in the motor system being an epiphenomenon that merely reflects the paucity of movement in people with PD. Secondly, semantic processing (Gabrieli, Poldrack, & Desmond, 1998) and episodic memory formation (Otten & Rugg, 2001) recruit the *left* IFC. This is important since dynamic modulation of beta has already been shown to be compromised in PD within the cortical-BG network including *right* IFC and subthalamic nucleus (STN) (Brittain et al., 2012; Swann et al., 2011; Swann et al., 2009). Given the coherent beta activity within cortico-BG circuitry (Hirschmann et al., 2011; Litvak et al., 2011) and bidirectional communication (Horschig et al., 2015; Lalo et al., 2008) within these circuits, we would predict that pathological beta would equally affect *left* IFC beta desynchronization and therefore impair episodic memory that recruits semantic encoding strategies. Intriguingly, it has been demonstrated behaviourally that PD patients do show a specific memory deficit when recollecting deep-encoded words, but no deficit in shallow-non-semantic encoding (Cohn et al., 2010). If this specific deficit can be shown to be associated with the inability to sufficiently desynchronize beta activity, it would demonstrate that impaired modulation of beta might underlie at least some of the higher cognitive symptoms associated with the disease. Finally, we have demonstrated a causal relationship between beta power desynchronization in left inferior prefrontal cortex and memory performance in young healthy adults (Hanslmayr, Matuschek, & Fellner, 2014). Elucidating a direct relationship between beta power ERD and episodic

memory performance in PD would therefore strongly argue for a causal role of hyper-synchronized beta oscillations in the symptoms of PD.

Given this background, the current study aimed to determine whether there is a direct relationship between impaired beta ERD and the long-term memory deficits observed in non-demented PD. The study design, hypotheses and analyses were pre-registered (MacDonald H, Jenkinson N, Hanslmayr S. Memory encoding and beta desynchronisation in Parkinson's disease [Internet]. 2016 Available from: <https://osf.io/vb64n/>). We recorded surface electroencephalography (EEG) during an established memory-encoding paradigm to examine beta oscillations in PD patients and healthy controls during deep-semantic and shallow-non-semantic encoding. We hypothesized that PD patients would exhibit impaired memory performance compared to healthy controls following deep-semantic encoding but that there would be no difference in memory performance between groups following shallow-non-semantic encoding. We further hypothesized that PD patients would show reduced beta ERD during deep-semantic encoding compared to healthy controls, but that there would be no difference in desynchronization between groups during shallow-non-semantic encoding.

Results

Participants

Twenty nine adults with PD and 34 healthy control adults with no known neurological impairment were recruited into the study from local PD community groups and research volunteer databases. This pre-registered recruitment target (see <https://osf.io/vb64n/>) was calculated to account for 10 % drop out and that some participants might be unable to adequately perform the memory task (e.g. insufficient number of remembered items) while still being sufficient to detect a large behavioural effect size (Cohn et al., 2010: Experiment 1) and obtain a power of 0.9. Data for 3 control participants were removed due to not being

able to perform the memory task correctly, and for 1 PD participant due to a change in diagnosis. Demographic information for the remaining 31 control and 28 PD participants is provided in Table 1. Patients were at an average disease duration of 6 ± 4 years (range 0.3 – 14) and tested on their normal medications to avoid the confound of exacerbated motor symptoms. See Table 2 for demographic and clinical data for each individual PD participant. All participants were native English speakers, had completed education at secondary or tertiary level, had no history of dementia, had normal or corrected-to-normal vision and completed the Oxford Cognitive Screen Plus questionnaire (Demeyere et al., 2016) as an assessment of global cognitive function. The two groups did not differ with respect to age, global cognitive function, or level of education (all $P > 0.254$). All results are shown as group means $\pm$ standard error.

Behavioural

Memory strength

In the deep-semantic encoding blocks, participants judged whether the presented word was animate i.e. whether it referred to the property of a living entity. In the shallow-non-semantic encoding blocks, participants judged whether the first and last letters of the word were in alphabetical order. These encoding instructions have been used previously to investigate subsequent memory effects (Hanslmayr et al., 2009; Otten & Rugg, 2001). Recognition testing at the end of each block required participants to rate their confidence as to whether a word presented was one encountered during encoding, or was a new word. This recognition stage was used to calculate memory strength.

Normal distributions were confirmed for all behavioural data sets (all $P > 0.423$). A mixed-effects repeated measures (RM) ANOVA on memory strength (d') revealed no main effect of Group ($F_{1,57} = 2.494$, $P = 0.120$) but a main effect of Encoding ($F_{1,57} = 183.499$, $P <$

0.001). Memory performance improved in both groups with the semantic processing strategy associated with deep encoding (2.524 ± 0.105) leading to greater memory strength (d') during recognition testing compared to shallow encoding (1.249 ± 0.057). There was a Group X Encoding interaction ($F_{1,57} = 4.885$, $P = 0.031$, Fig. 1A). One-tailed post-hoc t -tests revealed no difference in memory strength between groups following shallow-non-semantic encoding ($t_{57} = 0.130$, $P = 0.500$) but deep-semantic encoding lead to greater memory strength in control participants (2.739 ± 0.145) compared to PD (2.309 ± 0.153 ; $t_{57} = 2.042$, $P = 0.023$). Although both groups demonstrated memory benefits from the semantic processing required during deep encoding, controls benefited to a greater degree than PD participants.

When controlling for age, disease duration had a specific detrimental effect on mechanisms underlying memory formation when semantic processing was required in deep encoding. A LASSO regression was run for PD participants to correlate disease duration with deep-semantic and shallow-non-semantic memory strength as well as age. Only memory strength in the deep-semantic encoding condition was significantly correlated with disease duration (Fig. 1B, $F_{1,27} = 11.533$, $P = 0.002$, other $P > 0.242$). A similar regression analysis to correlate age and memory strength in controls was not performed as the assumption of normality was violated for age.

Encoding reaction time and accuracy

Reaction times and response accuracies were recorded during the *encoding stage* when participants were responding ‘yes’ or ‘no’ with button presses in response to deep-semantic and shallow-non-semantic judgements.

For reaction time, a mixed-effects RM ANOVA produced a main effect of Encoding ($F_{1,55} = 6.430$, $P = 0.014$) but no effect of Group ($F_{1,55} = 1.289$, $P = 0.261$) or Encoding X Group interaction ($F_{1,55} = 0.764$, $P = 0.386$). For both groups, reaction time was faster in shallow-

non-semantic encoding (1.12 ± 0.03 s) compared to deep-semantic encoding (1.17 ± 0.03 s) by an average of 50 ms. Similarly, for accuracy, there was a main effect of Encoding ($F_{1,55} = 139.156$, $P < 0.001$) but no effect of Group ($F_{1,55} = 0.044$, $P = 0.834$) or Encoding X Group interaction ($F_{1,55} = 0.119$, $P = 0.732$). Accuracy was higher in shallow-non-semantic encoding (90.9 ± 1.0 %) compared to deep-semantic encoding (75.2 ± 1.1 %) for both groups as expected. The lack of any main effects or interactions with group indicate the significant difference in memory strength between groups in the deep-semantic condition is therefore unlikely to be driven by perceptual differences during encoding.

EEG

All EEG analysis and presented data are from the *encoding stage*. EEG data from 1 control and 2 PD participants could not be used due to technical problems or large movements from dyskinesia, leaving 30 control and 26 PD EEG data sets for analysis. In alignment with previous EEG studies, and as per our pre-registered protocol, post-stimulus beta power decreases are expected to be associated with successful memory formation in healthy (Hanslmayr et al., 2009; Hanslmayr et al., 2011) and patient populations (Meconi et al., 2016). Therefore lower beta from 12 – 20 Hz was the main frequency range of interest for all dependent measures (see <https://osf.io/vb64n/>) over 0 – 1.5s relative to stimulus onset (i.e. word presentation).

As hypothesised, the cluster-based permutation testing on all electrodes showed that controls demonstrated greater beta ERD during deep-semantic encoding of subsequently remembered words (Hits) compared to PD participants (cluster stat = -150.1, $P = 0.014$, Fig. 2A & B show beta ERD for electrodes in significant cluster), however no difference between groups emerged during shallow-non-semantic encoding (cluster stat = -3.7, $P = 0.326$, Fig. 2C & D show beta ERD for electrodes in largest cluster that did not reach significance). A

mixed-effects RM ANOVA on averaged beta (over 0 – 1.5 s, 12 – 20 Hz) further supported this finding by producing a significant Encoding X Group interaction ($F_{1,54} = 6.959$, $P = 0.011$) that confirms the difference between groups in deep-semantic encoding ($t_{54} = 2.910$, $P = 0.005$) is significantly different to shallow-non-semantic encoding ($t_{54} = 1.030$, $P = 0.307$). There were no main effects of Encoding ($F_{1,54} = 0.612$, $P = 0.437$) or Group ($F_{1,54} = 3.946$, $P = 0.052$). Therefore, a difference in beta ERD between groups is seen only in the deep-semantic encoding condition, indicating that there is an ERD deficit in the PD group that occurs specifically during deep-semantic processing.

The relationship between beta ERD and the deep-semantic encoding condition is reinforced by the similar pattern of beta ERD seen during the encoding of words that were not successfully remembered (Misses). Misses in controls were associated with greater beta ERD during deep-semantic encoding when compared to PD participants (cluster stat = -54.1, $P = 0.031$), however no difference between groups emerged during shallow-non-semantic encoding (cluster stat = -3.8, $P = 0.330$). A mixed effects RM ANOVA similarly produced main effects of Encoding ($F_{1,54} = 5.450$, $P = 0.023$) and Group ($F_{1,54} = 6.155$, $P = 0.016$) and a significant Encoding X Group interaction ($F_{1,54} = 5.975$, $P = 0.018$). The interaction confirms the difference between groups in deep-semantic encoding ($t_{54} = 3.367$, $P = 0.001$) is significantly different to shallow-non-semantic encoding ($t_{54} = 0.919$, $P = 0.362$). The fact that a difference in beta ERD is seen between groups during encoding of both remembered and forgotten items implies the difference is related to deep-semantic encoding in general. This overall reduced beta desynchronization may lead to reduced memory performance in PD participants.

Successful memory formation specifically involving deep-semantic processing was associated with greater beta ERD. Within groups, controls demonstrated greater beta ERD for subsequently remembered words during deep-semantic compared to shallow-non-semantic

encoding (cluster stat = -94.4, $P = 0.012$, Fig. 2E & F show beta ERD for electrodes in significant cluster). Interestingly at a group level, PD participants did not show significantly greater ERD in deep-semantic encoding compared to shallow-non-semantic (no significant clusters were identified), although they did show a behavioural benefit of deep-semantic encoding, albeit to a lesser extent than controls. Based on findings of left IFC beta being specifically linked to memory strength in healthy controls (Hanslmayr et al., 2009; Hanslmayr et al., 2011; Meeuwissen et al., 2011), we did an additional correlational analysis focusing on left frontal beta in PD patients. Despite no group-level effect, linear regressions illustrated that PD participants who showed greater beta ERD over left frontal electrodes also had significantly greater memory strength during deep-semantic encoding ($P = 0.008$, $R^2 = 0.256$, Fig. 3A) but that disease duration negatively correlated with left frontal maximum beta ERD ($P = 0.007$, $R^2 = 0.263$, Fig. 3B). PD participants earlier in the disease who were able to achieve greater beta ERD in left frontal electrodes benefited more from deep-semantic encoding strategies of memory formation.

The secondary dependent measure was the subsequent-memory effect (SME) in beta power which compared power between high confidence hit (i.e. subsequently strongly remembered) and miss (i.e. subsequently forgotten) trials (Brewer, Zhao, Desmond, Glover, & Gabrieli, 1998; Hanslmayr et al., 2009; Otten et al., 2001). This categorization in the *encoding stage* depended on the participant's response in the *recognition stage* and their individualized receiver operating characteristic (ROC) curves (Hanslmayr et al. 2009; see Materials and Methods). The SME results broadly replicated a number of previous findings (Hanslmayr et al., 2009; Hanslmayr et al., 2011; Meconi et al., 2016) and further support the importance of beta ERD as the mechanism underlying successful memory formation through deep-semantic encoding strategies: there was a significant SME in deep-semantic encoding for controls (cluster stat = -42.2, $P = 0.027$, Fig. 4A & B illustrate beta ERD for electrodes in

significant cluster) and a SME approaching significance for PD participants (cluster stat = -22.3, $P = 0.097$, Fig. 4C & D illustrate beta ERD for electrodes in the largest cluster). Importantly, there was no significant SME associated with shallow-non-semantic encoding (controls: cluster stat = -2.1, $P = 0.698$, Fig. 4E & F illustrate beta ERD for electrodes in largest cluster that did not reach significance; PD: no clusters were identified). A mixed-effects RM ANOVA showed a main effect of Encoding ($F_{1,54} = 24.265$, $P < 0.000$), confirming that deep-semantic encoding produced a greater average SME ($-6 \pm 1\%$) compared to shallow-non-semantic encoding ($1 \pm 0.7\%$). There was no main effect of Group ($F_{1,54} = 0.007$, $P = 0.935$) or Encoding X Group interaction ($F_{1,54} = 0.023$, $P = 0.880$). The lack of an interaction was expected as, although PD participants remembered fewer items than controls following deep-semantic encoding, the remembered items in both groups should be accompanied by similar electrophysiological signatures (i.e. SME) as in both cases they lead to the same behavioural outcome – that of remembering (i.e. d' above zero).

Discussion

The study confirmed our pre-registered hypotheses and produced several novel findings that provide the first evidence of impaired beta modulation being associated with a non-motor symptom of PD. PD participants showed impaired memory strength compared to healthy controls but only following deep-semantic encoding of words. This behavioural finding was mirrored by the EEG results which demonstrated that PD participants exhibited reduced beta ERD compared to healthy controls but again only during deep-semantic memory formation. Furthermore, a correlation between disease duration and an increased deficit in deep-semantic encoding suggested the neuropathology of PD has a specific detrimental effect on the mechanisms underlying deep-semantic memory formation leading to both reduced beta ERD and reduced memory strength. This is reinforced by that fact that participants with PD who

showed greater beta ERD over left frontal electrodes benefited to a greater extent from the deep-semantic encoding memory strategy. There were no differences between the groups in age, global cognitive function, education or perception during encoding that could explain these behavioural or EEG results. Therefore, our results appear to be specific to episodic memory formation as a result of deep-semantic processing. Overall, our findings strengthen the idea that dysfunctional beta oscillations are likely to be the cause of PD symptoms in both motor and non-motor domains.

Parkinson's disease did not cause impaired memory performance in general, but rather a specific deficit in deep-semantic encoding of memory. Deep-semantic encoding in the context of the current study utilized general knowledge about the word to form an abstract representation and evaluate the representation as animate or inanimate. Age-related memory decline is a widely acknowledged fact that is seen across several subdomains, including episodic memory (e.g. see (Shing et al., 2010)). Over and above the aging-related decline, a further decline in episodic memory resulting from deep-semantic encoding appeared to be caused by the mechanisms underlying PD. Replicating previous findings, PD participants were able to employ the non-semantic encoding strategy to build a memory trace of equivalent strength to controls (Cohn et al., 2010). The difference in memory performance between groups was only elucidated following a deep-semantic encoding instruction. In contrast to Cohn and colleagues (Cohn et al., 2010), the current PD participants still showed a behavioural benefit from the deep-semantic encoding memory strategy and those who were less progressed in the disease benefited to a greater degree. People with PD struggle to spontaneously implement the optimal memory encoding strategy (Knoke, Taylor, & Saint-Cyr, 1998). However with explicit encoding instructions, PD participants managed to improve memory with the optimal deep-semantic encoding strategy, albeit to a lesser degree than controls. This finding suggests they are able to recruit the neural mechanisms to process

semantic information about the words in the deep encoding condition, but something prevents the formation of a robust memory trace. Overall, people with PD exhibited a limited deep-semantic processing capacity during memory encoding rather than a general deficit in recognition memory.

The deficit in episodic memory performance following a deep-semantic encoding strategy displayed by PD participants was associated with a reduced dynamic range of beta ERD during encoding. Brain oscillations are considered one of the core neural mechanisms for storage and retrieval of long-term memories (Buzsaki & Draguhn, 2004; Fell & Axmacher, 2011) and the extent of neural desynchronization is thought to relate to the degree of information stored in the brain (Hanslmayr, Staudigl, & Fellner, 2012). In the current study, the greater level of beta desynchronization for deep-semantic versus shallow-non-semantic encoding, and words that were subsequently remembered compared to those that weren't, further supports the importance of beta ERD as the mechanism underlying successful deep-semantic memory formation (Hanslmayr et al., 2009; Hanslmayr et al., 2011; Meconi et al., 2016; Meeuwissen et al., 2011; Sederberg et al., 2007). As both groups displayed similar behavioural outcomes of deep-semantic encoding (i.e. d' values above zero, although PD participants remembered fewer items than controls), it is not surprising that both groups displayed similar electrophysiological differences between high confidence hits and misses (i.e. a SME). Importantly however, overall beta ERD was significantly reduced in PD participants compared to controls during deep-semantic processing, but not for words encoded with a shallow-non-semantic strategy. This distinction implies that a reduced capacity to decrease beta power following stimulus presentation for PD participants reduced the richness of semantic information encoded in the brain and therefore weakened the memory strength, leading to fewer successfully recognized words and a lower d' value.

It has been proposed that the relative change in pre- to post-stimulus power is most important for memory performance, rather than absolute power levels (Klimesch, Doppelmayr, & Hanslmayr, 2006; Klimesch, Sauseng, & Gerloff, 2003). PD participants demonstrated decreases in the reactivity of their event-related beta power and therefore reduced encoding capacity. PD participants who were further progressed in the disease demonstrated further reductions in both beta reactivity and memory strength. A reduced dynamic range of BG-thalamocortical beta power in PD can therefore interfere with other neural mechanisms that operate in the beta frequency range apart from movement, including memory formation.

The neural changes causing episodic memory deficits in PD may be the same as those underlying motor symptoms. Memory formation recruits an extensive network of mainly left-lateralized regions for verbal material. This network includes the anterior temporal lobe for storage of conceptual representations and processing concepts at an abstract level (Jefferies & Lambon Ralph, 2006; Patterson, Nestor, & Rogers, 2007), and the IFC and temporoparietal region for strategic search and control processes that are necessary for semantic processing (Binder, Desai, Graves, & Conant, 2009; Jefferies, 2013; Jefferies & Lambon Ralph, 2006). The extent of beta ERD in left prefrontal cortex (PFC), specifically IFC, has been linked to memory performance (Hanslmayr et al., 2009; Hanslmayr et al., 2011). Function of the PFC is heavily influenced by the integrity of dopaminergic input onto frontostriatal connections. Therefore, it is not surprising that dopaminergic dysfunction seen in PD leads to impaired IFC function, observed in motor tasks that recruit the right IFC as part of the response inhibition network (Bokura, Yamaguchi, & Kobayashi, 2005; Gauggel, Rieger, & Feghoff, 2004; Obeso et al., 2011; Swann et al., 2011). We have extended these findings to also show impairment during a memory task that has been shown to recruit the left IFC during deep-semantic encoding. Previous studies have highlighted the ability of BG oscillatory activity to

influence cortical neuronal oscillations recorded with surface EEG (Chung et al., 2018; Horschig et al., 2015). We therefore propose that the same pathological BG beta mechanism causing the motor symptoms in PD is contributing to the deficit in deep-semantic encoding of memory seen in the current study. This would imply a common neural mechanism may underlie a variety of deficits in PD that involve cortico-BG processes which operate predominantly in the beta frequency range.

It is a matter of speculation as to the cause of altered memory-related beta oscillations within PD. However, there are potential candidate mechanisms that could be contributing to pathological beta within the memory domain. For example, long-term potentiation (LTP) in the hippocampus is proposed as the mechanism of synaptic plasticity playing a key role in the formation of long-term memories (Bliss & Collingridge, 1993). Neural oscillations are thought to shape synaptic plasticity by providing temporal windows for neural firing (Hanslmayr, Staresina, & Bowman, 2016), so the differences in cortical beta oscillations in the current study between PD patients and healthy participants might be linked to LTP-like mechanisms. However, the direct relationship between any one form of synaptic plasticity and a specific frequency range of oscillations or a particular type of memory is still unclear and highly speculative, especially in humans and cortical regions. Intriguingly, LTP-like mechanisms that are altered in the motor areas in people with PD (Kishore, Joseph, Velayudhan, Popa, & Meunier, 2012; Lago-Rodriguez et al., 2016; Suppa et al., 2011) are also suggested to be the mechanism behind the reduced modulation (in PD) of movement-related beta that is normally seen in the sensorimotor area during repetitive practice of arm movements in healthy controls (Moisello et al., 2015; Nelson et al., 2017). Therefore, future studies could investigate whether impaired LTP-like mechanisms are linked to the reduced memory performance and reduced beta modulation seen in PD patients in the current paradigm.

Identifying a common neural mechanism behind the motor and non-motor symptoms of PD has implications for treatment and disease monitoring. There are currently no standard treatment options for mild memory and cognitive problems in PD (i.e. mild cognitive impairment). Applying interventions previously shown to decrease hyper-synchronized beta activity such as deep brain stimulation or dopamine replacement therapy (Eusebio et al., 2011; Ray et al., 2008) should in theory also help with memory deficits caused by the same pathology. Considering the inverse relationship demonstrated in the current study between disease progression and both memory performance and beta ERD, it is feasible that this memory paradigm could be developed as a useful surrogate to measure functional beta reactivity. As such, the paradigm could be used as a new and convenient behavioural test to monitor disease progression, with specific applications in telemedicine.

It is important to note that while we present findings that the neural changes causing episodic memory deficits in PD may resemble those underlying motor symptoms, we do not posit that reduced beta de-synchronisation is the sole deficit that emerges in PD. Nor, in-fact, that there is a single source of beta that homogenises symptomology across domains (Spitzer & Haegens, 2017). Instead, we extend the impact of a deficit that has been identified in the motor domain to other (cognitive) areas. This will likely explain some symptoms well, but not all, and should be a consideration when titrating medications to alleviate different aspects of motor and/or cognitive performance. It is important to make this distinction as we are not claiming that beta observed in the motor system directly influences memory encoding – but that beta in memory-relevant areas is also deficient and, while these rhythms are likely to serve a similar functional role, deficits may indeed be graded across functional areas. Hence, motor deficits and memory deficits may be differentially influenced depending on the underlying pathophysiological state.

There are a few limitations to the current study that should be considered. Firstly, the relationship between beta ERD and the behavioural deficit in the PD group is correlational. However, it is the more parsimonious explanation that a common underlying neurological deficit (i.e. impaired beta desynchronization) causes both motor and memory problems than two unrelated behavioural symptoms producing the same epiphenomenon in the beta system. Furthermore, evidence exists for a causal relationship between the strength of beta desynchronization in left PFC and memory performance (Hanslmayr et al., 2014) so the direct relationship shown in the current study would support a causal role of pathological beta in PD symptomology. Extending the findings from Hanslmayr and colleagues, future studies could use transcranial magnetic stimulation to modulate left prefrontal beta in people with PD and look for a causal influence on their episodic memory performance. Secondly, beta desynchronization also plays a role in memory retrieval (Dujardin, Bourriez, & Guieu, 1994; Duzel et al., 2003) and people with PD are thought to use inefficient retrieval strategies (see (Zakzanis & Freedman, 1999)). However using recognition, which is one of the simplest ways to test episodic memory, greatly reduced retrieval demands in our task, e.g. compared to free or cued recall. A retrieval based explanation for our behavioural findings is therefore rather unlikely. Nevertheless, we cannot completely discount the contribution of impaired beta desynchronization during retrieval to the reduced recognition memory performance in our study. Our prior hypothesis and pre-registered protocol focused initially on memory encoding because encoding primarily recruits the IFC, while retrieval recruits parietal regions (Burgess & Gruzelier, 2000; Spitzer, Hanslmayr, Opitz, Mecklinger, & Bauml, 2009; Zion-Golumbic, Kutas, & Bentin, 2010). Due to the dopaminergic modulation of frontostriatal connections discussed previously, we expected pathological BG beta in PD would preferentially affect prefrontal cortical regions.

Despite displaying topographical maps in an effort to show the location of ERD differences between groups, the methods used in the current study cannot be used to form a robust conclusion about spatial differences in beta ERD. The location of beta ERD differences in deep-semantic encoding between patients and healthy participants seemed to indicate a widespread cortical deficit in beta desynchronization in PD patients, which included the left frontal region. This widespread difference is in contrast to, for example, more focal differences in beta ERD for healthy participants between deep-semantic and shallow-non-semantic encoding. However scalp-level EEG has limited spatial resolution. Subsequent studies using magnetoencephalography with a much higher spatial resolution would be needed to investigate these results further. Finally, when considering the generalizability of our results, it is worth noting that the PD patients in the current study were mild to moderately impaired in terms of disease severity. Our study therefore cannot directly speak to the relationship between memory impairments and beta oscillations in severely affected PD patients. However, our findings of an inverse relationship between disease duration and both memory performance and beta desynchronization speaks to a general characterisation that will likely extend (alongside other age-related factors) to those severely impaired patients.

Conclusion

This study provides the first evidence of impaired beta modulation being associated with a non-motor symptom of PD. PD participants showed impaired memory strength and beta ERD compared to healthy controls during deep-semantic encoding. The neuropathology of PD seemed to have a specific detrimental effect on the mechanisms underlying episodic memory formation in a deep-semantic encoding task leading to both reduced memory strength and reduced beta ERD. We propose that the neural changes causing memory deficits in PD may be the same as those underlying motor symptoms i.e. impaired modulation of beta activity

within BG– thalamocortical circuitry. Importantly the decrease in beta modulation shown in our study cannot be explained away as an epiphenomenon that scales with decreased movement in PD. Our findings strengthen the idea that dysfunctional beta oscillations are causal in PD symptomology, and extend their implications to non-motor symptoms of the disease.

Materials and Methods

The study was approved by the University of Birmingham Research Ethics Committee (ERN_09-528AP20) and written informed consent was obtained from each participant. Data collection was carried out during a single laboratory session for each participant at the University of Birmingham.

Behavioural task

Participants were seated approximately 1 m from a 19 inch computer monitor. Stimuli were presented in black text against a grey background using the Psychophysics Toolbox extension of Matlab (Brainard, 1997). The task was divided into eight blocks and each block into three stages (Fig. 5).

First, there was an *encoding stage*, which required either deep-semantic or shallow-non-semantic encoding of 30 words presented on the screen one at a time. All participants completed four blocks of each encoding. The order of presentation of each encoding-type was counterbalanced across participants. In the deep-semantic encoding blocks, participants judged whether the presented word was animate i.e. whether it referred to the property of a living entity. In the shallow-non-semantic encoding blocks, participants judged whether the first and last letters of the word were in alphabetical order. These encoding instructions have been used previously to investigate subsequent memory effects (Hanslmayr et al., 2009;

Otten & Rugg, 2001). Participants responded on each trial by pressing one of two response buttons (“yes” or “no”) on the keyboard using their index and middle finger. PD patients used fingers on their less affected hand and hand assignment was randomized (regardless of hand dominance) across healthy participants for comparison with patients. Button assignment was counterbalanced across patients and participants.

The encoding stimuli were taken from a pool of 240 English words, with a list of 120 per encoding condition selected from the MRC psycholinguistic database (Coltheart, 1981). Encoding lists were matched according to word frequency (10 - 93 per million), concreteness (252 - 593), imageability (452 – 615), number of syllables (1 – 4) and number of letters (3 – 10). Words were randomly drawn from the first encoding list for the first four blocks, and the second list for the last four blocks. The order of encoding instructions rather than encoding lists was counterbalanced across participants. A single trial began with a fixation cross for a variable duration of between 1500 and 2000 ms, followed by word presentation for 2000 ms and ended with a question mark to prompt the participant to respond (for which they were given 2500 ms). Participants were instructed not to react during word presentation but give their response during presentation of the question mark.

The second stage in each block consisted of a distracter task during which 20 faces of famous and non-famous people were presented to the participant one at a time. The participant was required to rate the attractiveness of each face using a 6-point rating scale. The distracter stage was intended to prevent the participants rehearsing the word lists, and also to familiarize participants with the 6-button ratings which were to be used in the subsequent recognition stage.

In the final *recognition stage* of each block, the 30 previously encoded words and 15 novel stimuli words drawn from the same pool were presented to participants one at a time. The

order of words was randomized and participants were required to rate their confidence as to whether the word was one encountered in the *encoding stage*, or was a new word. Ratings were given using the 6-point rating scale where response options were R1: recollect, R2: very familiar, R3: familiar, R4: unsure new, R5 sure new, R6: very sure new, using buttons pressed with the index, middle and ring fingers on both hands. The assignment of the buttons was counterbalanced across participants (i.e. R1 – R6 vs R6 – R1), and participants were explicitly instructed to use the full range of confidence ratings. The list of new words was matched to encoding lists for word frequency, concreteness, imageability, number of syllables and number of letters. A trial progressed in the same order and with the same timings as during the *encoding stage*, except that the question mark and button prompts remained on-screen until the participant responded.

EEG recording

Continuous EEG data were recorded using a 128 channel BioSemi ActiveTwo system (BioSemi) with electrodes positioned at the 128 standard equidistant BioSemi sites. Data were digitized using the BioSemi ActiView software, with a sampling rate of 1024 Hz and filtered between 0.1 and 100 Hz.

Behavioural data analysis

Reaction times and response accuracies were recorded during the *encoding stage*. Response times were calculated from the onset of the question mark which prompted the participant to respond until button press. Accuracy was calculated as the number of correct Yes or No responses during each type of encoding expressed as a percentage of all words presented for that encoding condition. All other behavioural analysis and presented data are from the *recognition stage*. Trials in the recognition stage were grouped into high confidence hit (HH), low confidence hit (LH) and miss (M) categories, depending on the participant's

response and their individualized receiver operating characteristic (ROC) curves (Hanslmayr *et al.* 2009). Using ROCs enabled objective quantification of individual response biases and corrected for participants' tendencies to use single buttons of the rating scale differently (Fig. 6). The primary dependent variable, memory strength (d'), was calculated from recognition responses using the following equation.

$$d' = Z[\%Hits] - Z[\%False\ alarms]$$

Z scores were calculated for each individual using MATLAB (The Mathworks). Hits refer to combined HH and LH responses when a word is correctly remembered. False alarms are responses where the participant has incorrectly identified a new word as remembered.

EEG data analysis

All EEG analysis and presented data are from the *encoding stage*. Offline analysis was performed in MATLAB using the open-source FieldTrip toolbox (Oostenveld, Fries, Maris, & Schoffelen, 2011) and in-house MATLAB functions. Raw EEG data were highpass (1 Hz) and lowpass filtered (40 Hz) with finite impulse response filters, re-referenced to the average reference, down-sampled to 500 Hz and epoched into 7000 ms segments around word presentation (3000 ms pre to 4000 ms post stimulus onset) for pre-processing. Independent component analysis allowed components related to ocular artefacts to be visually identified and removed before subsequent visual inspection and manual removal of remaining artefacts. If any channels had been removed during artefact rejection (mean of 0.6 channels removed, min: 0, max: 3), sensor data were interpolated via triangulation of nearest neighbour and then finally re-referenced to the average reference.

The EEG recording epochs extracted from individual encoding trials were grouped into HH, LH and M categories, depending on the participant's subsequent response in the

recognition stage. Epochs were further segmented from 750 ms pre-stimulus to 2000 ms post stimulus for the time-frequency analysis. The entire power spectrum was corrected for 1/f (Podvalny et al., 2015; Voytek et al., 2015) by fitting a linear function to the log-transformed data for every time point and then subtracting the linear fit. The 2.75 s epochs were then subjected to a Morlet wavelet transformation (width of 7 cycles) as implemented in Fieldtrip to extract time-frequency characteristics at frequencies 2 – 40 Hz in steps of 1 Hz. Average power values were calculated for each trial type (HH, LH and M) and baseline corrected (relative change, baseline -750 to -250 ms). This baseline duration is common to examine beta ERD in memory paradigms (e.g. (Hanslmayr et al., 2009; Meconi et al., 2016)) and the timing avoids filter smearing from post stimulus effects into the baseline period. The primary dependent measure was beta power decrease (i.e. ERD) for words that were subsequently successfully remembered, regardless of confidence level (i.e. during successful encoding of a memory resulting in a HH or LH trial in the recognition stage). The analysis of beta ERD between and within groups included an average of 101 (min 66/max 118) trials for controls and 98 trials (min 66/max 118) for PD participants in deep-semantic encoding, and 71 trials for both controls (min 33/max 98) and PD participants (min 25/max 103) in shallow-non-semantic encoding. The secondary dependent measure was the subsequent-memory effect (SME) in beta power which compared power between HH and M trials (Brewer et al., 1998; Hanslmayr et al., 2009; Otten et al., 2001).

Statistical analysis

For memory strength (d'), participants who had values outside 3 standard deviations of the group mean were removed using the median absolute deviation method. The Shapiro-Wilk test ensured normality before using a mixed-effects repeated-measures 2X2 analysis of variance (ANOVA) with factors Group (Controls, PD) and Encoding (Deep, Shallow) as per

our pre-registered protocol expecting a Group X Encoding interaction (MacDonald H, Jenkinson N, Hanslmayr S. Memory encoding and beta desynchronisation in Parkinson's disease [Internet]. 2016 Available from: <https://osf.io/vb64n/>). Post-hoc and planned comparisons were performed using *t*-tests. A least absolute shrinkage and selection operator (LASSO) regression was performed for the PD group to determine the capacity of age and/or disease duration to predict memory strength following deep-semantic and shallow-non-semantic encoding, accounting for collinearity between age and disease duration. A mixed-effects repeated-measures 2X2 ANOVA with factors Group (Controls, PD) and Encoding (Deep, Shallow) tested for differences between groups in encoding accuracy and reaction time for the two encoding conditions.

In alignment with previous EEG studies, and as per our pre-registered protocol, post-stimulus beta power decreases are expected to be associated with successful memory formation in healthy (Hanslmayr et al., 2009; Hanslmayr et al., 2011) and patient populations (Meconi et al., 2016). Therefore lower beta from 12 – 20 Hz was the main frequency range of interest for all dependent measures (see <https://osf.io/vb64n/>). Only negative clusters in this frequency range were expected so comparisons of scalp-wide group averaged data were subjected to one-tailed cluster-based permutation testing (2000 iterations) using the Monte-Carlo 'maxsum' method (Meconi et al., 2016), averaged over 12 – 20 Hz and 0 – 1.5 s relative to encoding stimulus onset. The time window of 0 – 1.5 s post encoding stimulus was chosen based on findings from previous studies investigating beta ERD using the same or similar memory paradigm (Hanslmayr et al., 2009; Meconi et al., 2016) and to avoid capturing any motor-related beta activity prior to the cue for a motor response (Pfurtscheller & Lopes da Silva, 1999) which appeared at the end of the encoding period (2 s after encoding stimulus). Data from all 128 electrodes are included in all EEG analyses. The only exception is for the additional correlational analyses in PD patients to further investigate the effect of encoding

on their beta ERD at an individual level, when a subset of only left frontal electrodes was used based on a literature-driven prior hypothesis (Hanslmayr et al., 2009; Hanslmayr et al., 2011; Meeuwissen et al., 2011). This subset consisted of the front left quadrant taken from left sagittal to vertex (D23 – A1 on BioSemi cap), and vertex down to mid frontal (A1 – C17). A 2x2 mixed-effects repeated-measures ANOVA also tested for an Encoding (Shallow, Deep) X Group (Controls, PD) interaction of beta ERD and SME averaged for each participant over 0 – 1.5 s, 12 – 20 Hz and significant cluster electrodes. Linear regression tested for a relationship between each PD individual’s maximum beta desynchronization over left frontal electrodes during deep-semantic encoding and i) memory strength and ii) disease duration.

The criterion for all statistical significance was $\alpha = 0.05$. Greenhouse-Geisser P values are reported for non-spherical data.

Data availability

Anonymized data, not published in the article, will be shared on reasonable request from a qualified investigator.

Acknowledgements

We thank Federica Meconi, Danying Wang and Sophie Watson for assistance with data collection.

Funding

H.J.M is supported by a Neurological Foundation of New Zealand Philip Wrightson Postdoctoral Fellowship. J.B. is supported by the Medical Research Council (MR/N003446/2). S.H. is supported by a Consolidator grant from the ERC (Grant #647954)

618 and is further supported by the Wolfson Society and the Royal Society. NJ receives ongoing
619 support from Parkinson's UK.

620 **Competing interests**

621 The authors declare no competing financial interests.

622

623 Tables

	HC	PD
Age (y)	67 (9)	65 (6)
Education	3.8 (0.4)	3.9 (0.4)
Gender	13F/18M	8F/20M
Disease Duration (y)	N/A	6 (4)
Handedness	3L/28R	5L/23R
OCS-Plus	9.7 (0.5)	9.7 (0.5)

624 **Table 1. Participant demographics and global cognitive function.**

625 Values are mean (standard deviation) unless otherwise specified. HC: healthy controls; PD:
626 Parkinson's disease; OCS-Plus: Oxford Cognitive Screen Plus questionnaire (max 10).
627 Education is grouped into 1: no formal education; 2: primary school; 3: secondary school; 4:
628 tertiary level.

629

Subject	Age (year)	Gender	PD Medication	LEDD (mg)	Disease Duration (year)	Side Most Affected
1	61	M	Stalevo: 375mg levodopa (5x75mg/18.75mg/200mg) Ropinirole 8mg Rasagiline 1mg	759	11	R
2	65	F	Rasagiline 1mg Madopar: 800mg levodopa (4x50mg/200mg)	900	8	L
3	76	F	Repinex 8mg Sinemet: 500mg levodopa (4x25mg/100mg, 1x25mg/100mg CR)	635	11	L
4	68	M	Sinemet: 300mg levodopa (3x25mg/100mg)	300	5	R
5	62	M	Stalevo: 200mg levodopa (4x50mg/12.5mg/200mg) Rasagiline 1mg Apomorphine 3mg	476	10	R

Repinex 4mg						
Madopar: 200mg levodopa						
6	67	M	(4x12.5mg/50mg)	300	2	L
Rasagiline 1mg						
Madopar: 400mg levodopa						
7	68	M	(4x25mg/100mg)	660	8	L
Rasagiline 1mg						
Repinex 8mg						
Madopar: 300mg levodopa						
8	58	F	(3x25mg/100mg)	326	6	L
Mirapexin 0.26mg						
Selegiline 5mg						
Sinemet: 500mg levodopa						
9	72	M	(5x25mg/100mg)	1090	13	L
ReQuipXL 12mg						
Amantadine 300mg						
10	79	M	Rasagiline 1mg	260	6	R

Ropinirole 8mg						
Stalevo: 700mg levodopa						
11	74	M	(3x200mg/50mg/200mg,	1711	14	L
			1x100mg/25mg/200mg)			
			Amantadine 300mg			
			Rotigotine 16mg			
12	64	M	Mirapexin 1.56mg	156	3	L
Rotigotine 8mg						
13	67	M	Rasagiline 1mg	1804	10	R
			Madopar: 400mg levodopa			
			(4x25mg/100mg)			
			Entacapone 800mg			
Rasagiline 1mg						
14	67	M	Pramipexole 2.1mg	610	4	R
			Sinemet: 300mg levodopa			
			(3x25mg/100mg)			
15	61	F	None	N/A	3	L

16	59	M	Rasagiline 1mg	100	1	R
Rasagiline 1mg						
Sinemet: 100mg levodopa						
(1x25mg/100mg)						
17	56	F	Stalevo: 250mg levodopa	773	5	L
(3x50mg/12.5mg/200mg						
1x100mg/25mg/200mg)						
Ropinirole 12mg						
Rotigotine 6mg						
18	75	M	Madopar: 500mg levodopa	680	3	L
(5x25mg/100mg)						
19	62	F	Sinemet: 150mg levodopa	150	0.33	L
(3x12.5mg/50mg)						
20	58	M	Sinemet: 150mg levodopa	150	1	L
(3x12.5mg/50mg)						
21	70	M	Sinemet: 400mg levodopa	400	4	L
(4x25mg/100mg)						
22	59	F	Ropinirole 12mg	640	7	L

Sinemet: 400mg levodopa (4x25mg/100mg)						
23	69	M	Madopar: 150mg levodopa (3x12.5mg/50mg)	150	0.5	L
Madopar: 700mg levodopa (6x25mg/100mg, 1x25mg/100mg CR)						
24	62	M	Ropinirole 16mg	1020	6	R
25	63	M	Requip 10mg	200	1	L
Ropinirole 8mg						
26	61	F	Madopar: 400mg levodopa (8x12.5mg/50mg)	560	4	L
Madopar: 400mg levodopa (4x25mg/100mg)						
27	54	M	Selegiline 25mg	650	1	L
Sinemet: 400mg levodopa (4x25mg/100mg)						
28	73	M		400	8	R

Table 2. Demographic and clinical data for Parkinson's participants.

PD: Parkinson's disease; LEDD: levodopa equivalent daily dose; CR: continuous release.

References

- Binder, J. R., Desai, R. H., Graves, W. W., & Conant, L. L. (2009). Where is the semantic system? A critical review and meta-analysis of 120 functional neuroimaging studies. *Cereb Cortex*, 19(12), 2767-2796. doi:10.1093/cercor/bhp055
- Bliss, T. V., & Collingridge, G. L. (1993). A synaptic model of memory: long-term potentiation in the hippocampus. *Nature*, 361(6407), 31-39. doi:10.1038/361031a0
- Bokura, H., Yamaguchi, S., & Kobayashi, S. (2005). Event-related potentials for response inhibition in Parkinson's disease. *Neuropsychologia*, 43(6), 967-975. doi:10.1016/j.neuropsychologia.2004.08.010
- Brainard, D. H. (1997). The Psychophysics Toolbox. *Spatial Vision*, 10(4), 433-436. doi:10.1163/156856897X00357
- Brewer, J. B., Zhao, Z., Desmond, J. E., Glover, G. H., & Gabrieli, J. D. (1998). Making memories: brain activity that predicts how well visual experience will be remembered. *Science*, 281(5380), 1185-1187.
- Brittain, J. S., & Brown, P. (2014). Oscillations and the basal ganglia: motor control and beyond. *NeuroImage*, 85 Pt 2, 637-647. doi:10.1016/j.neuroimage.2013.05.084
- Brittain, J. S., Watkins, K. E., Joundi, R. A., Ray, N. J., Holland, P., Green, A. L., . . . Jenkinson, N. (2012). A role for the subthalamic nucleus in response inhibition during conflict. *J Neurosci*, 32(39), 13396-13401. doi:10.1523/JNEUROSCI.2259-12.2012
- Burgess, A. P., & Gruzelier, J. H. (2000). Short duration power changes in the EEG during recognition memory for words and faces. *Psychophysiology*, 37(5), 596-606.
- Buzsaki, G., & Draguhn, A. (2004). Neuronal oscillations in cortical networks. *Science*, 304(5679), 1926-1929. doi:10.1126/science.1099745
- Chung, J. W., Burciu, R. G., Ofori, E., Coombes, S. A., Christou, E. A., Okun, M. S., . . . Vaillancourt, D. E. (2018). Beta-band oscillations in the supplementary motor cortex are modulated by levodopa and associated with functional activity in the basal ganglia. *NeuroImage Clin*, 19, 559-571. doi:10.1016/j.nicl.2018.05.021
- Cohn, M., Moscovitch, M., & Davidson, P. S. (2010). Double dissociation between familiarity and recollection in Parkinson's disease as a function of encoding tasks. *Neuropsychologia*, 48(14), 4142-4147. doi:10.1016/j.neuropsychologia.2010.10.013
- Coltheart, M. (1981). The MRC psycholinguistic database. *Q J Exp Psychol (Hove)*, 33, 497 - 505.
- Craik, F. I. M., & Lockhart, R. S. (1972). Levels of processing: A framework for memory research. *Journal of Verbal Learning and Verbal behavior*, 11, 671-684.
- Demeyere, N., Riddoch, M. J., Slavkova, E. D., Jones, K., Reckless, I., Mathieson, P., & Humphreys, G. W. (2016). Domain-specific versus generalized cognitive screening in acute stroke. *J Neurol*, 263(2), 306-315. doi:10.1007/s00415-015-7964-4
- Dujardin, K., Bourriez, J. L., & Guieu, J. D. (1994). Event-related desynchronization (ERD) patterns during verbal memory tasks: effect of age. *Int J Psychophysiol*, 16(1), 17-27.
- Dujardin, K., Moonen, A. J., Behal, H., Defebvre, L., Duhamel, A., Duits, A. A., . . . Leentjens, A. F. (2015). Cognitive disorders in Parkinson's disease: Confirmation of a spectrum of severity. *Parkinsonism Relat Disord*. doi:10.1016/j.parkreldis.2015.08.032
- Duzel, E., Habib, R., Schott, B., Schoenfeld, A., Lobaugh, N., McIntosh, A. R., . . . Heinze, H. J. (2003). A multivariate, spatiotemporal analysis of electromagnetic time-frequency data of recognition memory. *NeuroImage*, 18(2), 185-197.
- Edelstyn, N. M., John, C. M., Shepherd, T. A., Drakeford, J. L., Clark-Carter, D., Ellis, S. J., & Mayes, A. R. (2015). Evidence of an amnesia-like cued-recall memory impairment in nondementing idiopathic Parkinson's disease. *Cortex*, 71, 85-101. doi:10.1016/j.cortex.2015.06.021
- Eusebio, A., Thevathasan, W., Doyle Gaynor, L., Pogosyan, A., Bye, E., Foltynie, T., . . . Brown, P. (2011). Deep brain stimulation can suppress pathological synchronisation in parkinsonian patients. *J Neurol Neurosurg Psychiatry*, 82(5), 569-573. doi:10.1136/jnnp.2010.217489

- Fell, J., & Axmacher, N. (2011). The role of phase synchronization in memory processes. *Nat Rev Neurosci*, 12(2), 105-118. doi:10.1038/nrn2979
- Fellner, M. C., Bauml, K. H., & Hanslmayr, S. (2013). Brain oscillatory subsequent memory effects differ in power and long-range synchronization between semantic and survival processing. *NeuroImage*, 79, 361-370. doi:10.1016/j.neuroimage.2013.04.121
- Gabrieli, J. D., Poldrack, R. A., & Desmond, J. E. (1998). The role of left prefrontal cortex in language and memory. *Proc Natl Acad Sci U S A*, 95(3), 906-913.
- Gauggel, S., Rieger, M., & Feghoff, T. A. (2004). Inhibition of ongoing responses in patients with Parkinson's disease. *J Neurol Neurosurg Psychiatry*, 75(4), 539-544.
- Hanslmayr, S., Matuschek, J., & Fellner, M. C. (2014). Entrainment of prefrontal beta oscillations induces an endogenous echo and impairs memory formation. *Curr Biol*, 24(8), 904-909. doi:10.1016/j.cub.2014.03.007
- Hanslmayr, S., Spitzer, B., & Bauml, K. H. (2009). Brain oscillations dissociate between semantic and nonsemantic encoding of episodic memories. *Cereb Cortex*, 19(7), 1631-1640. doi:10.1093/cercor/bhn197
- Hanslmayr, S., Staresina, B. P., & Bowman, H. (2016). Oscillations and Episodic Memory: Addressing the Synchronization/Desynchronization Conundrum. *Trends Neurosci*, 39(1), 16-25. doi:10.1016/j.tins.2015.11.004
- Hanslmayr, S., Staudigl, T., & Fellner, M. C. (2012). Oscillatory power decreases and long-term memory: the information via desynchronization hypothesis. *Front Hum Neurosci*, 6, 74. doi:10.3389/fnhum.2012.00074
- Hanslmayr, S., Volberg, G., Wimber, M., Raabe, M., Greenlee, M. W., & Bauml, K. H. (2011). The relationship between brain oscillations and BOLD signal during memory formation: a combined EEG-fMRI study. *J Neurosci*, 31(44), 15674-15680. doi:10.1523/JNEUROSCI.3140-11.2011
- Hirschmann, J., Ozkurt, T. E., Butz, M., Homburger, M., Elben, S., Hartmann, C. J., . . . Schnitzler, A. (2011). Distinct oscillatory STN-cortical loops revealed by simultaneous MEG and local field potential recordings in patients with Parkinson's disease. *NeuroImage*, 55(3), 1159-1168. doi:10.1016/j.neuroimage.2010.11.063
- Horschig, J. M., Smolders, R., Bonnefond, M., Schoffelen, J. M., van den Munckhof, P., Schuurman, P. R., . . . Jensen, O. (2015). Directed Communication between Nucleus Accumbens and Neocortex in Humans Is Differentially Supported by Synchronization in the Theta and Alpha Band. *PLoS ONE*, 10(9), e0138685. doi:10.1371/journal.pone.0138685
- Jefferies, E. (2013). The neural basis of semantic cognition: converging evidence from neuropsychology, neuroimaging and TMS. *Cortex*, 49(3), 611-625. doi:10.1016/j.cortex.2012.10.008
- Jefferies, E., & Lambon Ralph, M. A. (2006). Semantic impairment in stroke aphasia versus semantic dementia: a case-series comparison. *Brain*, 129(Pt 8), 2132-2147. doi:10.1093/brain/awl153
- Jenkinson, N., & Brown, P. (2011). New insights into the relationship between dopamine, beta oscillations and motor function. *Trends Neurosci*, 34(12), 611-618. doi:10.1016/j.tins.2011.09.003
- Joundi, R. A., Brittain, J. S., Green, A. L., Aziz, T. Z., Brown, P., & Jenkinson, N. (2013). Persistent suppression of subthalamic beta-band activity during rhythmic finger tapping in Parkinson's disease. *Clin Neurophysiol*, 124(3), 565-573. doi:10.1016/j.clinph.2012.07.029
- Kishore, A., Joseph, T., Velayudhan, B., Popa, T., & Meunier, S. (2012). Early, severe and bilateral loss of LTP and LTD-like plasticity in motor cortex (M1) in de novo Parkinson's disease. *Clin Neurophysiol*, 123(4), 822-828. doi:10.1016/j.clinph.2011.06.034
- Klimesch, W., Doppelmayr, M., & Hanslmayr, S. (2006). Upper alpha ERD and absolute power: their meaning for memory performance. *Prog Brain Res*, 159, 151-165. doi:10.1016/S0079-6123(06)59010-7

- 730 Klimesch, W., Sauseng, P., & Gerloff, C. (2003). Enhancing cognitive performance with repetitive
731 transcranial magnetic stimulation at human individual alpha frequency. *Eur J Neurosci*, 17(5),
732 1129-1133.
- 733 Knoke, D., Taylor, A. E., & Saint-Cyr, J. A. (1998). The differential effects of cueing on recall in
734 Parkinson's disease and normal subjects. *Brain Cogn*, 38(2), 261-274.
735 doi:10.1006/brcg.1998.1042
- 736 Lago-Rodriguez, A., Ponzo, V., Jenkinson, N., Benitez-Rivero, S., Del-Olmo, M. F., Hu, M., . . . Cheeran,
737 B. (2016). Paradoxical facilitation after depotentiation protocol can precede dyskinesia onset
738 in early Parkinson's disease. *Exp Brain Res*, 234(12), 3659-3667. doi:10.1007/s00221-016-
739 4759-5
- 740 Lalo, E., Thobois, S., Sharott, A., Polo, G., Mertens, P., Pogosyan, A., & Brown, P. (2008). Patterns of
741 bidirectional communication between cortex and basal ganglia during movement in patients
742 with Parkinson disease. *J Neurosci*, 28(12), 3008-3016. doi:10.1523/JNEUROSCI.5295-
743 07.2008
- 744 Litvak, V., Jha, A., Eusebio, A., Oostenveld, R., Foltynie, T., Limousin, P., . . . Brown, P. (2011). Resting
745 oscillatory cortico-subthalamic connectivity in patients with Parkinson's disease. *Brain*,
746 134(Pt 2), 359-374. doi:10.1093/brain/awq332
- 747 McFarland, D. J., Miner, L. A., Vaughan, T. M., & Wolpaw, J. R. (2000). Mu and beta rhythm
748 topographies during motor imagery and actual movements. *Brain Topogr*, 12(3), 177-186.
- 749 Meconi, F., Anderl-Straub, S., Raum, H., Landgrebe, M., Langguth, B., Bauml, K. T., & Hanslmayr, S.
750 (2016). Aberrant prefrontal beta oscillations predict episodic memory encoding deficits in
751 schizophrenia. *Neuroimage Clin*, 12, 499-505. doi:10.1016/j.nicl.2016.08.017
- 752 Meeuwissen, E. B., Takashima, A., Fernandez, G., & Jensen, O. (2011). Evidence for human fronto-
753 central gamma activity during long-term memory encoding of word sequences. *PLoS ONE*,
754 6(6), e21356. doi:10.1371/journal.pone.0021356
- 755 Miller, K. J., Schalk, G., Fetz, E. E., den Nijs, M., Ojemann, J. G., & Rao, R. P. (2010). Cortical activity
756 during motor execution, motor imagery, and imagery-based online feedback. *Proc Natl Acad*
757 *Sci U S A*, 107(9), 4430-4435. doi:10.1073/pnas.0913697107
- 758 Moisello, C., Blanco, D., Lin, J., Panday, P., Kelly, S. P., Quartarone, A., . . . Ghilardi, M. F. (2015).
759 Practice changes beta power at rest and its modulation during movement in healthy subjects
760 but not in patients with Parkinson's disease. *Brain Behav*, 5(10), e00374.
761 doi:10.1002/brb3.374
- 762 Nelson, A. B., Moisello, C., Lin, J., Panday, P., Ricci, S., Canessa, A., . . . Ghilardi, M. F. (2017). Beta
763 Oscillatory Changes and Retention of Motor Skills during Practice in Healthy Subjects and in
764 Patients with Parkinson's Disease. *Front Hum Neurosci*, 11, 104.
765 doi:10.3389/fnhum.2017.00104
- 766 Obeso, I., Wilkinson, L., Casabona, E., Bringas, M. L., Alvarez, M., Alvarez, L., . . . Jahanshahi, M.
767 (2011). Deficits in inhibitory control and conflict resolution on cognitive and motor tasks in
768 Parkinson's disease. *Exp Brain Res*, 212(3), 371-384. doi:10.1007/s00221-011-2736-6
- 769 Oostenveld, R., Fries, P., Maris, E., & Schoffelen, J. M. (2011). FieldTrip: Open source software for
770 advanced analysis of MEG, EEG, and invasive electrophysiological data. *Comput Intell*
771 *Neurosci*, 2011, 156869. doi:10.1155/2011/156869
- 772 Otten, L. J., Henson, R. N., & Rugg, M. D. (2001). Depth of processing effects on neural correlates of
773 memory encoding: relationship between findings from across- and within-task comparisons.
774 *Brain*, 124(Pt 2), 399-412.
- 775 Otten, L. J., & Rugg, M. D. (2001). Task-dependency of the neural correlates of episodic encoding as
776 measured by fMRI. *Cereb Cortex*, 11(12), 1150-1160.
- 777 Patterson, K., Nestor, P. J., & Rogers, T. T. (2007). Where do you know what you know? The
778 representation of semantic knowledge in the human brain. *Nat Rev Neurosci*, 8(12), 976-
779 987. doi:10.1038/nrn2277

780 Pfurtscheller, G., & Lopes da Silva, F. H. (1999). Event-related EEG/MEG synchronization and
781 desynchronization: basic principles. *Clin Neurophysiol*, 110(11), 1842-1857.

782 Podvalny, E., Noy, N., Harel, M., Bickel, S., Chechik, G., Schroeder, C. E., . . . Malach, R. (2015). A
783 unifying principle underlying the extracellular field potential spectral responses in the
784 human cortex. *J Neurophysiol*, 114(1), 505-519. doi:10.1152/jn.00943.2014

785 Raskin, S. A., Borod, J. C., & Tweedy, J. (1990). Neuropsychological aspects of Parkinson's disease.
786 *Neuropsychol Rev*, 1(3), 185-221.

787 Ray, N. J., Jenkinson, N., Wang, S., Holland, P., Brittain, J. S., Joint, C., . . . Aziz, T. (2008). Local field
788 potential beta activity in the subthalamic nucleus of patients with Parkinson's disease is
789 associated with improvements in bradykinesia after dopamine and deep brain stimulation.
790 *Exp Neurol*, 213(1), 108-113. doi:10.1016/j.expneurol.2008.05.008

791 Sederberg, P. B., Kahana, M. J., Howard, M. W., Donner, E. J., & Madsen, J. R. (2003). Theta and
792 gamma oscillations during encoding predict subsequent recall. *J Neurosci*, 23(34), 10809-
793 10814.

794 Sederberg, P. B., Schulze-Bonhage, A., Madsen, J. R., Bromfield, E. B., McCarthy, D. C., Brandt, A., . . .
795 Kahana, M. J. (2007). Hippocampal and neocortical gamma oscillations predict memory
796 formation in humans. *Cereb Cortex*, 17(5), 1190-1196. doi:10.1093/cercor/bhl030

797 Shing, Y. L., Werkle-Bergner, M., Brehmer, Y., Muller, V., Li, S. C., & Lindenberger, U. (2010). Episodic
798 memory across the lifespan: the contributions of associative and strategic components.
799 *Neurosci Biobehav Rev*, 34(7), 1080-1091. doi:10.1016/j.neubiorev.2009.11.002

800 Spitzer, B., & Haegens, S. (2017). Beyond the Status Quo: A Role for Beta Oscillations in Endogenous
801 Content (Re)Activation. *eNeuro*, 4(4). doi:10.1523/ENEURO.0170-17.2017

802 Spitzer, B., Hanslmayr, S., Opitz, B., Mecklinger, A., & Bauml, K. H. (2009). Oscillatory correlates of
803 retrieval-induced forgetting in recognition memory. *J Cogn Neurosci*, 21(5), 976-990.
804 doi:10.1162/jocn.2009.21072

805 Suppa, A., Marsili, L., Belvisi, D., Conte, A., Iezzi, E., Modugno, N., . . . Berardelli, A. (2011). Lack of
806 LTP-like plasticity in primary motor cortex in Parkinson's disease. *Exp Neurol*, 227(2), 296-
807 301. doi:10.1016/j.expneurol.2010.11.020

808 Swann, N., Poizner, H., Houser, M., Gould, S., Greenhouse, I., Cai, W., . . . Aron, A. R. (2011). Deep
809 brain stimulation of the subthalamic nucleus alters the cortical profile of response inhibition
810 in the beta frequency band: a scalp EEG study in Parkinson's disease. *J Neurosci*, 31(15),
811 5721-5729. doi:10.1523/JNEUROSCI.6135-10.2011

812 Swann, N., Tandon, N., Canolty, R., Ellmore, T. M., McEvoy, L. K., Dreyer, S., . . . Aron, A. R. (2009).
813 Intracranial EEG reveals a time- and frequency-specific role for the right inferior frontal gyrus
814 and primary motor cortex in stopping initiated responses. *J Neurosci*, 29(40), 12675-12685.
815 doi:10.1523/JNEUROSCI.3359-09.2009

816 Voytek, B., Kramer, M. A., Case, J., Lepage, K. Q., Tempesta, Z. R., Knight, R. T., & Gazzaley, A. (2015).
817 Age-Related Changes in 1/f Neural Electrophysiological Noise. *J Neurosci*, 35(38), 13257-
818 13265. doi:10.1523/JNEUROSCI.2332-14.2015

819 Zakzanis, K. K., & Freedman, M. (1999). A neuropsychological comparison of demented and
820 nondemented patients with Parkinson's disease. *Appl Neuropsychol*, 6(3), 129-146.
821 doi:10.1207/s15324826an0603_1

822 Zgaljardic, D. J., Borod, J. C., Foldi, N. S., & Mattis, P. (2003). A review of the cognitive and behavioral
823 sequelae of Parkinson's disease: relationship to frontostriatal circuitry. *Cogn Behav Neurol*,
824 16(4), 193-210.

825 Zion-Golumbic, E., Kutas, M., & Bentin, S. (2010). Neural dynamics associated with semantic and
826 episodic memory for faces: evidence from multiple frequency bands. *J Cogn Neurosci*, 22(2),
827 263-277. doi:10.1162/jocn.2009.21251

830 **Figure legends**

831 **Figure 1. Memory performance.** A) Memory performance during encoding conditions

832 illustrating greater memory strength during deep-semantic encoding for healthy controls (N =
833 31) compared to Parkinson's disease (PD) participants (N = 28). Error bars denote standard
834 error of the mean. * $P < 0.05$. B) Correlation between deep-semantic encoding memory
835 performance and disease duration for PD participants ($P = 0.002$).

836 **Figure 2. Event related desynchronization.** Average beta (12 – 20 Hz) event related

837 desynchronization (ERD) for electrodes in significant and/or largest cluster identified during
838 cluster-based statistical analysis. Top row: between group differences during deep-semantic
839 encoding of remembered words; middle row: between group differences during shallow-non-
840 semantic encoding of remembered words; bottom row: differences within healthy participants
841 between deep-semantic and shallow-non-semantic encoding of remembered words. Grey
842 dashed squares indicate time window used in statistical analysis to identify significant
843 electrode clusters over 12 – 20 Hz. Time course of beta ERD averaged over electrodes
844 contributing to significant and/or largest cluster during encoding of subsequently successfully
845 remembered words for controls (blue, N = 30) compared to Parkinson's disease (PD)
846 participants (red, N = 26) in the deep-semantic encoding (A) and shallow-non-semantic
847 encoding (C) conditions. A power decrease is denoted with negative values. Only deep-
848 semantic encoding showed a significant difference between groups (electrodes contributing to
849 significant cluster black in panel B). Topographical maps show the location of the ERD
850 differences between groups in deep-semantic (B) and shallow-non-semantic (D) encoding,
851 with colder colours indicating significantly greater ERD in controls compared to PD
852 participants. Cluster shown for shallow-non-semantic encoding in C and D did not reach
853 significance. E) Time course of beta ERD averaged over electrodes contributing to significant

cluster during encoding of subsequently successfully remembered words for deep-semantic (green) compared to shallow-non-semantic encoding (magenta) in controls. A power decrease is denoted with negative values. Only controls showed a significant difference between encoding conditions (electrodes contributing to significant cluster black in F). Topographical map in F shows the location of ERD differences between encoding conditions, with colder colours indicating significantly greater ERD in deep-semantic compared to shallow-non-semantic encoding. No cluster identified between encoding conditions for PD patients.

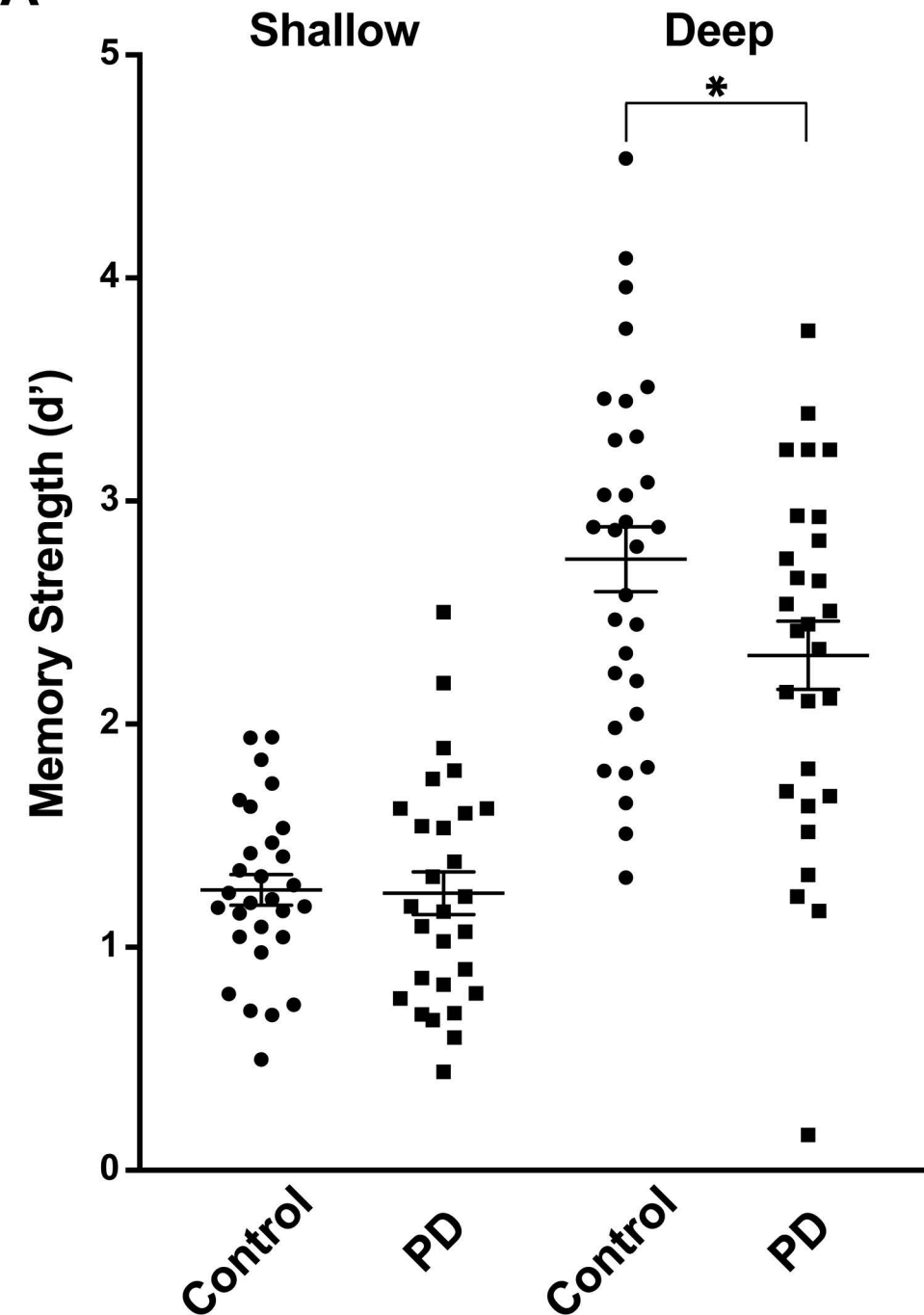
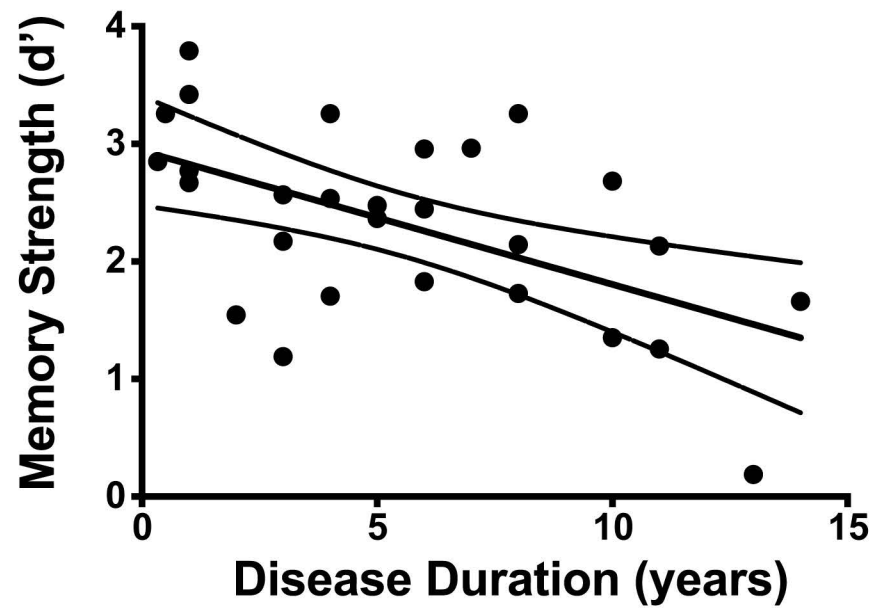
Figure 3. Correlations in Parkinson's disease patients. A) Correlation between deep-semantic encoding memory performance and maximum beta ERD over left frontal electrodes for PD participants ($N = 26$, $P = 0.008$, $R^2 = 0.256$). B) Correlation between maximum beta ERD over left frontal electrodes and disease duration for PD participants ($N = 26$, $P = 0.007$).

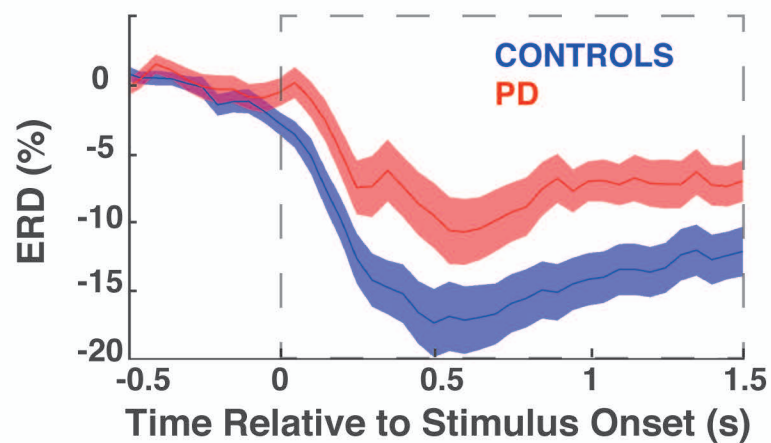
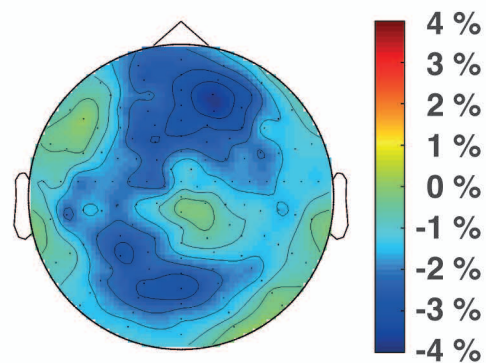
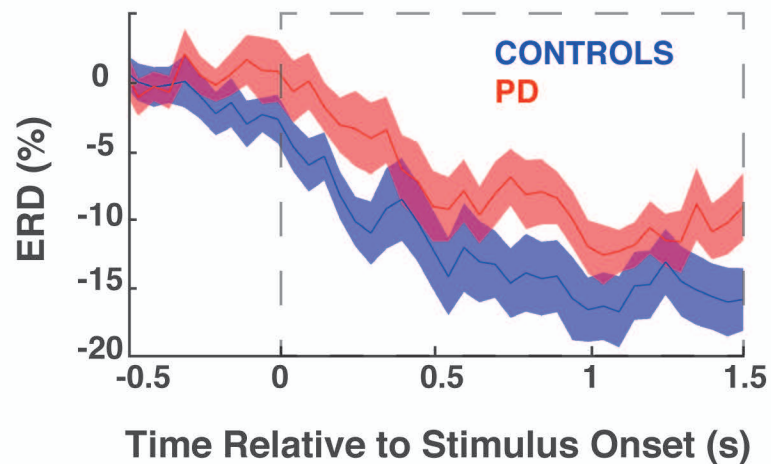
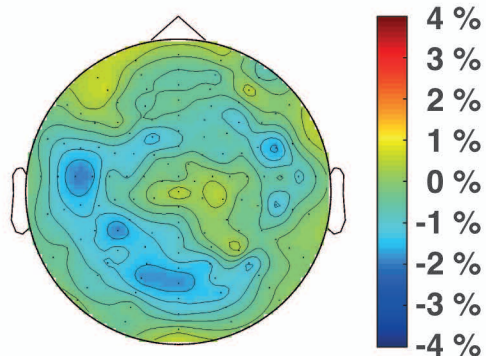
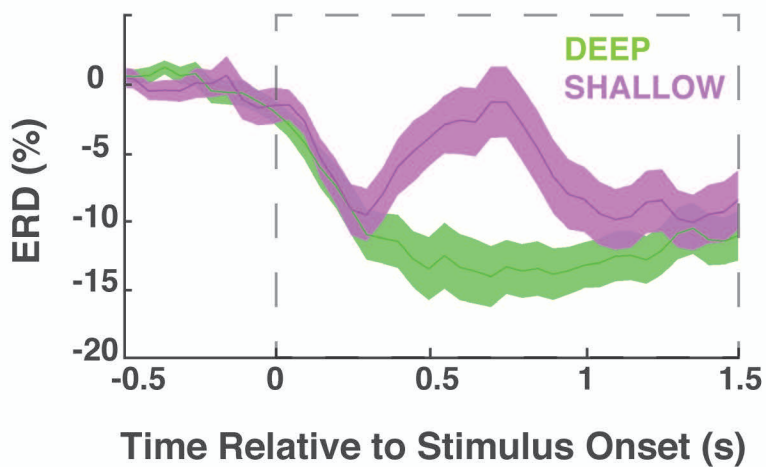
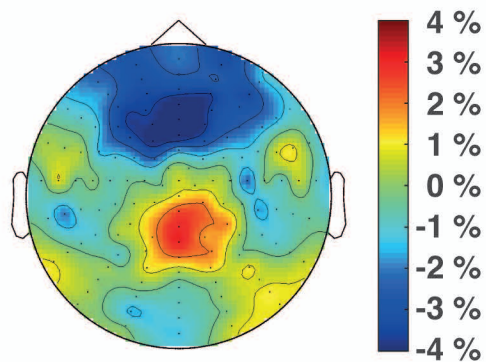
Figure 4. Subsequent memory effects. Average beta (12 – 20 Hz) event related desynchronization (ERD) for electrodes in significant and/or largest cluster identified during cluster-based statistical analysis. Top row: differences within healthy participants between remembered and forgotten words during deep-semantic encoding; middle row: differences within PD patients between remembered and forgotten words during deep-semantic encoding; bottom row: differences within healthy participants between remembered and forgotten words during shallow-non-semantic encoding. Grey dashed squares indicate time window used in statistical analysis to identify significant electrode clusters over 12 – 20 Hz. Time course of beta ERD averaged over electrodes contributing to significant and/or largest cluster during high confidence hit (HH, cyan) compared to miss (M, yellow) trials in deep-semantic encoding for controls (A, $N = 30$) and PD participants (C, $N = 26$). Both groups demonstrated greater ERD during encoding of subsequently remembered (HH) compared to forgotten (M) words, but only the cluster in controls reached significance (electrodes contributing to significant cluster black in B). Topographical maps show the location of the

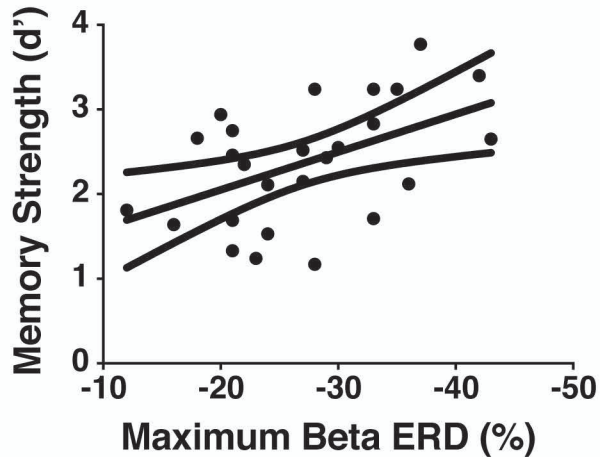
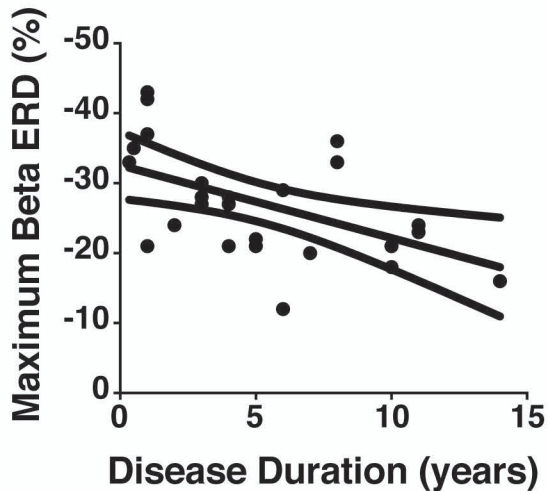
ERD differences between words in deep-semantic encoding for controls (B) and PD patients (D), with colder colours indicating greater ERD for remembered compared to forgotten words. Time course (E) and location (F) of beta ERD averaged over electrodes contributing to largest, non-significant cluster during high confidence hit (HH, cyan) compared to miss (M, yellow) trials in shallow-non-semantic encoding for controls (N = 30). No cluster identified between remembered and forgotten words in shallow-non-semantic encoding for PD patients.

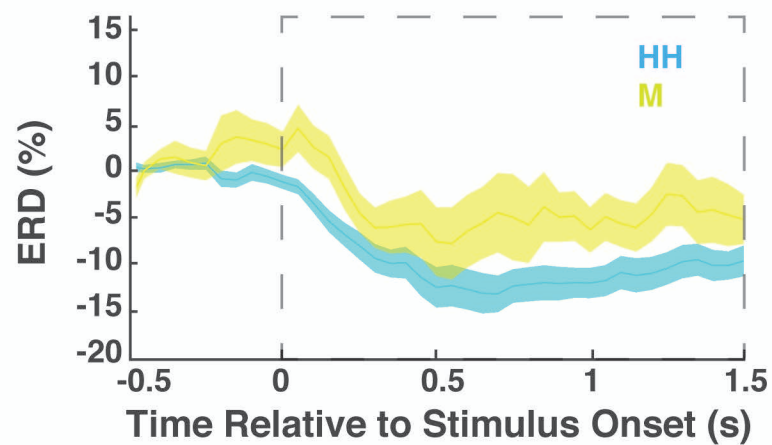
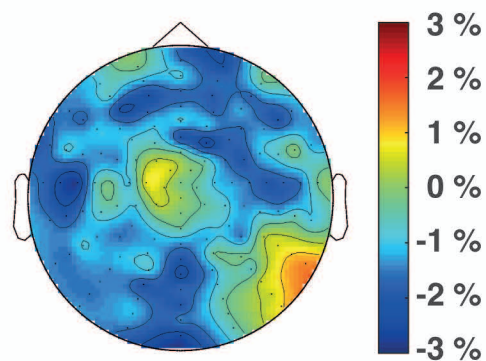
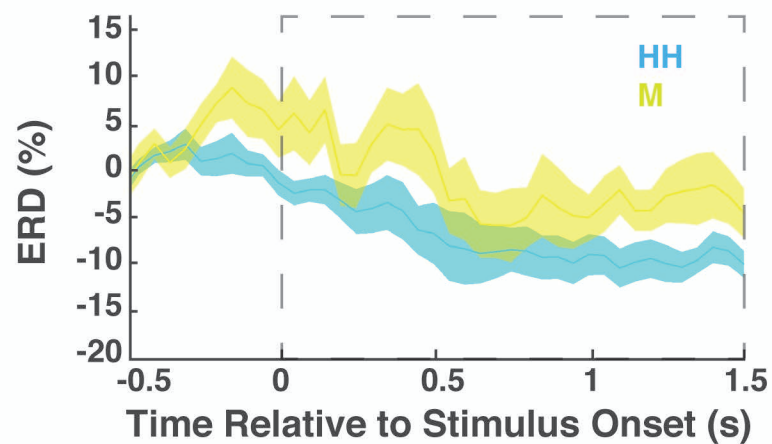
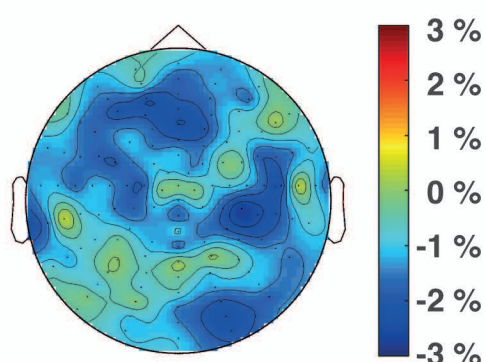
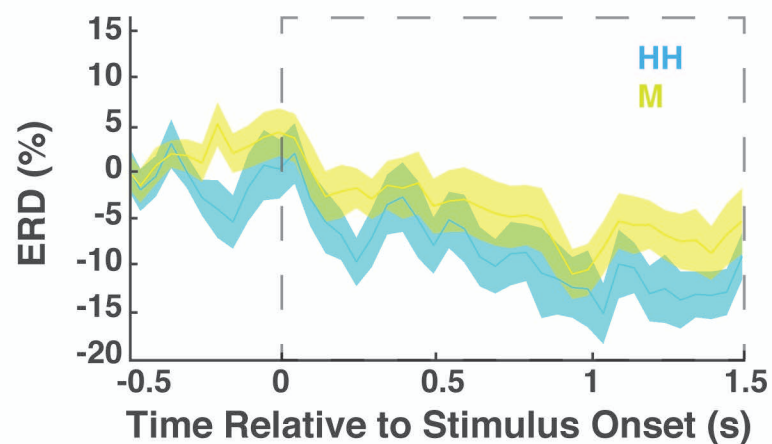
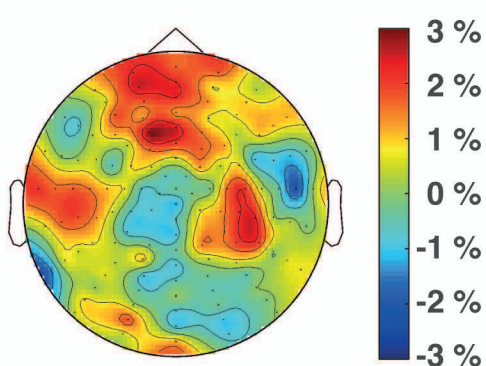
Figure 5. Three stages of memory task. The letters in brackets indicated to participants which button on the keyboard corresponded to which response. In the final screen for a recognition trial participants saw assigned responses (i.e. recollection, very familiar etc.) rather than R1 – R6 which are shown here due to space constraints.

Figure 6. Receiver operating characteristic curves. ROC curves for a representative control (A) and Parkinson's disease participant (B) in deep-semantic and shallow-non-semantic encoding conditions. The false alarm rate is cumulative. The responses given on the 6-point rating scale are grouped into the following conditions: high confidence hit (HH); low confidence hit (LH); miss (M).

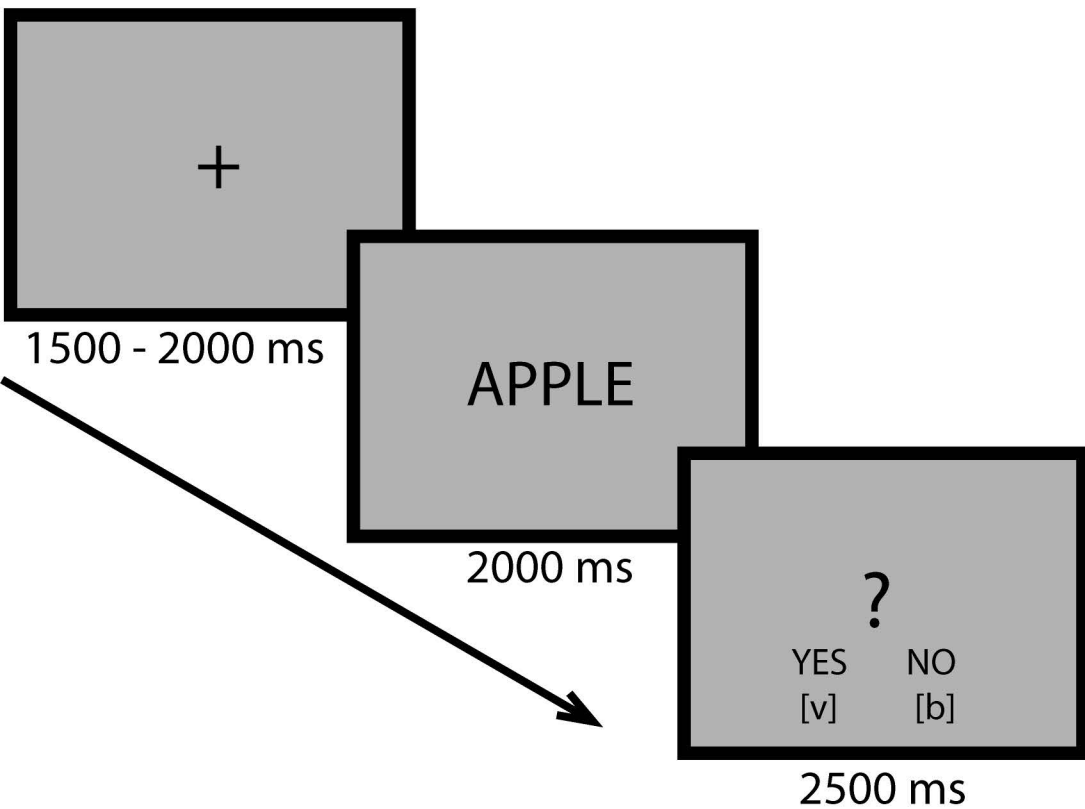
A**B**

A**B****C****D****E****F**

A**B**

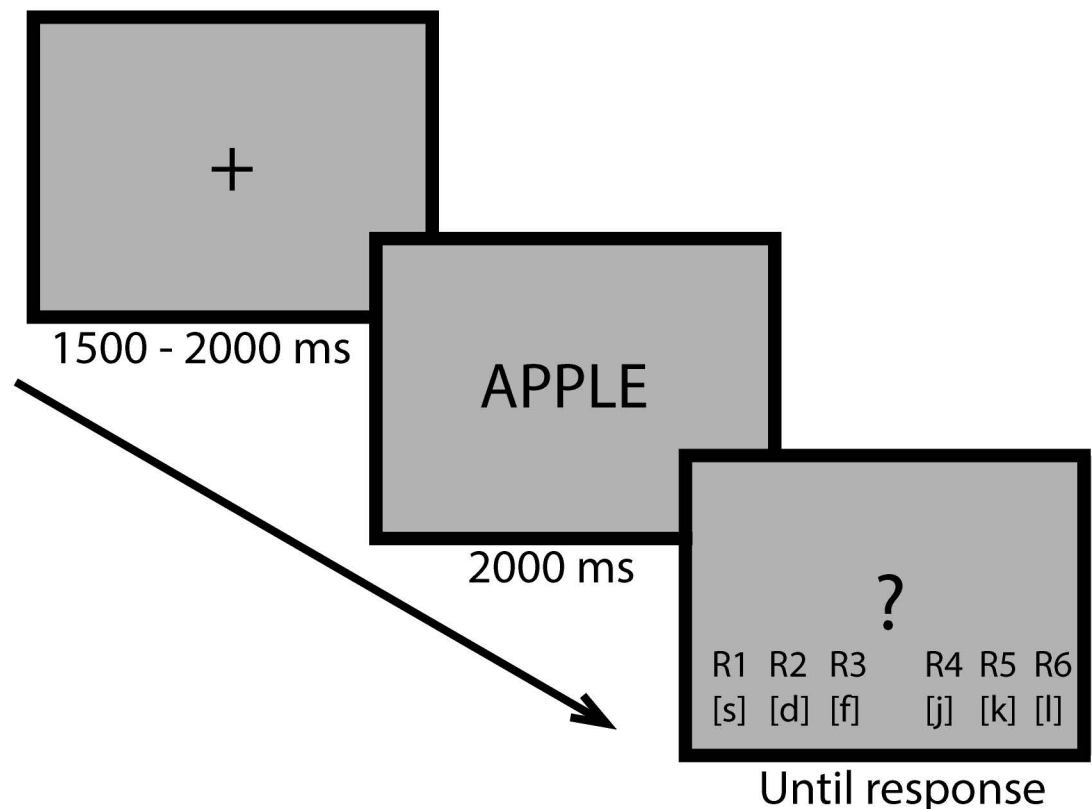
A**B****C****D****E****F**

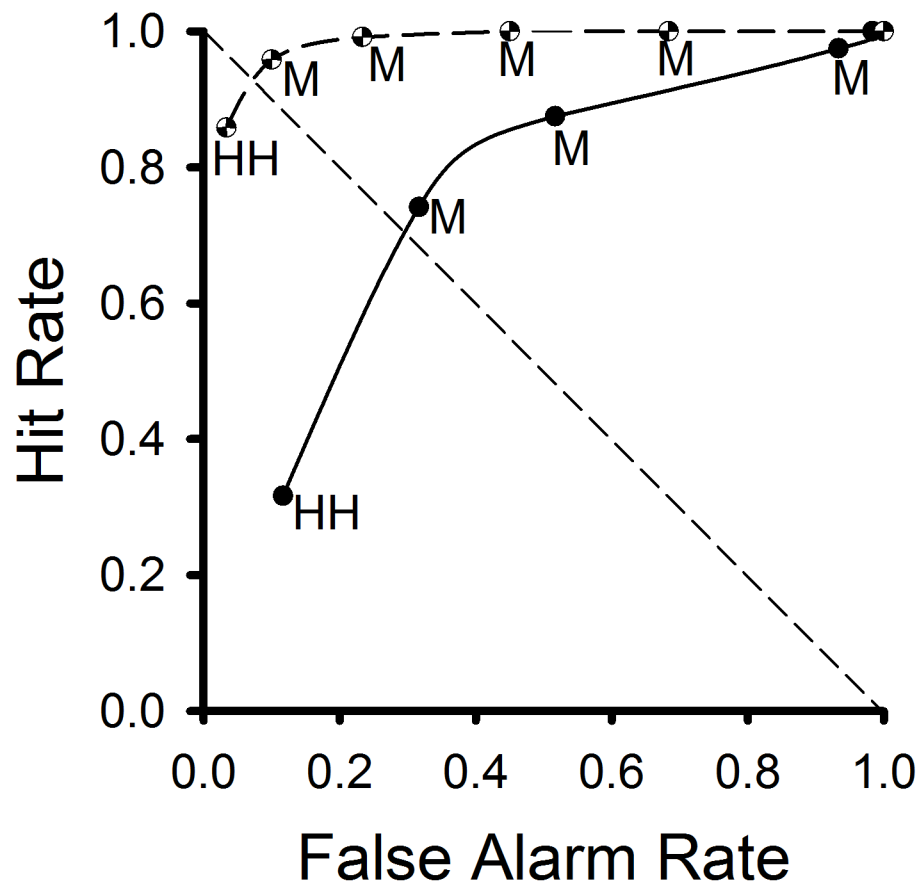
Encoding



DISTRACTOR TASK

Recognition



A**B**