

Oscillatory brain activity and maintenance of verbal and visual working memory: A systematic review

Yuri G. Pavlov^{1,2}  | Boris Kotchoubey¹ 

¹Institute of Medical Psychology and Behavioral Neurobiology, University of Tübingen, Tübingen, Germany

²Department of Psychology, Ural Federal University, Ekaterinburg, Russian Federation

Correspondence

Yuri G. Pavlov, Institute of Medical Psychology and Behavioral Neurobiology, University of Tübingen, Tübingen 72076, Germany.

Email: pavlovug@gmail.com

Funding information

Russian Foundation for Basic Research, Grant/Award Number: #19-013-00027

Abstract

Brain oscillations likely play a significant role in the storage of information in working memory (WM). Despite the wide popularity of the topic, current attempts to summarize the research in the field are narrative reviews. We address this gap by providing a descriptive systematic review, in which we investigated oscillatory correlates of maintenance of verbal and visual information in WM. The systematic approach enabled us to challenge some common views popularized by previous research. The identified literature (100 EEG/MEG studies) highlighted the importance of theta oscillations in verbal WM: frontal midline theta enhanced with load in most verbal studies, while more equivocal results have been obtained in visual studies. Increasing WM load affected alpha activity in most studies, but the direction of the effect was inconsistent: the ratio of studies that found alpha increase versus decrease with increasing load was 80/20% in the verbal WM domain and close to 60/40% in the visual domain. Alpha asymmetry (left < right) was a common finding in both verbal and visual WM studies. Beta and gamma activity studies yielded the least convincing data: a diversity in the spatial and frequency distribution of beta activity prevented us from making a coherent conclusion; gamma rhythm was virtually neglected in verbal WM studies with no systematic support for sustained gamma changes during the delay in EEG studies in general.

1 | INTRODUCTION

Working memory (WM) is the ability to maintain and manipulate information for a short period of time (Baddeley, 2003). The temporal structure of WM can be subdivided into a period of information encoding, maintenance, and retrieval. The maintenance interval (delay period) is a defining component of WM differentiating it from other types of memory.

Dissecting the temporal structure of WM delay using behavioral measures as well as by the employment of functional magnetic resonance imaging (fMRI) is compounded with a lot of assumptions and limitations (Gitelman et al., 2003; Lindquist et al., 2009; Steinbrink et al., 2006). Behavioral

measures such as reaction time and accuracy only reflect the result of information processing in WM. fMRI studies of WM focus on the spatial distribution of activation in the brain and have limited time resolution.

Qualitatively different types of information about WM are available from M/EEG (magneto-/electroencephalography) signals. Brain oscillations derived from M/EEG play an important role in human cognition and represent an energy-efficient mechanism for communication within the brain (Buzsaki, 2004; Fell & Axmacher, 2011). An important question is how different oscillations relate to the maintenance of information in WM. The neural activity that persists over the delay period is a subject of extensive research in psychology (D'Esposito & Postle, 2015).

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited and is not used for commercial purposes.

© 2020 The Authors. *Psychophysiology* published by Wiley Periodicals LLC on behalf of Society for Psychophysiological Research

At a first glance, the amount of literature on oscillatory mechanisms supporting WM performance is enormous. The excessive amount of literature may be one reason why there is no systematic review of this wide topic. Previous numerous attempts to summarize the findings in the field have led to a few influential narrative reviews (van Ede, 2018; Hsieh & Ranganath, 2014; Klimesch, 1999; Roux & Uhlhaas, 2014). Despite their impactful role in advancing the field, the data in narrative reviews can be selectively reported, favoring the most visible studies, or focusing on studies supporting the narrative (Bushman & Wells, 2001; Pae, 2015). While we acknowledge that there is a place for both types of research synthesis—systematic and narrative reviews—most researchers in the field will likely agree that a comprehensive review of the extant findings is needed for building theories and searching for oscillatory mechanisms underlying WM. In the present review, we did not select data to support or reject any theoretical claims. We aimed to provide a systematic overview of the empirical findings in two domains of WM—a panoramic view of the state of the art in research on verbal and visual WM.

Being a cross-modal construct, WM can depend on stimulus modality (Baddeley et al., 2019; Daniel et al., 2016; Postle, 2006). A non-exhaustive list of the differences between verbal and visual WM may include the speed of memory decay, spatial and temporal distribution of associated neural activity, and the time of encoding and consolidation (Oberauer et al., 2018). A somewhat special place of verbal WM has been identified in a collaborative expert report summarizing the findings in WM research (Oberauer et al., 2018). Therefore, one can expect that some findings would differ between visual and verbal WM. To characterize the generalizability versus specificity of the findings in verbal and visual modalities, we compared the descriptive information between the corresponding studies.

The main objective of the review was to systematically study oscillatory activity during maintenance of verbal and visual information in WM in healthy young individuals under normal conditions.

2 | METHOD

2.1 | Search strategy

Two databases (Pubmed and Web of Science) were included in the search. The first author extracted data from the records. A first search was conducted with the query (meg OR magnetoencephalography OR eeg OR electroencephalography OR electrophysiology*) AND ((working memory) OR (short-term memory)) on 22.04.2019. The second search with the same query was conducted on 02.01.2020 and restricted to 2019 and 2020 publication years. The results were amended by screening references in the articles identified in the databases search, and in review articles.

2.2 | Definition of the eligibility criteria

Among verbal WM tasks, the n-back task is one of the most frequently used. In a n-back task, a continuous sequence of items (e.g., letters or digits) is presented to the participant. At the presentation of each display, the participant makes the decision with a button press whether the item in the current display matches the item presented N displays back. Thus, in the 1-back task, the participant has to compare the stimulus being currently displayed and the immediately previous stimulus stored in memory and press the button if the two are identical.

In the n-back task, memory-related brain activity is confounded with the activity related to the motor response. Therefore, the results of the time-frequency analysis obtained in studies employing this paradigm are likely to be biased. Alpha/beta activity related to the preparation of the response and the rebound after its execution overlaps with the activity related to the maintenance of information in WM and decision making (Pesonen et al., 2007). For instance, Chen and Huang (2015) reported a significant effect of WM load on the activity in the beta frequency band. However, the difference in the onset of beta activity between 1-back and 2-back conditions was the same as the difference in reaction time between the two conditions. Beta suppression latency in the n-back task correlated with reaction time in another study (Palomaki et al., 2012). Therefore, the effect observed by the authors is more likely to be related to the preparation of the response than to WM processes. Even using longer inter-stimulus intervals (ISI) as in Deiber et al. (2007) (3.5 s) does not protect from the rebound of alpha-beta activity after the response. Even if no response trials are analyzed, the EEG activity may be contaminated by response suppression. The difficulty in the isolation of directly WM related cognitive operations makes the task hardly suitable for EEG. Therefore, only studies with a clear separation between maintenance and other cognitive operations were included in the review. All n-back studies were excluded.

Patterns of oscillatory activity obtained from EEG, MEG and intracranial EEG (iEEG) data can be similar. However, some differences exist (Leijten et al., 2003; Malmivuo, 2012; Sharon et al., 2007; Stefan et al., 1994). Particularly, a number of properties make iEEG studies less generalizable. First, the sample of human iEEG studies is much smaller than that of studies using MEG and EEG. Second, physical properties of the signal are different, because more local neuronal populations are recorded by means of iEEG. Third, iEEG is most frequently recorded in epilepsy patients. The patients regularly take painkillers, while recovering from the implantation of electrodes, they also stop taking antiepileptic drugs to provoke seizures. The mild cognitive impairment that may take place in this population requires the cognitive tasks to be modified accordingly. For these reasons, we decided to exclude iEEG literature from the analysis.

Oscillatory brain activity during the delay was the focus of the review. Short delay periods, typical for some EEG

research, may complicate the interpretation of the findings in the context of numerous behavioral studies. Furthermore, the first few hundred milliseconds after a stimulus are usually covered by phase-locked evoked and event-related responses (ERP), which were explicitly not the target of the present work. In this review, we aimed to study sustained oscillatory brain activity as opposed to transient short-lived oscillatory surges elicited by the stimulus onset.

Therefore, we included in the analysis only WM studies with the delay period equal to or longer than 1 s. This inclusion criterion will not give a full guarantee that the sustained oscillatory activity is not contaminated by ERP. We consider this decision a reasonable trade-off to include as many studies as possible to the review to increase the sample size, while avoiding mixing up different processes with different underlying mechanisms.

2.3 | Screening on eligibility criteria abstracts/titles

Inclusion criteria:

1. EEG or MEG study
2. human subjects
3. study oscillatory brain activity
4. population of healthy young adults

Exclusion criteria:

1. animal research
2. ERP study
3. iEEG study
4. special population (such as children, elderly, patients, etc.)
5. motor (e.g., sequential motor task), tactile, or auditory WM task
6. mathematical modeling or methods paper
7. review, editorial, letter to the editor
8. brain stimulation studies (e.g., transcranial electric stimulation, transcranial magnetic stimulation (TMS))
9. interventional study (e.g., the influence of physical exercise, alcohol, caffeine, nicotine consumption, or other drugs)
10. concurrent real-life tasks (e.g., driving, flight simulator, jogging, cycling)

2.4 | Full texts screening and data extraction

The inclusion/exclusion criteria were refined to satisfy the objective to study oscillatory activity during the maintenance of information in WM. To achieve this goal, we extracted data on absolute or relative spectral power in theta (~4–8 Hz),

alpha (~8–13 Hz), beta (~13–30 Hz), and gamma (>30 Hz) frequency bands. Delta frequency band was not included.

Brain oscillations are characterized not only by frequency but also by spatial distribution. Among different types of theta rhythms, the most closely related to the current work theta is frontal midline theta (FMT) rhythm. In the rest of the manuscript, any mention of theta rhythm or FMT will refer to frontal midline theta unless stated otherwise. Similarly, alpha activity, when the separation was possible, was sourced from posterior cortical areas (as opposed, for example, to central or frontal alpha). We did not restrict our analyses of beta and gamma activity to particular spatial locations.

An inclusion criterion that was difficult to infer from the abstracts was the type of analysis. Only those studies were included where spectral power was calculated and statistical output was reported. If one article comprised multiple experiments in different samples, they were treated as independent studies. In the opposite situation, if an article used the dataset reported in another article, the article was either excluded or combined with the previous one into a single study if the results were complementary. In some of the studies, the delay period was not analyzed statistically but we were able to infer the absence or existence of the effects from the figures (e.g., using bar graphs or distribution of spectral power in time with reported measurement error). These studies were also included in the review if at least 1 s of the visual depiction was presented.

Verbal information can be presented auditorily or visually. However, since we aimed to explore the generalizability and specificity of WM modality, verbal WM studies were included only if the information was presented visually.

Thus, the additional inclusion criteria were:

1. the task should follow a model with a separate delay period
2. the duration of the delay interval should be longer than or equal to 1 s. As a consequence, the studies were excluded in which the duration of delay was not reported.
3. spectral power in either theta, alpha, beta, or gamma frequency band during the delay period should be analyzed (e.g., studies reporting the only index of lateralized alpha activity were excluded)
4. the analyzes should include a comparison either with baseline or between levels of WM load
5. type of WM task: verbal or visual (visuospatial)
6. the information is presented visually

Exclusion criteria:

1. connectivity analysis
2. machine learning studies
3. concurrent activity during the delay (n-back task, distractors, mental manipulations)

TABLE 1 Verbal WM studies included in the review

Reference	N	Delay duration, s	Task	Levels of difficulty	MEG/EEG	theta	alpha	beta	gamma	Alpha laterality
Altamura et al. (2010)	20	1	Sternberg (sim)	1:5 letters	MEG	-	-	↑	-	-
Bashivan et al. (2014)	15	3	Sternberg (sim)	4:2, 4, 6, 8 letters	EEG	-	↑	0	-	R
Bonnefond and Jensen (2012)	18	1.1	Sternberg (suc)	1:4 letters	MEG	-	↑*	-	-	-
Brookes et al. (2011)	8	8	Sternberg (suc)	3:2, 5, 8 letters	MEG	↑	-	↓	-	-
Gao et al. (2018)	26	3	Sternberg (sim)	2:4, 6 letters	EEG	-	-	0	0	-
Griesmayr et al. (2010)	15	2.5	DMTS	1:3 letters	EEG	↑	-	-	-	-
Harmony et al. (1996)	10	2	Sternberg (sim)	2:3, 5 digits	EEG	*	↓	-	-	-
Heinrichs-Graham and Wilson (2015)	16	3	Sternberg (sim)	1:6 letters	MEG	0	↑	↑	-	R
Hu et al. (2019) (study 1: DMTS)	23	5	DMTS	3:1, 3, 5 syllables	EEG	-	↑	-	-	R
Hu et al. (2019) (study 2: Sternberg)	20	3	Sternberg (sim)	3:1, 3, 5 digits	EEG	-	↑	-	-	R
Ithipiripat et al. (2013)	13	1.5	mod. Sternberg (suc)	1:4 letters	EEG	↑	↑	↓	0	-
Jensen and Tesche (2002)	10	3	Sternberg (suc)	4:1, 3, 5, 7 digits	MEG	↑	-	-	-	-
Jensen et al. (2002)	10	2.8	Sternberg (sim)	3:2, 4, 6 letters	EEG	↑	↑	-	-	R
Khader et al. (2010)	17	5-7	DMTS	1:6 letters	EEG	↑	↑	-	-	Both
Klimesch et al. (1999)	14	8	Sternberg (suc)	2:5, 10 letters and digits	EEG	↑	↑	-	-	R
Klimesch et al. (1993)	16	2	Sternberg (sim)	2:5, 10 letters and digits	EEG	-	↑	-	-	R
Kottlow et al. (2015)	22	3.5	Sternberg (sim)	2:2, 5 letters	EEG	↑	*	*	-	R
Kustermann et al. (2018)	15	2	Sternberg (sim)	3:1, 3, 5 letters	MEG	0	↑	-	0	R
Kwon et al. (2015)	17	5	Simple span	2:5, 10 words	EEG	↑	↓	-	-	-
Meltzer et al. (2007)	18	10.85	Sternberg (sim)	3:2, 4, 6 digits	EEG	0	0	-	-	both
Michels et al. (2008)	18	3	Sternberg (sim)	3:4, 6, 8 letters	EEG	↑	*	-	-	R
Michels et al. (2010)	16	3.5	Sternberg (sim)	2:2, 5 letters	EEG	↑	↑	↑	↑	R
Okuhata et al. (2013)	10	3.2	Sternberg (sim, suc)	2:3, 5 digits	EEG	-	*	-	-	-
Onton et al. (2005)	23	2-4	mod. Sternberg (suc)	3:3, 5, 7 letters	EEG	↑	-	↑	-	-
Park et al. (2013)	22	1.5	Sternberg (suc)	3:3, 5, 7 digits	EEG	↑	-	0	-	-
Pavlov and Kotchoubey (2017)	56	6.7	mod. Sternberg (sim)	2:5, 7 letters	EEG	↑	*	↑	-	R
Poch et al. (2010)	11	1.2	Sternberg (sim)	1:8 letters	MEG	↑	↑	↑	↑	-

TABLE 1 (Continued)

Reference	N	Delay duration, s	Task	Levels of difficulty	MEG/EEG	theta	alpha	beta	gamma	Alpha laterality
Prokovec, Heinrichs-Graham, et al. (2019)	26	3	Sternberg (sim)	2:4, 6 letters	MEG	0	↑	-	-	R
Schack and Klimesch (2002)	10	1.8	Sternberg (suc)	4:1, 2, 3, 4 digits presented as words	EEG	↑	↑	-	-	-
Scheeringa et al. (2009)	20	7	Sternberg (sim)	3:3, 5, 7 letters	EEG	↑	↑	-	-	R
Stephane et al. (2010)	19	2.05	Sternberg (suc)	1:5 letters	MEG	0	↓	↓	↓	both
Wianda and Ross (2019)	25	3	mod. Sternberg (suc)	1:5 letters	MEG	-	↑	*	-	R
Wilson et al. (1999)	10	4	simple span	5:1, 3, 5, 7, 8 digits	EEG	↑	↓	↓	-	R
Xie et al. (2016)	16	2	Sternberg (suc)	4:2, 4, 6, 8 letters	EEG	-	↑	-	-	-
Zakrzewska and Brzezicka (2014)	69	2.5	Sternberg (suc)	4:2, 3, 4, 5 letters	EEG	↑	-	-	-	-

Note: * complex results described in the main text of the systematic review, - no information available, ↑ increase, ↓ decrease, 0 no changes in the parameter, ↑* the information acquired by personal communication. Levels of difficulty: Number of levels; number of items presented at each level of load. Task: suc, successive; sim, simultaneous presentation of the stimuli; mod. Sternberg, modified Sternberg task. Alpha laterality: R, stronger alpha power changes in the right hemisphere; both—no clear difference between the hemispheres.

3 | RESULTS

The full sample of 100 studies is presented in Table 1 (verbal WM) and Table 2 (visual WM). The flow chart with stages of the selection process is depicted in Figure 1. The 100 articles included 100 studies: one article had two datasets (studies), and one study was published in two articles.

3.1 | General results

Of the 100 selected studies, the number of verbal and visual WM studies was 35 and 65, respectively (see Tables 1 and 2, respectively). Among verbal studies, ten studies employed MEG and 25 EEG. Among visual studies, there were 24 MEG studies, 40 EEG studies, and one study with concurrent EEG/MEG recording.

The number of participants did not significantly differ between the studies carried out in verbal ($mean \pm SD = 19.3 \pm 12$, $median = 17$) and visual modalities ($mean \pm SD = 23.3 \pm 21.9$, $median = 18$) ($t(98) = 1.02$, $p = .311$; see Figure 2 for the distribution).

In verbal studies, the duration of the delay period varied in a range of 1–10.85 s, in the visual modality from 1 to 20 s. Some studies used variable delay duration. In these cases, we included the longest intervals in the analysis. To account for outliers, the median instead of the mean value was used as a central tendency measure. The median duration of the delay period across all 100 studies was 2.8 s, and the difference between the modalities (verbal: 3 s; visual: 2.5 s) did not attain significance ($U = 1,394$, $p = .063$; see Figure 2).

Fifty-eight of the 100 studies (58%) varied set-size and consequently the WM load. The proportion of the verbal WM studies exploring more than one level of load (74%, 26/35) was larger in comparison with visual modality (48%, 31/65) ($\chi^2 = 6.56$, $p = .01$). Verbal experiments never used a single item to encode without additional levels of load (0%, 0/35). The proportion was significantly smaller than in visual modality (26%, 17/65) ($\chi^2 = 11$, $p < .001$).

Figure 2 depicts the distribution of the levels of load in the two WM domains. As can be seen in the figure, load variation starting from one item is quite typical in visual (25%, 30/118) but significantly less common in verbal (7%, 6/84) domain WM research ($\chi^2 = 11.2$, $p < .001$).

3.2 | WM tasks

71% of verbal WM studies (25/35) used the classic version of the Sternberg task (Sternberg, 1966). In this task, the temporal subprocesses in WM such as encoding, maintenance, and retrieval are separated. In the classic version of the task at the encoding phase one or several items are presented either

TABLE 2 Visual WM studies included in the review

Reference	N	Delay duration, s	Content	Levels of difficulty	MEG/EEG	Theta	Alpha	Beta	Gamma	Alpha laterality
Babiloni et al. (2005)	20	3.5–5.5	Relative height of two bars	1:2	MEG	–	↓	–	–	Both
Babiloni et al. (2004)	12	3.5–5.5	Relative height of two bars	1:2	EEG	↓	↓	–	–	R
Bastiaansen et al. (2002)	174	1 or 4	Location	1:1	EEG	↓	↑	–	–	R
Berger et al. (2019)	25	2	Location	2:1, 4	EEG	↑	–	–	–	–
Blacker et al. (2016)	18	2	Location	1:1	EEG	–	↑	–	–	Both
Boonstra et al. (2013)	8	6	Object, location	3:1, 3, 5	EEG	↓	↓	↓	–	Both
Brookes et al. (2012)	10	6	Shape (serial order)	1:2	MEG	0	↓	↓	0	–
Busch and Herrmann (2003)	16	3	Shape, color, texture	3:1, 3, 6	EEG	–	↑	–	–	–
Crespo-Garcia et al. (2013)	29	2.5–5	Location	2:3, 5	EEG	–	↑	–	–	Both
Daume, Graetz, et al. (2017)	29	3	Object	1:1	MEG	0	*	↑	↑	Both
Daume, Gruber, et al. (2017)	27	3	Object	1:1	MEG	0	*	↑	↑	–
de Vries et al. (2017)	20	1.8	Color	2:1, 2	EEG	–	↓	–	–	Both
de Vries et al. (2019)	22	1.4	Color	1:1	EEG	↓	↓	–	–	Both
Ellmore et al. (2017)	15	6	Image (serial order)	1:2	EEG	0	↑	–	–	R
Erickson et al. (2019)	60	2	Color, location	3:2, 4, 6	EEG	–	↓	↓	–	Both
Eschmann et al. (2018)	27	2	Location	2:1, 4	EEG	↑	–	–	–	–
Fukuda et al. (2015)	33	1–4	Color, location	6:1, 2, 3, 4, 6, 8	EEG	–	↓	–	–	–
Grimault et al. (2009)	27	1.2	Color, location	2:2, 4	MEG	–	↑	–	–	–
Heinz and Johnson (2017)	18	1.4	Shape	2:2, 4	EEG	–	↑	–	–	–
Herrmann et al. (2004)	22	3–4	Angle	1:1	EEG	–	↑	–	–	Both
Honkanen et al. (2015)	13	2.05	Shape	2:2, 4	MEG/EEG concurrent	0	↓	↑	↑	–
Ichihara-Takeda et al. (2015)	10	5	Location	1:1	MEG	–	↑	–	–	R
Ikkai et al. (2014)	18	2	Location	1:2	EEG	–	↑	–	–	–
Johnson et al. (2011)	12	3	Shape, location	1:4	EEG	–	↑	–	–	R
Jokisch and Jensen (2007)	10	3	Face	1:1	MEG	–	↑	–	↑	Both
Kaplan et al. (2016)	16	20	Location, object (serial order)	1:5	MEG	↑	–	–	–	–
Kawasaki and Yamaguchi (2012)	19	2	Shape, location	3:2, 4, 6	EEG	↑	↓	–	–	–

TABLE 2 (Continued)

Reference	N	Delay duration, s	Content	Levels of difficulty	MEG/EEG	Theta	Alpha	Beta	Gamma	Alpha laterality
Kawasaki and Yamaguchi (2013)	14	2	Color, location	2:3, 6	EEG	↑	↓	-	-	-
Khader et al. (2010)	17	5-7	Object	1:1	EEG	↑	↑	-	-	Both
Kulashekhar et al. (2016)	13	1.1-1.3	Color	1:1	MEG	↓	↑	↓	0	-
Manza et al. (2014)	31	2	Color, location	3:1, 3, 6	EEG	-	↑	-	-	Both
Map elli and Ozkurt (2019)	28	1-1.5	Object	1:4	EEG	-	↑	-	-	R
Maurer et al. (2015)	24	3.5	Shape	2:2, 4	EEG	↑	*	-	-	R
Medendorp et al. (2007)	8	2,4	Location (serial order)	2:1, 2	MEG	-	↓	↓	-	-
Moran et al. (2010)	14	1.2	Color, location	4:2, 3, 4, 6	MEG	↑	↑	-	-	-
Morgan et al. (2011)	16	2	Color, angle	2:1, 2	MEG	-	-	-	↑	-
Nenert et al. (2012)	16	1-3	Location, orientation	3:2, 4, 6	EEG	-	↑	-	-	Both
Okada and Salenius (1998)	4	2.8	Location	1:4	MEG	-	0	-	-	-
Olsen et al. (2013)	12	2	Object, location	1:3	MEG	↑	-	-	-	-
Pahor and Jausovec (2017)	60	1	Color, location	3:4, 6, 8	EEG	-	↓	-	↓	R
Park et al. (2011)	11	2.5-3.5	Location	1:2	MEG	0	↑	*	*	Both
Poch et al. (2014)	17	1	Orientation, location	2:1, 2	MEG	-	↑	-	0	-
Poch et al. (2018)	36	1	Orientation, location	3:2, 3, 4	EEG	-	↓	-	-	-
Proskovec et al. (2018, Proskovec, Heinrichs-Graham, et al., 2019) combined	22	2.5	Location	2:2, 4	MEG	↓	↓	↓	0	-
Rawle et al. (2012)	20	1	Color, location	3:1, 2, 4	EEG	↓	↓	↓	0	-
Román-López et al. (2019)	26	2,5	Color, shape	1:2	EEG	↑	↑	↑	-	-
Rominger et al. (2019)	52	3	Shape	1:4	EEG	-	↑	-	-	R
Roux et al. (2012)	25	1.2	Location	2:3, 6	MEG	0	↑	↑	↑	-
Sauseng et al. (2005)	23	2.5	Location	1:3	EEG	-	↑	-	-	Both
Schroeder et al. (2018)	30	1.2	Orientation	2:1, 2	EEG	-	↓	-	-	-
Seemuller et al. (2012)	17	1	Angle	1:1	EEG	0	↓	↓	-	Both
Smyrnis et al. (2014)	10	3.5-4.5	Location	1:1	EEG	-	-	↓	↓	-
Spitzer and Blankenburg (2012)	28	3	Light blinking frequency	1:1	EEG	0	↑	-	-	Both
Spitzer et al. (2014)	24	3	Quantity	6:3, 4, 5, 6, 7, 8	EEG	-	↑	↑	-	Both

(Continues)

TABLE 2 (Continued)

Reference	N	Delay duration, s	Content	Levels of difficulty	MEG/EEG	Theta	Alpha	Beta	Gamma	Alpha laterality
Takase et al. (2019)	29	1.5	Orientation (serial order)	1:7	MEG	↑	↑	-	-	-
Tallon-Baudry et al. (1999)	13	0.8–1.6	Location	1:1	EEG	-	-	↑	*	-
Trubutschek et al. (2017)	13	2.5–4	Location	1:1	MEG	-	↑	↓	-	-
Trubutschek et al. (2019)	30	1.5–3	Location	1:1	MEG	-	-	↓	-	-
Tuladhar et al. (2007)	5	2.7	Face (serial order)	4:1, 2, 3, 4	MEG	-	↑	-	-	-
van Dijk et al. (2010)	18	1.5	Location	1:1	MEG	-	0	-	-	-
van Ede et al. (2017)	16	4	Location (serial order)	2:2, 4	MEG	↑	↓	↓	↑	Both
Vandenbroucke et al. (2015)	25	1	Orientation, location	4:2, 4, 6, 8	EEG	0	↓	-	-	-
Visser et al. (2016)	31	1	Color	2:2, 4	EEG	↑	↓	↑	-	-
Yin et al. (2012)	12	1	Shape	1:3	EEG	↓	↑	-	-	-
Zhang et al. (2016)	16	3	Object (serial order)	6:1, 2, 3, 4, 5, 6	EEG	↑	-	-	0	-

Note: * complex results described in the main text of the systematic review, – no information available, ↑ increase, ↓ decrease, 0 no changes in the parameter. Levels of difficulty: Number of levels: number of items presented at each level of load. Alpha laterality: R, stronger alpha power changes in the right hemisphere; both—no clear difference between the hemispheres. Content: type of content presented for encoding. Serial order—the items presented in the way the rehearsal in a sequence is likely to occur.

simultaneously or in a succession, which is followed by a delay period (maintenance interval). After the delay, a single-item probe is presented at the retrieval stage where participants have to decide whether the probe matches any of the items in the encoded memory set. The Sternberg task shares the advantage of temporal separation of subprocesses with the delayed matching-to-sample (DMTS) paradigm. In verbal DMTS, however, (1) only simultaneous presentation of items is possible at the encoding phase, and (2) all the items are probed simultaneously at the retrieval stage. In three studies the DMTS paradigm was used. Two other studies employed a simple span paradigm with free recall.

Five other studies used modified Sternberg paradigms. The modifications included: instead of encoding all letters the participants had to memorize only the marked ones and ignored the others (Onton et al., 2005); the task was to memorize only the letters presented in the previously cued hemifield (Kustermann et al., 2018), in a fashion similar to the lateralized change detection paradigm (Vogel & Machizawa, 2004); at the retrieval phase the probe was replaced by the task to reproduce the sequence of four items by clicking on the presented items in the right order (Itthipuripat et al., 2013); a similar manipulation with the probe was performed in another study where the probe consisted of two items: a letter and a digit representing the serial number of the presented letter, and the participants had to indicate whether the letter was on the presented serial position in the encoded letters string (Pavlov & Kotchoubey, 2017); in one study the probe was a pair of letters, each of them was presented at the encoding but the order of the letters was either correct or reversed (Wianda & Ross, 2019).

Letters (24/35), digits (6/34), words (2/35), a combination of digits and letters (2/35), and syllables (1/35) were used in verbal WM tasks.

Visual WM tasks were much more diverse. In an attempt to systematically categorize the tasks, we used a two-factor model. A typical task involves the presentation of memory items sequentially or simultaneously, followed by the delay period and a probe. An important distinction between the tasks can be made by (1) the type of content for memory maintenance (e.g., location, shape, color, etc., of the stimuli or images or faces to memorize) and (2) relevance of the presentation order to solve the task. For example, a typical change detection task (Luck & Vogel, 1997) requires to memorize the color and location of squares occupying all visual field. The squares are presented simultaneously making the memory content essentially a single-complex item, not a sequence of independent items. In the opposite case, a sequence of complex images (e.g., everyday objects) more likely to be stored as a sequence, not as one complex object.

In 12/65 studies the stimuli were presented sequentially but in three of them, there was no need to maintain the order of presentation along with the identity of stimuli (see

Table 2). As an example, Olsen et al. (2013) presented simple objects sequentially at different non-overlapping positions on the screen. Although the objects were presented in a succession, they may easily be stored as a whole. Thus, the majority (86%, 56/65) of visual WM studies required participants to maintain only the content.

The following content was used in the visual WM tasks: complex objects such as images, recognizable everyday objects (9), faces (2), shapes (geometric shapes or symbols) (10), location (of dots, shapes, or objects) (39), color (16), orientation (7), texture (1), angle (2), frequency of a blinking light (1) and the number of the blinks (1), the relative height of two bars (2). The sum of the above numbers is larger than 65 because some studies used items with more than one feature.

3.3 | Theta

In 25 (18 EEG/ 7 MEG) of 35 verbal WM studies, theta rhythm was featured in the results or depicted in figures (see Table 1 and Figure 3 for a summary of the results). The theta has never been found to decrease in EEG verbal WM tasks with an exception of one study (Harmony et al., 1996). The study reported a decrease of theta compared with baseline, but only in a low load condition (memorizing three digits). One EEG study reported null findings on the group level (Meltzer et al., 2007). All the other EEG studies reported an increase in theta activity during the delay. Another common finding frequently reported in the EEG literature is a stepwise increase of theta activity with load. The review identified 15 studies using the tasks with more than one level of WM load. The effect of the stepwise increase of theta was found in ten out of the 15 studies. In addition to the above-mentioned study of Harmony et al. (1996), in one study the stepwise increase of theta activity was observed only in high-performance participants (Pavlov & Kotchoubey, 2017). No observable and/or statistically significant stepwise increase was found in three studies (Kwon et al., 2015; Meltzer et al., 2007; Schack & Klimesch, 2002). As we can see, EEG verbal WM research strongly supports the hypothesis of the involvement of theta oscillations in WM maintenance processes.

The picture is less clear when MEG data are taken into account. There is only one MEG study where stepwise theta increase with WM load was demonstrated (Jensen & Tesche, 2002). Another study found theta increase from 2- to 5-letters conditions but no further increase in the 8-letters condition (Brookes et al., 2011). Poch et al. (2010) reported an increase of theta in comparison with baseline. Three studies did not report any sustained increase of theta during the delay period (Kustermann et al., 2018; Proskovec, Heinrichs-Graham, et al., 2019; Stephane et al., 2010).

Finally, one MEG study did not find any theta change either during encoding or during the delay period (Heinrichs-Graham & Wilson, 2015). These authors explicitly noticed the discrepancy between their findings and the results of previous MEG and EEG studies. They explained the inconsistency by the difference between the simultaneous presentation of stimuli in their study and the successive presentation in those MEG studies where the effect was found (Brookes et al., 2011; Jensen & Tesche, 2002). However, the hypothesis is not supported by our data. The increase of theta with load was reproducible in both simultaneous (7/11 studies) and successive (8/9 studies) varieties of the Sternberg paradigm ($\chi^2 = 1.68$, $p = .19$). Inclusion of the studies using the simple span (successive presentation, two studies) and DMTS (simultaneous presentation, two studies) paradigms did not change the result (9/13 vs. 10/11 studies in the successive and simultaneous categories, respectively, $\chi^2 = 1.69$, $p = .19$).

Elucidating the role of theta rhythm in the maintenance of WM, Hsieh et al. (2011) argued that theta is important for the maintenance of temporal order and less relevant to the storage function. In the classic Sternberg task, only one item is probed. It means that maintaining the order of the presented items is irrelevant to the task. However, a piece of evidence is provided by a MEG study that compared two conditions: (1) memorizing only content and (2) memorizing the content as well as the location of the presented stimuli (Poch et al., 2010). In this study, the information about the order can be seen as the spatial dimension added to the verbal content. The authors found a stronger theta increase in the verbal-spatial condition in comparison with the verbal condition.

It appears to be difficult to design a task that could provide sufficient evidence for or against this hypothesis in the verbal domain. Moreover, even presented simultaneously verbal content is rehearsed in a sequence. It can be less of a problem for the visual WM domain. Moreover, in a visual Sternberg paradigm using unpronounceable shapes, theta effects were still observable (Maurer et al., 2015). Given the obvious limitation of verbal WM studies in the context of this hypothesis, it can be beneficial to broaden the scope of this part of the review to the visual WM studies.

19/40 EEG and MEG 13/25 visual WM studies featured theta in the results. Among the 19 + 13=32 studies, nine EEG and five MEG studies reported an increase in theta activity during the delay, six EEG and two MEG studies found a decrease of theta, and four EEG and six MEG studies reported null effects. Because, as compared with verbal WM studies, a demonstration of a decrease in theta power appears unexpected, we looked at the corresponding set of studies more closely.

We first hypothesized that this result may be explained by a broadband power suppression of alpha also occupying neighboring frequencies. Although, in five studies the theta

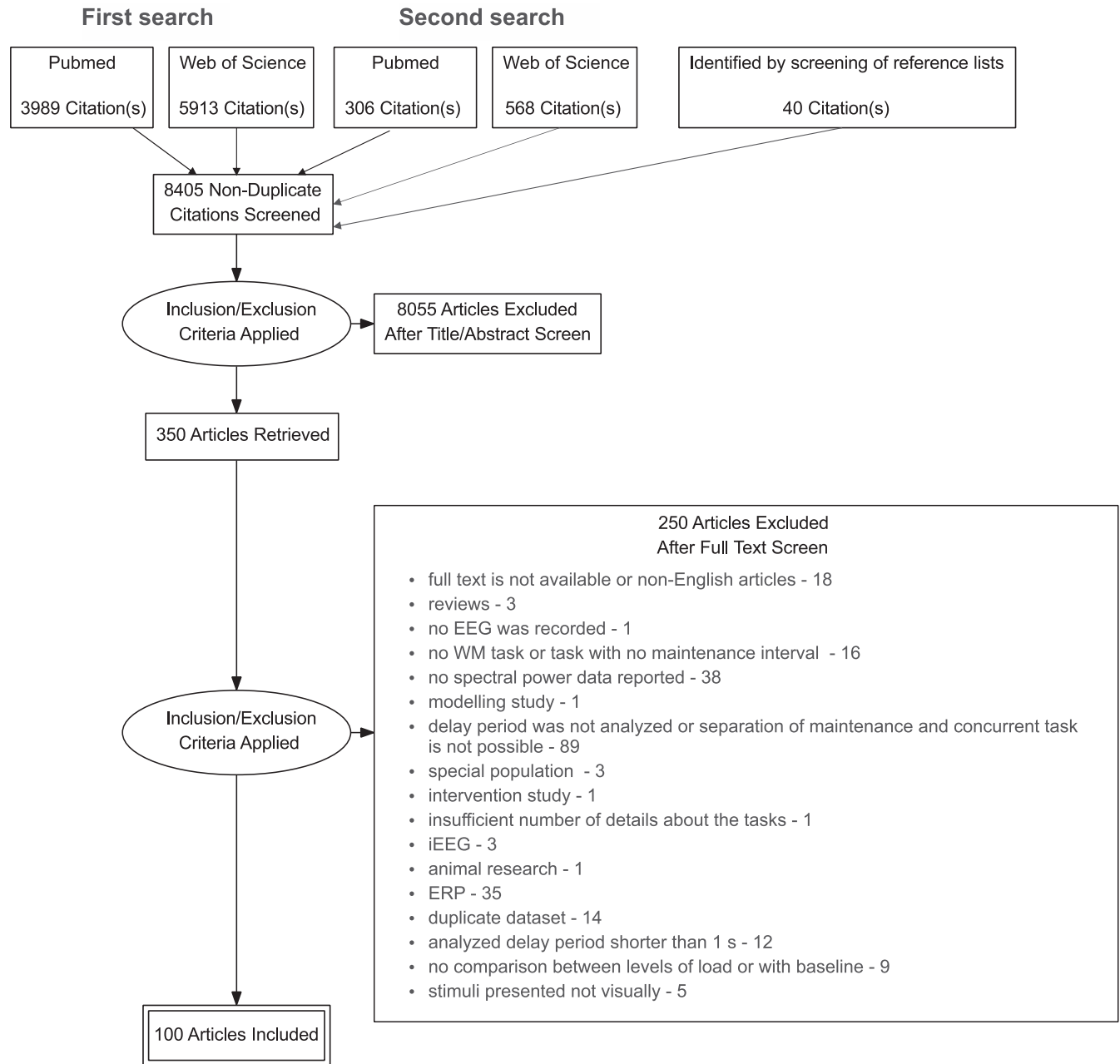


FIGURE 1 Flow chart of the systematic review

decrease was, indeed, accompanied by a similar alpha decrease; in the other three studies, alpha increased with load. Thus, the hypothesis was not supported by the data (5/8 vs. 3/8: $\chi^2 = 1$, $p = .317$).

One of the eight studies that reported theta decrease was the study by Boonstra et al. (2013). However, the data shown in the bottom panel of the first figure in the study (see Figure 4 below) challenge the conclusion of theta suppression during the delay. One can notice that the 4–8 Hz frequency band designated by the authors as theta largely overlaps with alpha and shares its behavior. Taking into account this observation, sustained theta activity in this work was in fact absent.

Proskovec et al. (2018) in a MEG study localized theta in the left dorsolateral prefrontal cortex and reported an increase, but their Figure 3 shows the opposite pattern: after a short transient increase, theta decreased below baseline level and maintained in this state until probe presentation. In contrast to the results by Proskovec, theta continuously increased during the delay period in two EEG studies in exactly the same paradigm (Berger et al., 2019; Eschmann et al., 2018). The spatial WM task employed by these three studies was first published in the EEG study by Griesmayr et al. (2014),¹ which also reported a theta increase.

¹not included in the review because of the clinical population.

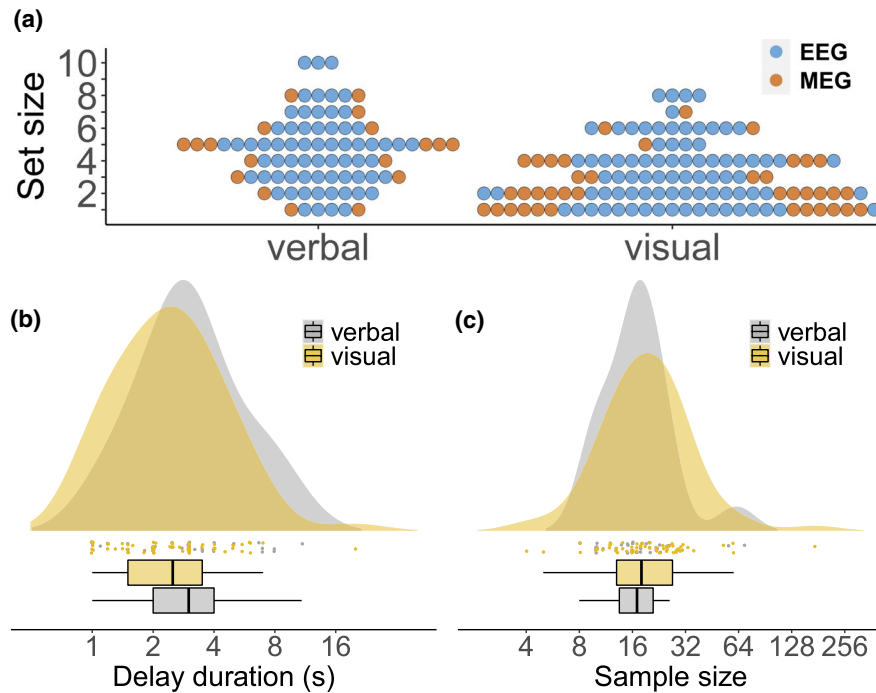


FIGURE 2 (a) Distribution of set-sizes by WM modality. Each dot represents one level of load in a study. Note that there are many more dots than studies because one study could use several levels of load. (b) Distribution of the delay period durations. (c) Distribution of sample sizes. Raincloud plots (Allen et al., 2018) were used to visualize the distributions in b and c panels. Note the logarithmic scale in panels b and c

Similar to the verbal WM study by Harmony et al. (1996), four visual WM studies reported a decrease of theta only in conditions with very low WM load. Babiloni et al. (2004) used a task where only the result of the height comparison of two clearly different bars had to be maintained, not the sensory information per se. Three other studies used a one item WM paradigm (Bastiaansen et al., 2002; de Vries et al., 2019; Kulashekhar, Pekkola, Palva, & Palva, 2016).

A similar pattern was found in the study by Rawle et al. (2012, p. 8) who wrote: “The load-dependent depression of theta during the memory delay, in a frontal distribution, agrees with the findings of Bastiaansen et al. (2002), who proposed that delay-period frontal desynchronization of theta activity is related to visuospatial working memory function.” This hypothesis is partially supported by the data above: verbal WM studies reported an increase in theta significantly more frequently than visual ones (19/24 and 14/32, respectively, $\chi^2 = 7.12$, $p = .008$). Nevertheless, Khader et al. (2010) used both modalities and found no significant differences in the theta expression.

No specific details can differentiate the last study with theta decrease (Yin et al., 2012) from similar ones where the opposite pattern was observed. Like most visual WM the study used a DMTS paradigm. Temporal order information is relevant in the Sternberg task but redundant in the DMTS type of task. This gets us back to the hypothesis on the role of theta oscillations and its relation to the maintenance of items' order in WM (Hsieh et al., 2011). If theta is, indeed,

important for the maintenance of the temporal relationship between the items in WM, then looking at the type of task may help to clarify the answer to this question.

A systematic analysis of the experimental paradigms used in the 32 visual WM studies, allowed us to subdivide them into two main groups: (1) DMTS type of task where no temporal information is stored in WM (26/32), (2) other paradigms with the temporal order information required to complete the task (6/32). 4/6 serial order paradigms and 10/26 DMTS-like paradigms reported an increase in theta activity during the delay. The proportions were not significantly different ($\chi^2 = 1.58$, $p = .209$).

3.4 | Alpha

In 27 of 35 verbal WM studies, alpha rhythm was featured in the results or depicted in figures. Twenty studies varied WM load in two or more levels. A change in alpha activity as a function of increasing load (stepwise change) was observed in 15 of them. In two cases no stepwise load-dependent alpha change in any direction was observed (Harmony et al., 1996; Kwon et al., 2015). A saturation effect was observed in two studies (Bashivan et al., 2014; Xie et al., 2016, see below). In one study the effect was neither statistically tested nor presented in a graphical form (Klimesch et al., 1993). These results generally replicate the findings in the theta frequency band—most studies report stepwise changes in alpha power.

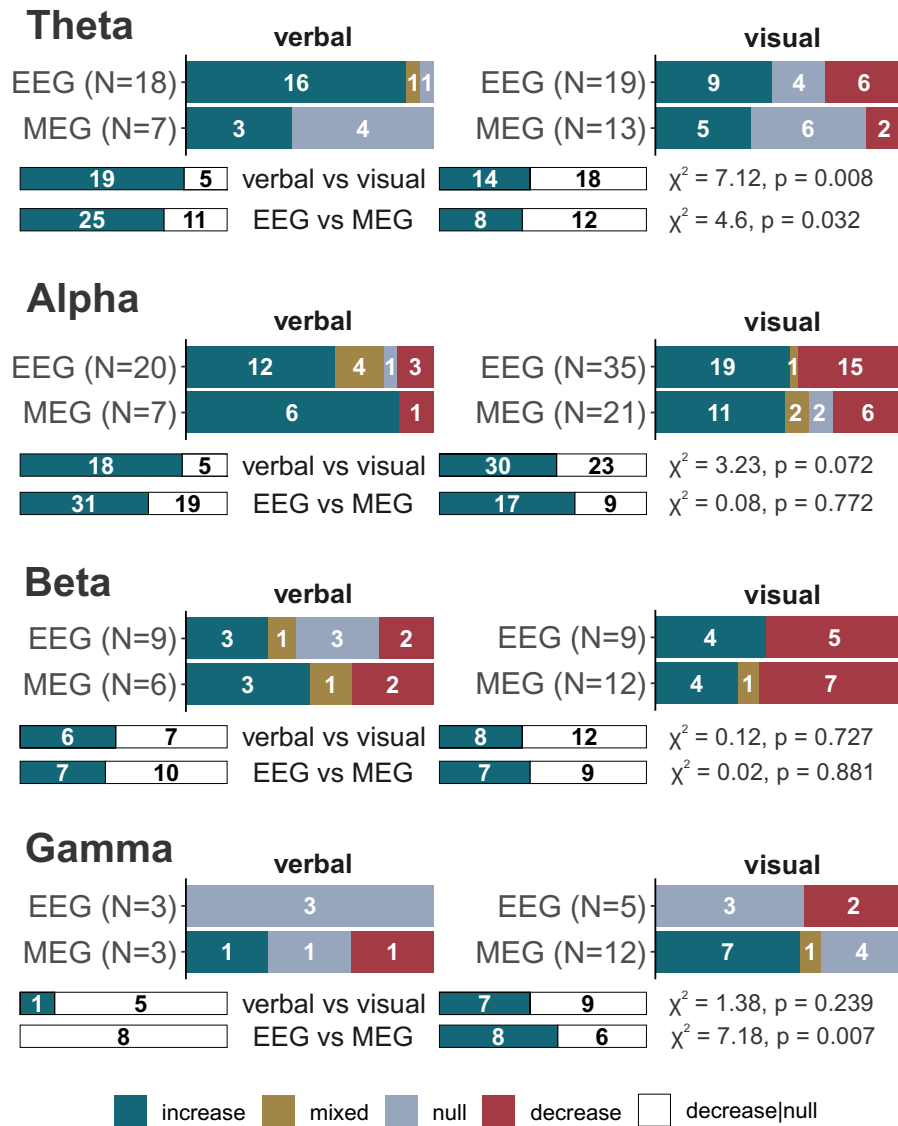


FIGURE 3 Summary of the findings in the reviewed verbal and visual WM studies. increase/decrease—changes in relative/absolute spectral power as compared with either baseline or other levels of WM load; null—no significant changes; mixed—the direction of the effect differed in either subbands, electrode locations, or groups of participants. N—number of studies in the category

However, when compared with theta activity, there was more discrepancy concerning the direction of the effects. A clear load-related increase of alpha activity was observed in 18/27 studies. In four studies there was a decrease of alpha during the delay period, and one study did not find significant changes in alpha during the delay. In the remaining six studies the results were more complex. In two studies alpha increased in the 4-items condition as compared with the 2-items condition, but no further increase in the 6- and 8-items conditions was observed (saturation effect) (Bashivan et al., 2014; Xie et al., 2016). Michels et al. (2008) distinguished two groups of subjects ($N = 9$ in each group): a group where alpha increased with WM load and another group where alpha decreased. Interestingly, but not surprisingly, in two other studies conducted by the same group on the same topic, no individual differences in alpha reactivity were

reported anymore (Kottlow et al., 2015; Michels et al., 2010). One study reported an increase of alpha in a right occipital channel but a decrease in a midline parietal channel (Kottlow et al., 2015). Similarly, in another study alpha activity increased with load in the right posterior channels but decreased in all other channels (Pavlov & Kotchoubey, 2017). Finally, Okuhata et al. (2013) found that alpha modulation depended on the type of stimulus presentation.

As already said above, stimuli in verbal WM tasks can be presented either simultaneously or successively. Okuhata et al. (2013) directly compared the two types of presentation and reported a stepwise increase of alpha in the successive presentation condition, but a decrease in the simultaneous presentation condition. In line with Okuhata et al. (2013), another group of authors also suggested that alpha activity mainly occurs in the tasks requiring the maintenance of

simultaneously presented information (Hsieh et al., 2011). We found that the majority (20/35) of the studies used simultaneous presentation and the other 14 used successive presentation. The above-mentioned study of Okuhata et al. (2013) compared both presentation types. Using only 23 studies yielding unambiguous findings, we found no significant difference in the direction of alpha modulation between simultaneous (12/14 showed an increase) and successive (6/9 showed an increase) presentation ($\chi^2 = 1.17, p = .279$). Similarly, although the number of studies in the two categories was very disproportional, visual WM studies using DMTS (26/46 showed an increase) versus serial order (4/7 showed an increase) paradigms did not differ in terms of frequency of an increase in alpha ($\chi^2 < 0.01, p = .975$).

Another hypothesis was put forward in a narrative review of van Ede (2018) who selectively compared a few studies using either visual or verbal WM tasks. The author suggested that the sign of alpha modulation depends on the nature of memoranda: visual content maintenance results in alpha suppression and verbal content maintenance leads to alpha enhancement. We tested this hypothesis in the current work. As stated above, after the exclusion of mixed results, alpha increase was found in 78% (18/23) verbal WM studies and in 57% (30/53) visual WM studies, with the difference approaching significance ($\chi^2 = 3.23, p = .072$). Thus, the increase of alpha activity is generally a frequent finding (almost 2/3 studies), but the modality of WM might partially contribute to the discrepant results.

In three visual WM studies (excluded from the analysis above) higher frequency alpha increased with load but lower frequency alpha decreased (Daume, Graetz, et al., 2017; Daume, Gruber, et al., 2017; Maurer et al., 2015). We tested whether there is a clear distinction between the directionality of the effect between lower alpha (below 10 Hz) and higher alpha (above 10 Hz) in verbal WM studies that reported a decrease during the delay. None of the six studies with alpha decrease separated alpha into subbands. In one study the time-frequency maps allowed to track alpha in the subbands but no visually observable difference between them was noticeable (Kwon et al., 2015). Then we inspected all studies

where time-frequency maps allowed us to observe changes in both alpha subbands. In short, no studies reported the sought difference between the subbands. Instead, the frequency range of the alpha effect tended to vary between studies: the effect covered both subbands in some studies (Bonfond & Jensen, 2012; Heinrichs-Graham & Wilson, 2015; Hu et al., 2019; Jensen et al., 2002; Kustermann et al., 2018), whereas it occupied higher alpha frequencies in others (Hu et al., 2019; Michels et al., 2010; Proskovec, Heinrichs-Graham, et al., 2019; Scheeringa et al., 2009; Wianda & Ross, 2019), but no studies reported specific alpha changes in the lower alpha subband. Interestingly, Proskovec, Wiesman, et al. (2019) and Heinrichs-Graham and Wilson's (2015) studies were conducted in exactly the same paradigm by the same group but the alpha effect was found either in both alpha bands or only in the higher alpha, without an explanation. Similarly, Hu et al., 2019 reported two studies with different alpha bands in one and the same article.

In some of the reviewed studies, the alpha increase during the delay was prominently right side lateralized (i.e., more alpha in the right hemisphere, see Tables 1 and 2). Note that the stimuli in these studies were always presented in the middle of the screen and there were no lateralized or spatial cues. The results of the systematic review demonstrate a prevalence of right-lateralized alpha in verbal WM tasks (see Table 1). In the EEG studies where topographical maps were reported, the alpha increase effect had no asymmetry in 3/19, right asymmetry in 16/19 studies, and no studies reported left side asymmetry. Moreover, of the 28 visual studies with a symmetrical presentation of the stimuli and available topoplots, 19 studies reported no asymmetry, and 9 reported right hemisphere asymmetry. The proportion of studies reported right side asymmetry was larger in verbal than in visual WM ($\chi^2 = 12.3, p < .001$).

Bashivan et al. (2014) among others noticed that alpha saturated with no further modulation after four items load. We tested whether this effect characterized other studies that varied load at four and higher levels of load (e.g., at least compared four vs. five items load). One MEG study and 11 EEG studies in the verbal WM domain were identified applying this criterion.

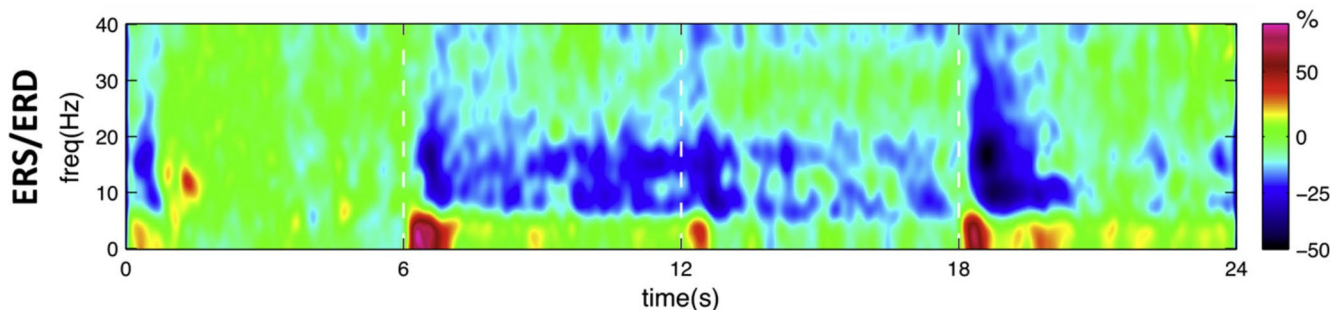


FIGURE 4 Time-frequency representation of EEG spectral power during task performance. The delay period is from 12 to 18 s. The figure is taken from Boonstra et al. (2013) with permission from Elsevier

Xie et al. (2016), similarly to Bashivan et al. (2014), showed a saturation of alpha changes in four items load. Scheeringa et al. (2009) found a significant difference between three and five but not between five and seven items load. Although Michels et al. (2008) claimed that the effect of alpha was modulated by individual differences (see above), the effect of load (any of the four vs. six vs. eight items comparisons) was not significant on the group level. Meltzer et al. (2007) failed to identify a stepwise modulation of alpha but no pair-wise comparisons were conducted. Moreover, Wilson et al. (1999) found that alpha suppression was stronger at seven and eight items load as compared with five items load. It is unclear whether there was a significant difference between seven and eight items. Five versus seven items contrast was significant in another study (Pavlov & Kotchoubey, 2017). The increase of alpha from four to six items load was significant in Jensen et al. (2002), but the effect was manifested over the central-frontal-temporal area instead of the alpha-typical posterior areas of the cortex. Similarly, another study reported that alpha increased significantly in the comparison of four and six items load (Proskovec, Heinrichs-Graham, et al., 2019). Klimesch et al. (1993), Klimesch et al. (1999) in two studies used a combination of letters and digits (5 vs. 10 items load). Both studies showed the effect of load on alpha activity during the delay. Thus, both tendencies—a saturation of alpha with load and continuation of changes—were present in the verbal WM subset.

To test the specificity of this result, we inspected the corresponding studies in the visual WM subset. Seven of 55 studies (one MEG and six EEG studies) compared at least two levels of load above three items. However, in four of the seven studies, the effect was neither formally tested by pair-wise comparisons nor it was possible to infer the effect from the figures or other analyses (Erickson et al., 2019; Moran et al., 2010; Pahor & Jausovec, 2017; Spitzer et al., 2014). In the remaining three studies the saturation effect took place. In two studies the comparison of four with six items load did not yield significance (Kawasaki & Yamaguchi, 2012; Nenert et al., 2012). In another study alpha monotonically decreased up to three item load (not even 4) (Fukuda et al., 2015).

3.5 | Beta

Fifteen of 35 studies featured beta frequency band (13–30 Hz) in the results or depicted in figures. Of these 15, three studies reported null findings, five studies showed beta suppression, and four studies reported beta increase during the delay. One study reported an increase of beta in occipital and a decrease in frontal areas (Altamura et al., 2010). In another study, beta increased with load at occipital channels but decreased with load at parietal ones (Kottlow et al., 2015). In one study higher beta increased at frontal sites in the high-performance

group but increased at posterior sites in the low-performance group (Pavlov & Kotchoubey, 2017).

In contrast to theta rhythm that primarily covers frontal cortical areas and alpha that covers posterior areas of the brain, beta rhythm has no typical spatial distribution. In five studies beta was analyzed at posterior sites (Bashivan et al., 2014; Heinrichs-Graham & Wilson, 2015; Kottlow et al., 2015; Michels et al., 2010; Pavlov & Kotchoubey, 2017), in other five studies at frontal sites (Brookes et al., 2011; Itthipuripat et al., 2013; Onton et al., 2005; Park et al., 2013; Wilson et al., 1999), and in four studies both frontal and posterior sites were included into the analysis (Altamura et al., 2010; Gao et al., 2018; Poch et al., 2010; Stephane et al., 2010). In one study (Wianda & Ross, 2019) continuous beta suppression over the left central area was shown, whereas the occipital beta behaved similarly to alpha and increased during WM delay.

Frequencies and spatial distributions of the beta effects varied substantially. Only lower beta or beta1 (typically 13–20 Hz) was analyzed in four studies (Heinrichs-Graham & Wilson, 2015; Michels et al., 2010; Onton et al., 2005; Park et al., 2013), higher beta or beta2 (20–30 Hz) in three studies (Brookes et al., 2011; Pavlov & Kotchoubey, 2017; Wilson et al., 1999), and in eight studies the whole frequency band or both subbands were inspected (Altamura et al., 2010; Bashivan et al., 2014; Gao et al., 2018; Itthipuripat et al., 2013; Kottlow et al., 2015; Poch et al., 2010; Stephane et al., 2010; Wianda & Ross, 2019).

No particular pattern of beta activation/suppression in these studies was related to the spatial distribution: beta could alternate in the frontal as well as in the posterior areas equiprobably; likewise, the frequency subband was not predictive of the direction of the effect.

The comparative results from the visual WM literature were as diverse as the results of verbal WM studies. Out of 21 visual WM studies featuring beta in the results, the majority (12) reported a decrease, but eight studies reported an increase of beta activity during the delay. One study found a decrease in the central beta but an increase of posterior beta (Park et al., 2011). In the same study alpha activity also decreased under WM load. Based on this finding we hypothesized that posterior beta might be an alpha harmonic and, therefore, share the behavior of the alpha rhythm. There were eight verbal and 16 visual WM studies where the data were specifically reported for the posterior alpha and beta. In seven verbal and 13 visual WM studies (i.e., in 83% of all studies) the alpha and beta varied in the same direction.

3.6 | Gamma

Gamma activity in verbal WM tasks is even less well studied than beta activity. Gamma results were featured

in three of 25 EEG and three of 10 MEG studies. In two EEG studies, null findings were reported. One study reported an increase in gamma activity during the delay. Of the three MEG studies, one demonstrated a decrease of beta/low gamma oscillatory power with increasing WM load (Brookes et al., 2011), one study showed, in contrast, an increase of gamma activity (Poch et al., 2010), and one study did not reveal any changes in gamma with load (Kustermann et al., 2018).

In the visual domain, the sample was larger. Five visual WM studies using EEG featured gamma activity in the results. Two of them yielded null findings (Rawle et al., 2012; Zhang et al., 2016). Gamma decreased in two other studies (Pahor & Jausovec, 2017; Smyrnis et al., 2014). In one study gamma quickly disappeared after an initial (~800 ms) increase (Tallon-Baudry et al., 1999). Twelve similar studies were carried out with MEG, seven of which reported an increase in gamma during WM maintenance. No changes were found in four studies. In one MEG study, gamma increased over posterior and right temporal regions but decreased over left temporal and central areas (Park et al., 2011).

Thus, summarizing over verbal and visual domain studies, the results reveal a strong difference between the two recording modalities: a sustained increment of gamma activity was obtained in most (8/14) MEG studies but in one (1/8) EEG study ($\chi^2 = 4.19, p = .04$).

4 | DISCUSSION

4.1 | Theta

4.1.1 | Temporal order hypothesis

The importance of frontal midline theta in verbal WM maintenance was supported by the majority of the reviewed studies: theta power was higher during the delay period in comparison with baseline.

Presumably, theta activity plays a role in the maintenance of the temporal relationship between the items in memory: storage of a larger number of items makes the relationship more complex (Hsieh & Ranganath, 2014). Another study conducted by the same group supports this idea (Roberts et al., 2013). In this study, the participants required to remember either location of four presented sequentially abstract images or the temporal order of the images. The order condition generated stronger theta power.

It appears to be difficult to design a task that could provide sufficient evidence for or against this hypothesis in the verbal domain. Moreover, even presented simultaneously verbal content is rehearsed in a sequence. Nonetheless, we suggest that the hypothesis can be tested in the verbal WM

domain in three ways. First, by comparison of single-item WM load, when no temporal relationship between items exists, with higher load conditions. Onton et al. (2005) demonstrated the expected pattern of load-dependent increase of theta power (see Figure 5). Unfortunately, the delay period after the presentation of the items was too short to make a reliable conclusion. Out of 56 verbal and visual WM available studies, this is the only one where this kind of comparison could be made. Second, the explicit task to memorize order and content in one condition and to memorize only content in another one could shed some light on the matter. For example, sufficient evidence could come out of a comparison of the classic Sternberg paradigm with a single item probe and a modified Sternberg paradigm with a requirement to remember both content and order, similar to the ones used in Pavlov and Kotchoubey (2017) or Wianda and Ross (2019). To date, there are no such studies. Third, more studies manipulating WM load in a wide range can provide additional evidence in support for the hypothesis, provided that the difficulty of the task is equalized over conditions.

4.2 | Alpha

4.2.1 | Directionality

Alpha activity was another popular target in the reviewed studies. In contrast to theta, the directionality of alpha modulation varied: after the exclusion of mixed and null findings, about 20% of the verbal WM studies detected a decrease and 80% an increase in alpha power during the delay. The proportion of visual WM studies was close to 40/60%. To explain this discrepancy, we tested several hypotheses and rejected all of them with different degrees of certainty. On the one hand, the modality (visual vs. verbal) and type of presentation (simultaneous vs. successive) failed to fully explain the alpha directionality. On the other hand, there is not enough information on the impact of alpha subband frequency (lower vs. upper alpha) and individual differences in the direction of changes in alpha power.

Alpha suppression has been seen as a sign of cortical engagement allowing either the encoding of information into WM or decoding the information for retrieval (Jensen & Mazaheri, 2010; Klimesch et al., 2007). When simultaneously presented verbal stimuli remain on the screen for a long time (3s or longer) alpha is continuously suppressed over the encoding period (Bailey et al., 2014; Murphy et al., 2019; Scheeringa et al., 2009; Segrave et al., 2010). Whereas alpha increase during the delay may reflect sensory gating through the disengagement of certain cortical areas to protect memory representations from interference (Klimesch et al., 2007; Payne & Sekuler, 2014; Roux & Uhlhaas, 2014). In the case of

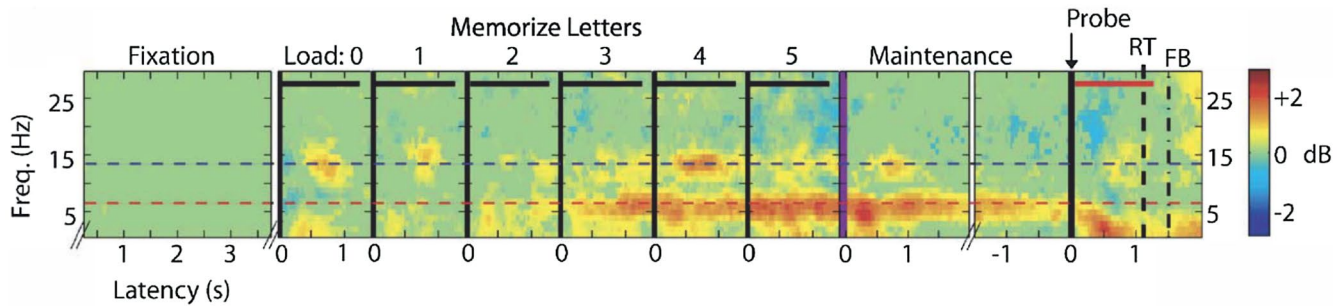


FIGURE 5 Development of FMT enhancement with increasing load from 0 to 5 items in a verbal WM task. Black horizontal bars represent the encoding period (the figure is adapted from Onton et al. (2005) with permission from Elsevier)

visual presentation, alpha supports the storage of the memory items by blocking visual input. We can hypothesize that the cases when alpha activity is suppressed during the delay are the cases when visual information processing is still ongoing.

Attention is able to selectively suppress alpha activity spatially related to a specific memory representation (Fukuda et al., 2015; Schneider et al., 2019; de Vries et al., 2018). For example, when internal attention is directed with a previously presented cue (retro-cue) to the left hemifield items maintained in WM, right posterior alpha activity is attenuated (Myers et al., 2015; Sauseng et al., 2009; de Vries et al., 2018; Wolff et al., 2017; Worden et al., 2000). Thus, although visual input is blocked during the delay, previously presented spatial cues may direct internal attention and affect alpha activity.

A few properties of these studies complicate the interpretation of the alpha suppression in WM tasks with retro-cues. First, most of the above-mentioned studies used short delays which do not help to separate pure oscillatory effects from ERPs. Second, a lot of these kinds of studies only report the index of lateralized alpha (for this reason they were excluded from the review). Without having access to spectral power data from both hemispheres, it is difficult to say whether alpha was generally suppressed during the delay or less strongly enhanced in the contralateral hemisphere.

We can assume that in a typical WM task without retro-cues (e.g., Sternberg task), the item currently in the focus of attention suppresses alpha, while the items currently outside the focus enhance alpha at the same time. Both operations may engage such local neuronal populations that M/EEG cannot spatially differentiate them. But what is happening when there is only one item to store? Then alpha shows a small magnitude increase anyway (e.g., see Figure 6).

The existing data exhibit a striking contrast between a large number of available studies and the inability to predict, not the magnitude, but even the direction of alpha modulation. Before we start building a mechanistic foundation of the role of alpha activity in the maintenance of WM, more replication studies are deemed necessary.

4.2.2 | Asymmetry

A stronger increase of alpha in the right hemisphere is common in many verbal WM tasks. If we accept the mainstream interpretation, an increase of alpha during the delay serves to prevent the sensory cortex from further processing of information, thus protecting stored memory from interference (Bonfond & Jensen, 2012; Foxe & Snyder, 2011; Jensen & Mazaheri, 2010; Klimesch et al., 2007). Putting it differently, the current view suggests that alpha represents a mechanism of the active inhibition of irrelevant cortical areas (Klimesch et al., 2007).

But why should the activity of the right temporo-parieto-occipital region during a WM task performance be irrelevant? We suggest that the maintenance of verbal information presented visually dissociates two simultaneous demands: blockage of visual input by disengaging visual cortex and activation of the language cortex for rehearsal. The rehearsal processes involve left posterior regions, which are reflected in the suppression of alpha in the left hemisphere. The opposite task to inhibit distracting visual input enhances alpha activity in both hemispheres.

The same kind of alpha asymmetry was found in visual WM studies, though significantly less frequently. The mechanism generating the asymmetry in non-verbal tasks is less clear. Johnson et al. (2011) used tasks requiring either memorizing shapes of objects or their location. The right-side (left < right) alpha asymmetry revealed itself only in the shape condition, while alpha was symmetrical in the location condition. Moreover, alpha was generally stronger in the shape condition. Perhaps, maintenance of spatial information suppressed alpha in the right hemisphere thus equalizing it over the posterior cortical areas. However, Bastiaansen et al. (2002) showed stronger alpha over the right occipital area in a spatial task. Although the prevalence of right-side alpha asymmetry among all reviewed studies associated with the content stored in memory (verbal or visuospatial), the origin of the alpha asymmetry in visual WM demands more systematic investigation.

The natural alpha asymmetry in the resting state also might contribute to the effect (Allen & Cohen, 2010; Ocklenburg et al., 2019; Stewart et al., 2014; van der Vinne et al., 2017). However, the effect was also present in many studies where alpha power was normalized on the baseline. Thus, the asymmetry during the delay was a sign of a genuine increase in alpha asymmetry, not preservation of the status quo.

4.2.3 | Saturation

In some of the reviewed studies, alpha power did not change in WM load beyond the theoretically predicted (Cowan, 2001; Oberauer et al., 2018) average individual WM capacity limit of four items. This reminds the behavior of P300 that also stops to change in higher levels of WM load in the classical version of the Sternberg task (Kotchoubey, 2002). Another ERP component, contralateral delay activity (CDA), also saturated in the change detection paradigm (Luria et al., 2016). Similarly, fMRI BOLD signal increased monotonically with load up to the individual's WM capacity that averaged at four items set-size (Todd & Marois, 2004). Although some studies confirmed the saturation of alpha in verbal WM, the effect was not as consistent as in the case of CDA.

In the visual domain, all the identified studies showed the saturation of non-lateralized alpha (as opposed to lateralized alpha in the lateralized change detection paradigm) at higher levels of load. It should be taken into account, however, that only a few visual WM studies used set-sizes above four items.

If it is true that the saturation of alpha is necessary in visual WM tasks but can sometimes be avoided in verbal WM tasks, this difference may reflect an effect of chunking, that is, the grouping of WM items (chunks) into larger units. Thus, for example, the task to encode a string of nine digits is feasible because the string is automatically converted into 3 three-digit numbers. The four items WM limit proposed by Cowan (2001) refers to a condition when rehearsal and chunking are restricted. When rehearsal of verbal material such as digits or letters is not blocked, then the capacity limit reaches the number of items an individual can rehearse in about 2 s (Baddeley, 1996; Cowan, 2001). The capacity increases further if chunking is possible. These are, probably, the reasons why set-sizes above four items, as frequently used in verbal WM research, do not overload average WM capacity.

4.3 | Beta

Beta oscillations in verbal WM research are much less studied than alpha and theta. The review has shown a wide diversity of effects in the beta frequency band. An

assumption that there is probably more than one beta rhythm may contribute to the explanation of the discrepancy in the effects. These beta rhythms may occupy different frequencies, have different temporal and spatial distributions, and, therefore, different functional meanings. The data present strong evidence that occipital beta may be a harmonic of alpha and does not play an independent functional role. The role of beta oscillations outside occipital regions in WM remains unclear.

4.4 | Gamma

Even though intracranial EEG (iEEG) and MEG repeatedly demonstrated the robustness of gamma responses in cognitive tasks (Herrmann et al., 2010; Jerbi et al., 2009), gamma obtained from the EEG signal is less robust. The reason for this is a strong contamination of EEG by muscle artifacts mainly occupying higher frequencies including the gamma frequency band (Muthukumaraswamy, 2013). Generally worse signal-to-noise ratio in the gamma band of the EEG signal may explain less attention to gamma oscillations in EEG in verbal WM tasks. Interestingly, only a single EEG study reported a gamma increase during the delay in the combined whole dataset of 65 studies employing this method. Contradicting our expectations, a few EEG studies even reported a decrease in gamma power.

To understand how the signal-to-noise ratio might have affected our results we extracted data from a representative sample of iEEG studies. To achieve that, we used similar criteria applied to the M/EEG dataset to identify the iEEG studies featuring gamma rhythm in the results. In sharp contrast to EEG, the increase of gamma power with load was confirmed in six of nine available verbal WM iEEG studies (Bahramisharif et al., 2018; Gehrig et al., 2019; Howard et al., 2003; Kambara et al., 2017; Mainy et al., 2007; Meltzer et al., 2008). Even this high percentage (67% “positive” findings) can be an underestimation. Of the three studies with “negative” findings, two studies recorded iEEG from the hippocampus (Boran et al., 2019; Leszczyński et al., 2015), which cannot be directly compared with the cortical iEEG. Finally, the last study (Raghavachari et al., 2001) briefly mentioned that gamma activity was affected by load but the results are reported elsewhere. Unfortunately, we were unable to identify the referenced study.

The present systematic review includes only studies performed with healthy participants unaffected by any drugs. The compared group of the nine iEEG studies involved, in contrast, neurological (mainly epileptic) patients, most of them were medicated. Although the difference is important, we cannot imagine how epilepsy or antiepileptic drugs might improve gamma responses making them more clear and consistent. Moreover, the iEEG data are corroborated by the

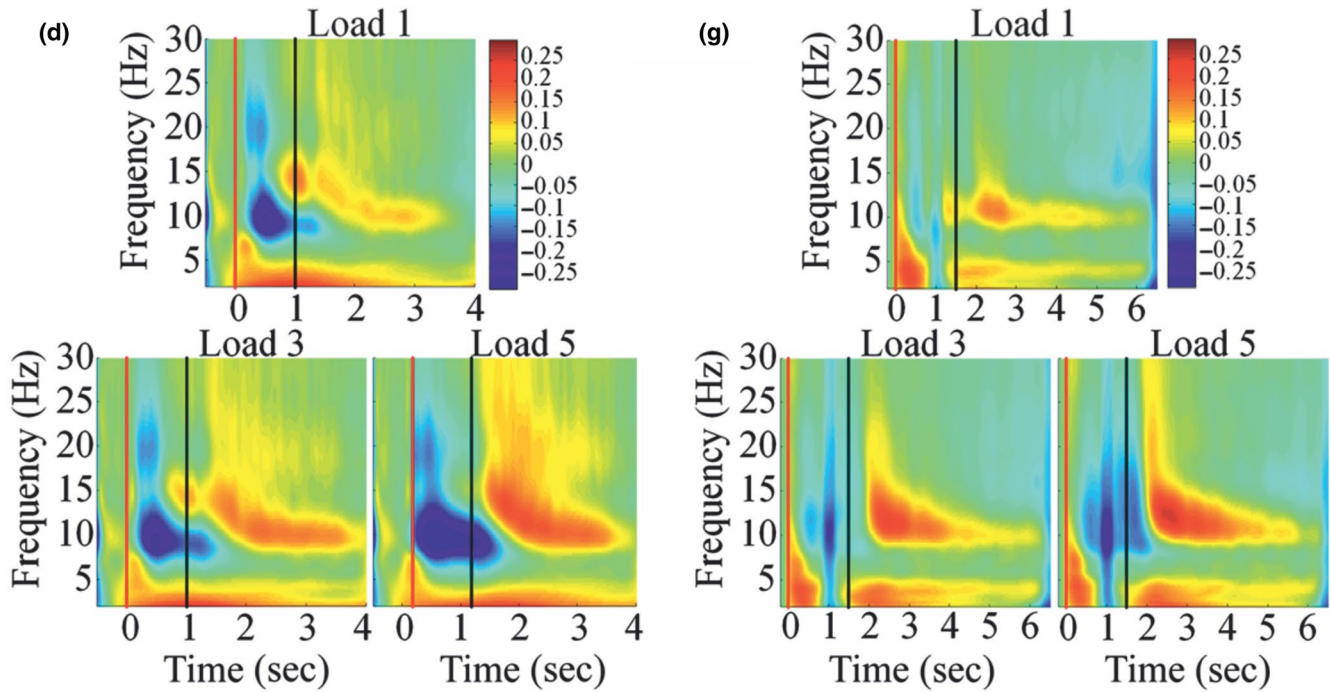


FIGURE 6 Alpha activity during the delay in two verbal WM paradigms (D: Sternberg, G: DMTS). Adopted from figure 3 in Hu et al. (2019) with permission from MIT Press. The black vertical lines mark the beginning of the delay period

results of MEG studies, 60% of which showed an increase in gamma activity during the delay. However, most of those included in this analysis studies used visual WM tasks, whereas the sample of verbal WM studies featuring gamma was small.

4.5 | EEG versus MEG

Alpha and beta bands did not differentiate the results obtained by any of the methods but theta and gamma did.

Whereas iEEG and partially MEG linked sustained gamma with the maintenance of WM, EEG failed to provide such evidence. Only a single EEG study reported some gamma modulation during WM delay. Until more convincing evidence is available, the whole enterprise of extracting meaningful quality data from the EEG in the gamma frequency band raises doubt. Perhaps, we even have a reason to be skeptical about the studies employing different approaches such as theta-gamma cross-frequency coupling.

EEG and MEG disagreed less strikingly but sufficiently in the theta band: EEG verbal WM studies almost unanimously showed an effect of increased theta during the delay, but MEG findings exhibited less consistency. A possible reason for this difference may be the specific sensitivity of the MEG signal to the orientation of the current dipole. MEG is sensitive to the activity generated in sulci of the cortex (i.e., in sources tangential to the head surface) but blind to radially oriented sources. The strongest theta sources are located in the medial prefrontal cortex and directed perpendicularly to the scalp.

This distribution of the magnetic fields prevents it from being registered by MEG. A simultaneous M/EEG recording in the Simon task (a task that reliably generates FMT in EEG) showed a less focused, more lateral, and individually different patterns of theta activity in MEG (Zuure et al., 2020).

The results in gamma and theta bands highlight the difference between EEG and MEG in the methodology and probably in the underlying biophysical mechanisms. At least in some cases, results obtained by MEG and EEG should be interpreted interchangeably with caution.

4.6 | Verbal versus Visual WM

In the final section, we ask: Does WM for visually presented verbal stimuli differ from visual WM? Or can verbal WM, on the basis of the electrophysiological data, be considered as a special case of visual WM? The answer is, paralleling behavioral studies (Oberauer et al., 2018), electrophysiology of verbal WM differs from that of visual WM.

Verbal WM studies reported an increase in theta significantly more frequently than visual ones (19/24 and 14/32, respectively, $\chi^2 = 7.12$, $p = .008$). Moreover, eight visual WM studies even indicated a significant decrease in theta. No such finding was found in the verbal WM domain. A hypothesis is that theta is important for the maintenance of temporal order which is of stronger demand in the verbal tasks.

In contrast to the significant effect of modality on theta modulation, a similar difference in the alpha band was less

convincing. However, right hemisphere asymmetry in alpha power (right > left) was reported more frequently in the verbal domain. This effect is the most prominent influence of the verbal content that supposedly activates primarily left located language cortical areas, thus suppressing alpha in the left, while enhancing alpha in the right hemisphere. Another difference may be the saturation effect, which was less prominent in the verbal than in the visual domain; but we should keep in mind that the sample size of this subset of studies was rather small.

Considering a large diversity in frequency and spatial characteristics of beta activity it might be too preliminary to make any certain conclusion about verbal/visual differences in beta oscillatory responses. Nevertheless, the delay beta power showed a similar increase in 7/20 of visual and 4/15 of verbal studies ($\chi^2 = 0.12$, $p = .727$).

Finally, as compared to verbal, more visual gamma studies (1) featured the rhythm in the results, and (2) reported an increase of gamma power during the delay (although not significantly, verbal: 2/6, visual: 7/16, $\chi^2 = 0.19$, $p = .658$). We cannot rule out that the latter depended on the former, that is, the null results were simply not reported. The question of whether gamma activity is more associated with visual than with verbal information processing remains difficult to answer. Such studies cannot be conducted in animals, and human iEEG study would require recordings from larger areas than possible and ethically justifiable (Parvizi & Kastner, 2018). More MEG studies engaging different WM modalities may provide such evidence.

To summarize, a number of differences between the modalities exist: verbal WM is left-lateralized (left < right alpha power), probably, has a larger capacity, and is more dependent on theta activity.

4.7 | Limitations, challenges, and perspectives

A certain level of simplification and generalization is inevitable in any review of the literature. The issue is even more severe in a systematic review aiming to accommodate as many studies as possible, while maintaining objectivity and precision in the quantification of the results. This holds true even if the quantification involves only identifying the direction of the effect with three viable options. More thorough data quantification would necessarily mean taking steps to decrease heterogeneity, resulting in a much smaller sample and narrower scope of the review. We admit that we are only scratching the surface of the literature by the application of this simplistic approach, and the current work might be only a small step away from a pure narrative review. But reading those hundreds of M/EEG papers published over the years has left an impression that most discussion sections mainly

compare their results with similar previous findings, avoiding even to mention studies that obtained the opposite effect. Our review creates a context for future research to make this behavior less likely to occur.

A review can only analyze variables that vary from study to study systematically, but not chaotically. We thus considered only factors that could build reasonably large groups of studies. Even this cautious approach resulted in some unbalanced comparisons. Other smaller factors may contribute to the inconsistencies in the reviewed findings. Potentially, the decisions in the analysis pipeline such as length of the time window used in Fourier transform (or other similar parameters for Hilbert transform or wavelet analysis), analysis time window, filter settings, number of trials, selection of the channels, and reference scheme, duration of baseline, definition of the frequency bands (e.g., alpha can vary in the range between 8 and 13, 7 and 14, 10 and 12 Hz, or can be selected on the basis of individual alpha frequency), application of independent component analysis (ICA) (to source or sensor space data) for source separation and many other factors could affect the results. The heterogeneity of the studies included in the review prevented us from assessing the impact of these factors.

A pure description of the analysis pipelines is also beyond the scope of the current review. Descriptive data are only helpful if the experimental paradigms in the original studies are almost identical. Thus, selecting studies using the same experimental paradigm is an essential step in a search of the methodological aspects contributing to the variability of results. A promising example of such a study looked at all possible ways to analyze N400 ERP in the visual modality in a sample of 132 studies (Šoškić et al., 2020). Another possibility to solve this problem is to apply a multiverse analysis to the raw data. Going in this direction would involve the deep participation of the original authors and/or broader adoption of the data-sharing practices.

Conflicting findings even within one study, missing methodological details, and a vague description of the results may lead to misrepresentation or imprecise the quantification of studies in any review of the literature. We believe that—devised by an authoritative board of M/EEG and WM experts—methodological recommendations would benefit the field by mitigating the negative effects of reporting inconsistency among studies. To ensure an extensive adoption, these recommendations should necessarily be supplemented by simple machine- and human-readable checklists/interactive forms. Of course, the present article never aimed to develop such recommendations.

Apart from widely adopted analytic approaches to brain oscillations, such as FFT and wavelet analysis, novel techniques have been proposed, some of them quite recently (Cole & Voytek, 2019; Donoghue et al. (in press); Kosciessa et al., 2020). On the one hand, we cannot rule it out that one of these methods

would make a breakthrough in the study of cognitive functions, and particularly working memory; on the other hand, singular adoption of them will further increase the already high methodological variability, and further decrease the comparability of results. The experience tells us that cognitive neuroscientists rarely hasten to embrace novel computational techniques after the release. Whether we should regard this conservatism as an advantage or disadvantage remains unclear.

4.8 | Conclusions

1. In the pool of 35 identified verbal and 65 visual WM studies, oscillatory EEG responses in theta, alpha, beta, and gamma frequency bands were analyzed in 25, 27, 15, 6 verbal and 32, 56, 21, 17 visual WM studies, respectively. A significant amount of the reviewed literature supports the role of theta and alpha oscillations in WM though much less is known about the role of beta and particularly gamma activity.
2. Theta activity increases with WM load in verbal tasks, whereas this trend is less pronounced in visual WM tasks. This may be related to the larger role of theta in building a temporal structure for the maintenance of multiple items in verbal WM.
3. Alpha activity changes with increasing WM load, but the direction of the effect is not consistent across the studies. In approximately 20% and 40% of the verbal and visual studies, respectively, alpha decreased with load, while in the other studies it increased. The direction of the modulation is not fully explained by the stimulus modality, type of the stimulus presentation (simultaneous or successive), and, as far as it is possible to judge on the basis of few studies, on individual differences. The reason for this fact remains to be elucidated.
4. The distribution of posterior alpha power had right hemisphere asymmetry (left < right) in the majority of the studies, even if sensory input was symmetrical. Significantly smaller amounts of visual WM studies showed this asymmetry. The effect is probably related to the leading role of the left hemisphere in verbal information processing.
5. The saturation of alpha at the theoretical limit of WM capacity (i.e., alpha responses change with WM load from one to four items, but remain stable above the 4-item level) was found in several, but not in all verbal WM studies, and the reasons of the differences remain unclear. The saturation effect at four items load is more consistent in the visual WM literature.
6. Many MEG and iEEG studies report incrementing gamma activity during WM performance whereas EEG studies do not. This may indicate that the scalp EEG is not the best tool to study gamma oscillations in WM tasks.

ACKNOWLEDGMENTS

The study was supported by the Russian Foundation for Basic Research (RFBR) #19-013-00027. Open access funding enabled and organized by Projekt DEAL.

AUTHOR CONTRIBUTIONS

Yuri G. Pavlov: Conceptualization; Data curation; Formal analysis; Funding acquisition; Investigation; Methodology; Visualization; Writing-original draft; Writing-review & editing. **Boris Kotchoube:** Writing-review & editing.

ORCID

Yuri G. Pavlov  <https://orcid.org/0000-0002-3896-5145>

Boris Kotchoube  <https://orcid.org/0000-0002-6605-4744>

REFERENCES

- Allen, J. J. B., & Cohen, M. X. (2010). Deconstructing the “resting” state: Exploring the temporal dynamics of frontal alpha asymmetry as an endophenotype for depression. *Frontiers in Human Neuroscience*, 4. <https://doi.org/10.3389/fnhum.2010.00232>
- Allen, M., Poggiali, D., Whitaker, K., Marshall, T. R., & Kievit, R. (2018). *Raincloud plots: A multi-platform tool for robust data visualization* (No. e27137v1). <https://doi.org/10.7287/peerj.preprints.27137v1>
- Altamura, M., Goldberg, T. E., Elvevag, B., Holroyd, T., Carver, F. W., Weinberger, D. R., & Coppola, R. (2010). Prefrontal cortex modulation during anticipation of working memory demands as revealed by magnetoencephalography. *International Journal of Biomedical Imaging*, 2010, 1–10. <https://doi.org/10.1155/2010/840416>
- Babiloni, C., Babiloni, F., Carducci, F., Cappa, S. F., Cincotti, F., Del Percio, C., Miniussi, C., Moretti, D. V., Rossi, S., Sosta, K., & Rossini, P. M. (2004). Human cortical responses during one-bit short-term memory. A high-resolution EEG study on delayed choice reaction time tasks. *Clinical Neurophysiology*, 115(1), 161–170. [https://doi.org/10.1016/S1388-2457\(03\)00286-4](https://doi.org/10.1016/S1388-2457(03)00286-4)
- Babiloni, C., Babiloni, F., Carducci, F., Cincotti, F., Del Percio, C., Della Penna, S., Franciotti, R., Pignotti, S., Pizzella, V., Rossini, P. M., Sabatini, E., Torquati, K., & Romani, G. L. (2005). Human alpha rhythms during visual delayed choice reaction time tasks: A magnetoencephalography study. *Human Brain Mapping*, 24(3), 184–192. <https://doi.org/10.1002/hbm.20079>
- Baddeley, A. (2003). Working memory: Looking back and looking forward. *Nature Reviews Neuroscience*, 4(10), 829–839. <https://doi.org/10.1038/nrn1201>
- Baddeley, A. D., Hitch, G. J., & Allen, R. J. (2019). From short-term store to multicomponent working memory: The role of the modal model. *Memory & Cognition*, 47(4), 575–588. <https://doi.org/10.3758/s13421-018-0878-5>
- Bahramsharif, A., Jensen, O., Jacobs, J., & Lisman, J. (2018). Serial representation of items during working memory maintenance at letter-selective cortical sites. *PLOS Biology*, 16(8), e2003805. <https://doi.org/10.1371/journal.pbio.2003805>
- Bailey, N. W., Segrave, R. A., Hoy, K. E., Maller, J. J., & Fitzgerald, P. B. (2014). Impaired upper alpha synchronisation during working memory retention in depression and depression following traumatic

- brain injury. *Biological Psychology*, 99, 115–124. <https://doi.org/10.1016/j.biopsycho.2014.03.008>
- Bashivan, P., Bidelman, G. M., & Yeasin, M. (2014). Spectrotemporal dynamics of the EEG during working memory encoding and maintenance predicts individual behavioral capacity. *European Journal of Neuroscience*, 40(12), 3774–3784. <https://doi.org/10.1111/ejn.12749>
- Bastiaansen, M. C. M., Posthuma, D., Groot, P. F. C., & de Geus, E. J. C. (2002). Event-related alpha and theta responses in a visuo-spatial working memory task. *Clinical Neurophysiology*, 113(12), 1882–1893. [https://doi.org/10.1016/S1388-2457\(02\)00303-6](https://doi.org/10.1016/S1388-2457(02)00303-6)
- Berger, B., Griesmayr, B., Minarik, T., Biel, A. L., Pinal, D., Sterr, A., & Sauseng, P. (2019). Dynamic regulation of interregional cortical communication by slow brain oscillations during working memory. *Nature Communications*, 10(1), 1–11. <https://doi.org/10.1038/s41467-019-12057-0>
- Blackler, K. J., Ikkai, A., Lakshmanan, B. M., Ewen, J. B., & Courtney, S. M. (2016). The role of alpha oscillations in deriving and maintaining spatial relations in working memory. *Cognitive, Affective & Behavioral Neuroscience*, 16(5), 888–901. <https://doi.org/10.3758/s13415-016-0439-y>
- Bonfond, M., & Jensen, O. (2012). Alpha oscillations serve to protect working memory maintenance against anticipated distracters. *Current Biology*, 22(20), 1969–1974. <https://doi.org/10.1016/j.cub.2012.08.029>
- Boonstra, T. W., Powell, T. Y., Mehrkanoon, S., & Breakspear, M. (2013). Effects of mnemonic load on cortical activity during visual working memory: Linking ongoing brain activity with evoked responses. *International Journal of Psychophysiology*, 89(3), 409–418. <https://doi.org/10.1016/j.ijpsycho.2013.04.001>
- Boran, E., Fedele, T., Klaver, P., Hilfiker, P., Stieglitz, L., Grunwald, T., & Sarnthein, J. (2019). Persistent hippocampal neural firing and hippocampal-cortical coupling predict verbal working memory load. *Science Advances*, 5(3), eaav3687. <https://doi.org/10.1126/sciadv.aav3687>
- Brookes, M. J., Liddle, E. B., Hale, J. R., Woolrich, M. W., Luckhoo, H., Liddle, P. F., & Morris, P. G. (2012). Task induced modulation of neural oscillations in electrophysiological brain networks. *NeuroImage*, 63(4), 1918–1930. <https://doi.org/10.1016/j.neuroimage.2012.08.012>
- Brookes, M. J., Wood, J. R., Stevenson, C. M., Zumer, J. M., White, T. P., Liddle, P. F., & Morris, P. G. (2011). Changes in brain network activity during working memory tasks: A magnetoencephalography study. *NeuroImage*, 55(4), 1804–1815. <https://doi.org/10.1016/j.neuroimage.2010.10.074>
- Busch, N. A., & Herrmann, C. S. (2003). Object-load and feature-load modulate EEG in a short-term memory task. *NeuroReport*, 14(13), 1721–1724. <https://doi.org/10.1097/00001756-200309150-00013>
- Bushman, B. J., & Wells, G. L. (2001). Narrative impressions of literature: The availability bias and the corrective properties of meta-analytic approaches. *Personality and Social Psychology Bulletin*, 27(9), 1123–1130. <https://doi.org/10.1177/0146167201279005>
- Buzsaki, G. (2004). Neuronal oscillations in cortical networks. *Science*, 304(5679), 1926–1929. <https://doi.org/10.1126/science.1099745>
- Chen, Y., & Huang, X. (2015). Modulation of alpha and beta oscillations during an n-back task with varying temporal memory load. *Frontiers in Psychology*, 6, 2031. <https://doi.org/10.3389/fpsyg.2015.02031>
- Cole, S., & Voytek, B. (2019). Cycle-by-cycle analysis of neural oscillations. *Journal of Neurophysiology*, 122(2), 849–861. <https://doi.org/10.1152/jn.00273.2019>
- Cowan, N. (2001). The magical number 4 in short-term memory: A reconsideration of mental storage capacity. *Behavioral and Brain Sciences*, 24(1), 87–114. <https://doi.org/10.1017/S0140525X01003922>
- Crespo-Garcia, M., Pinal, D., Cantero, J. L., Diaz, F., Zurrón, M., & Atienza, M. (2013). Working memory processes are mediated by local and long-range synchronization of alpha oscillations. *Journal of Cognitive Neuroscience*, 25(8), 1343–1357. https://doi.org/10.1162/jocn_a_00379
- D'Esposito, M., & Postle, B. R. (2015). The cognitive neuroscience of working memory. *Annual Review of Psychology*, 66(1), 115–142. <https://doi.org/10.1146/annurev-psych-010814-015031>
- Daniel, T. A., Katz, J. S., & Robinson, J. L. (2016). Delayed match-to-sample in working memory: A BrainMap meta-analysis. *Biological Psychology*, 120, 10–20. <https://doi.org/10.1016/j.biopsycho.2016.07.015>
- Daume, J., Graetz, S., Gruber, T., Engel, A. K., & Fries, U. (2017). Cognitive control during audiovisual working memory engages frontotemporal theta-band interactions. *Scientific Reports*, 7(1), 12585. <https://doi.org/10.1038/s41598-017-12511-3>
- Daume, J., Gruber, T., Engel, A. K., & Fries, U. (2017). Phase-amplitude coupling and long-range phase synchronization reveal frontotemporal interactions during visual working memory. *The Journal of Neuroscience*, 37(2), 313–322. <https://doi.org/10.1523/JNEUROSCI.2130-16.2016>
- de Vries, I. E. J., Savran, E., van Driel, J., & Olivers, C. N. L. (2019). Oscillatory mechanisms of preparing for visual distraction. *Journal of Cognitive Neuroscience*, 31(12), 1873–1894. https://doi.org/10.1162/jocn_a_01460
- de Vries, I. E. J., van Driel, J., Karacaoglu, M., & Olivers, C. N. L. (2018). Priority switches in visual working memory are supported by frontal delta and posterior alpha interactions. *Cerebral Cortex*, 28(11), 4090–4104. <https://doi.org/10.1093/cercor/bhy223>
- de Vries, I. E. J., van Driel, J., & Olivers, C. N. L. (2017). Posterior α EEG dynamics dissociate current from future goals in working memory-guided visual search. *Journal of Neuroscience*, 37(6), 1591–1603. <https://doi.org/10.1523/jneurosci.2945-16.2016>
- Deiber, M.-P., Missonnier, P., Bertrand, O., Gold, G., Fazio-Costa, L., Ibanez, V., & Giannakopoulos, P. (2007). Distinction between perceptual and attentional processing in working memory tasks: A study of phase-locked and induced oscillatory brain dynamics. *Journal of Cognitive Neuroscience*, 19(1), 158–172. <https://doi.org/10.1162/jocn.2007.19.1.158>
- Donoghue, T., Haller, M., Peterson, E., Varma, P., Sebastian, P., & Gao, R., & Voytek, B. (in press). Parameterizing neural power spectra into periodic and aperiodic components. *Nature Neuroscience*.
- Ellmore, T. M., Ng, K., & Reichert, C. P. (2017). Early and late components of EEG delay activity correlate differently with scene working memory performance. *PLoS One*, 12(10), e0186072. <https://doi.org/10.1371/journal.pone.0186072>
- Erickson, M. A., Smith, D., Albrecht, M. A., & Silverstein, S. (2019). Alpha-band desynchronization reflects memory-specific processes during visual change detection. *Psychophysiology*, 56(11), <https://doi.org/10.1111/psyp.13442>
- Eschmann, K. C. J., Bader, R., & Mecklinger, A. (2018). Topographical differences of frontal-midline theta activity reflect functional



- differences in cognitive control abilities. *Brain and Cognition*, *123*, 57–64. <https://doi.org/10.1016/j.bandc.2018.02.002>
- Fell, J., & Axmacher, N. (2011). The role of phase synchronization in memory processes. *Nature Reviews Neuroscience*, *12*(2), 105–118. <https://doi.org/10.1038/nrn2979>
- Foxe, J. J., & Snyder, A. C. (2011). The role of alpha-band brain oscillations as a sensory suppression mechanism during selective attention. *Frontiers in Psychology*, *2*. <https://doi.org/10.3389/fpsyg.2011.00154>
- Fukuda, K., Mance, I., & Vogel, E. K. (2015). Alpha power modulation and event-related slow wave provide dissociable correlates of visual working memory. *The Journal of Neuroscience*, *35*(41), 14009–14016. <https://doi.org/10.1523/jneurosci.5003-14.2015>
- Gao, C., Sun, J., Yang, X., & Gong, H. (2018). Gender differences in brain networks during verbal Sternberg tasks: A simultaneous near-infrared spectroscopy and electro-encephalography study. *Journal of Biophotonics*, *11*(3), e201700120. <https://doi.org/10.1002/jbio.201700120>
- Gehrig, J., Michalareas, G., Forster, M.-T., Lei, J., Hok, P., Laufs, H., Senft, C., Seifert, V., Schoffelen, J.-M., Hanslmayr, S., & Kell, C. A. (2019). Low-frequency oscillations code speech during verbal working memory. *The Journal of Neuroscience*, *39*(33), 6498–6512. <https://doi.org/10.1523/JNEUROSCI.0018-19.2019>
- Gitelman, D. R., Penny, W. D., Ashburner, J., & Friston, K. J. (2003). Modeling regional and psychophysiological interactions in fMRI: The importance of hemodynamic deconvolution. *NeuroImage*, *19*(1), 200–207. [https://doi.org/10.1016/S1053-8119\(03\)00058-2](https://doi.org/10.1016/S1053-8119(03)00058-2)
- Griesmayr, B., Berger, B., Stelzig-Schoeler, R., Aichhorn, W., Bergmann, J., & Sauseng, P. (2014). EEG theta phase coupling during executive control of visual working memory investigated in individuals with schizophrenia and in healthy controls. *Cognitive, Affective, & Behavioral Neuroscience*, *14*(4), 1340–1355. <https://doi.org/10.3758/s13415-014-0272-0>
- Griesmayr, B., Gruber, W. R., Klimesch, W., & Sauseng, P. (2010). Human frontal midline theta and its synchronization to gamma during a verbal delayed match to sample task. *Neurobiology of Learning and Memory*, *93*(2), 208–215. <https://doi.org/10.1016/j.nlm.2009.09.013>
- Grimault, S., Robitaille, N., Grova, C., Lina, J.-M., Dubarry, A.-S., & Jolicœur, P. (2009). Oscillatory activity in parietal and dorsolateral prefrontal cortex during retention in visual short-term memory: Additive effects of spatial attention and memory load. *Human Brain Mapping*, *30*(10), 3378–3392. <https://doi.org/10.1002/hbm.20759>
- Harmony, T., Fernández, T., Silva, J., Bernal, J., Díaz-Comas, L., Reyes, A., Marosi, E., Rodríguez, M., & Rodríguez, M. (1996). EEG delta activity: An indicator of attention to internal processing during performance of mental tasks. *International Journal of Psychophysiology*, *24*(1–2), 161–171. [https://doi.org/10.1016/S0167-8760\(96\)00053-0](https://doi.org/10.1016/S0167-8760(96)00053-0)
- Heinrichs-Graham, E., & Wilson, T. W. (2015). Spatiotemporal oscillatory dynamics during the encoding and maintenance phases of a visual working memory task. *Cortex*, *69*, 121–130. <https://doi.org/10.1016/j.cortex.2015.04.022>
- Heinz, A. J., & Johnson, J. S. (2017). Load-dependent increases in delay-period alpha-band power track the gating of task-irrelevant inputs to working memory. *Frontiers in Human Neuroscience*, *11*, 250. <https://doi.org/10.3389/fnhum.2017.00250>
- Herrmann, C. S., Fründ, I., & Lenz, D. (2010). Human gamma-band activity: A review on cognitive and behavioral correlates and network models. *Neuroscience & Biobehavioral Reviews*, *34*(7), 981–992. <https://doi.org/10.1016/j.neubiorev.2009.09.001>
- Herrmann, C. S., Senkowski, D., & Rottger, S. (2004). Phase-locking and amplitude modulations of EEG alpha: Two measures reflect different cognitive processes in a working memory task. *Experimental Psychology*, *51*(4), 311–318. <https://doi.org/10.1027/1618-3169.51.4.311>
- Honkanen, R., Rouhinen, S., Wang, S. H., Palva, J. M., & Palva, S. (2015). Gamma oscillations underlie the maintenance of feature-specific information and the contents of visual working memory. *Cerebral Cortex*, *25*(10), 3788–3801. <https://doi.org/10.1093/cercor/bhu263>
- Howard, M. W., Rizzuto, D. S., Caplan, J. B., Madsen, J. R., Lisman, J., Aschenbrenner-Scheibe, R., & Kahana, M. J. (2003). Gamma oscillations correlate with working memory load in humans. *Cerebral Cortex*, *13*(12), 1369–1374. <https://doi.org/10.1093/cercor/bhg084>
- Hsieh, L. T., Ekstrom, A. D., & Ranganath, C. (2011). Neural oscillations associated with item and temporal order maintenance in working memory. *The Journal of Neuroscience*, *31*(30), 10803–10810. <https://doi.org/10.1523/jneurosci.0828-11.2011>
- Hsieh, L.-T., & Ranganath, C. (2014). Frontal midline theta oscillations during working memory maintenance and episodic encoding and retrieval. *NeuroImage*, *85*, 721–729. <https://doi.org/10.1016/j.neuroimage.2013.08.003>
- Hu, Z., Barkley, C. M., Marino, S. E., Wang, C., Rajan, A., Bo, K. E., Samuel, I. B. H., & Ding, M. (2019). Working memory capacity is negatively associated with memory load modulation of alpha oscillations in retention of verbal working memory. *Journal of Cognitive Neuroscience*, *31*(12), 1933–1945. https://doi.org/10.1162/jocn_a_01461
- Ichihara-Takeda, S., Yazawa, S., Murahara, T., Toyoshima, T., Shinozaki, J., Ishiguro, M., Shiraishi, H., Ikeda, N., Matsuyama, K., Funahashi, S., & Nagamine, T. (2015). Modulation of alpha activity in the parieto-occipital area by distractors during a visuospatial working memory task: A magnetoencephalographic study. *Journal of Cognitive Neuroscience*, *27*(3), 453–463. https://doi.org/10.1162/jocn_a_00718
- Ikkai, A., Blacker, K. J., Lakshmanan, B. M., Ewen, J. B., & Courtney, S. M. (2014). Maintenance of relational information in working memory leads to suppression of the sensory cortex. *Journal of Neurophysiology*, *112*(8), 1903–1915. <https://doi.org/10.1152/jn.00134.2014>
- Itthipuripat, S., Wessel, J. R., & Aron, A. R. (2013). Frontal theta is a signature of successful working memory manipulation. *Experimental Brain Research*, *224*(2), 255–262. <https://doi.org/10.1007/s00221-012-3305-3>
- Jensen, O., Gelfand, J., Kounios, J., & Lisman, J. E. (2002). Oscillations in the alpha band (9–12 Hz) increase with memory load during retention in a short-term memory task. *Cerebral Cortex*, *12*(8), 877–882. <https://doi.org/10.1093/cercor/12.8.877>
- Jensen, O., & Mazaheri, A. (2010). Shaping functional architecture by oscillatory alpha activity: Gating by inhibition. *Frontiers in Human Neuroscience*, *4*, 186. <https://doi.org/10.3389/fnhum.2010.00186>
- Jensen, O., & Tesche, C. D. (2002). Frontal theta activity in humans increases with memory load in a working memory task. *European Journal of Neuroscience*, *15*(8), 1395–1399. <https://doi.org/10.1046/j.1460-9568.2002.01975.x>
- Jerbi, K., Ossandón, T., Hamamé, C. M., Senova, S., Dalal, S. S., Jung, J., & Lachaux, J.-P. (2009). Task-related gamma-band dynamics

- from an intracerebral perspective: Review and implications for surface EEG and MEG. *Human Brain Mapping*, 30(6), 1758–1771. <https://doi.org/10.1002/hbm.20750>
- Johnson, J. S., Sutterer, D. W., Acheson, D. J., Lewis-Peacock, J. A., & Postle, B. R. (2011). Increased alpha-band power during the retention of shapes and shape-location associations in visual short-term memory. *Frontiers in Psychology*, 2, 128. <https://doi.org/10.3389/fpsyg.2011.00128>
- Jokisch, D., & Jensen, O. (2007). Modulation of gamma and alpha activity during a working memory task engaging the dorsal or ventral stream. *Journal of Neuroscience*, 27(12), 3244–3251. <https://doi.org/10.1523/JNEUROSCI.5399-06.2007>
- Kambara, T., Brown, E. C., Jeong, J.-W., Ofen, N., Nakai, Y., & Asano, E. (2017). Spatio-temporal dynamics of working memory maintenance and scanning of verbal information. *Clinical Neurophysiology*, 128(6), 882–891. <https://doi.org/10.1016/j.clinph.2017.03.005>
- Kaplan, R., Bush, D., Bisby, J. A., Horner, A. J., Meyer, S. S., & Burgess, N. (2016). Medial prefrontal-medial temporal theta phase coupling in dynamic spatial imagery. *Journal of Cognitive Neuroscience*, 29(3), 507–519. https://doi.org/10.1162/jocn_a_01064
- Kawasaki, M., & Yamaguchi, Y. (2012). Individual visual working memory capacities and related brain oscillatory activities are modulated by color preferences. *Frontiers in Human Neuroscience*, 6, 318. <https://doi.org/10.3389/fnhum.2012.00318>
- Kawasaki, M., & Yamaguchi, Y. (2013). Frontal theta and beta synchronizations for monetary reward increase visual working memory capacity. *Social Cognitive and Affective Neuroscience*, 8(5), 523–530. <https://doi.org/10.1093/scan/nss027>
- Khader, P. H., Jost, K., Ranganath, C., & Rosler, F. (2010). Theta and alpha oscillations during working-memory maintenance predict successful long-term memory encoding. *Neuroscience Letters*, 468(3), 339–343. <https://doi.org/10.1016/j.neulet.2009.11.028>
- Klimesch, W. (1999). EEG alpha and theta oscillations reflect cognitive and memory performance: A review and analysis. *Brain Research Reviews*, 29(2), 169–195. [https://doi.org/10.1016/S0165-0173\(98\)00056-3](https://doi.org/10.1016/S0165-0173(98)00056-3)
- Klimesch, W., Doppelmayr, M., Schwaiger, J., Auinger, P., & Winkler, T. (1999). “Paradoxical” alpha synchronization in a memory task. *Cognitive Brain Research*, 7(4), 493–501. [https://doi.org/10.1016/S0926-6410\(98\)00056-1](https://doi.org/10.1016/S0926-6410(98)00056-1)
- Klimesch, W., Sauseng, P., & Hanslmayr, S. (2007). EEG alpha oscillations: The inhibition–timing hypothesis. *Brain Research Reviews*, 53(1), 63–88. <https://doi.org/10.1016/j.brainresrev.2006.06.003>
- Klimesch, W., Schimke, H., & Pfurtscheller, G. (1993). Alpha frequency, cognitive load and memory performance. *Brain Topography*, 5(3), 241–251. <https://doi.org/10.1007/BF01128991>
- Kosciessa, J. Q., Grandy, T. H., Garrett, D. D., & Werkle-Bergner, M. (2020). Single-trial characterization of neural rhythms: Potential and challenges. *NeuroImage*, 206, 116331. <https://doi.org/10.1016/j.neuroimage.2019.116331>
- Kotchoubey, B. (2002). Do event-related brain potentials reflect mental (cognitive) operations? *Journal of Psychophysiology*, 16(3), 129–149. <https://doi.org/10.1027//0269-8803.16.3.129>
- Kottlow, M., Schlaepfer, A., Baenninger, A., Michels, L., Brandeis, D., & Koenig, T. (2015). Pre-stimulus BOLD-network activation modulates EEG spectral activity during working memory retention. *Frontiers in Behavioral Neuroscience*, 9, 111. <https://doi.org/10.3389/fnbeh.2015.00111>
- Kulashekhar, S., Pekkola, J., Palva, J. M., & Palva, S. (2016). The role of cortical beta oscillations in time estimation. *Human Brain Mapping*, 37(9), 3262–3281. <https://doi.org/10.1002/hbm.23239>
- Kustermann, T., Rockstroh, B., Miller, G. A., & Popov, T. (2018). Neural network communication facilitates verbal working memory. *Biological Psychology*, 136, 119–126. <https://doi.org/10.1016/j.biopsycho.2018.05.018>
- Kwon, G., Lim, S., Kim, M.-Y., Kwon, H., Lee, Y.-H., Kim, K., Lee, E.-J., & Suh, M. (2015). Individual differences in oscillatory brain activity in response to varying attentional demands during a word recall and oculomotor dual task. *Frontiers in Human Neuroscience*, 9, 381. <https://doi.org/10.3389/fnhum.2015.00381>
- Leijten, F. S. S., Huiskamp, G.-J.-M., Hilgersom, I., & van Huffelen, A. C. (2003). High-resolution source imaging in mesiotemporal lobe epilepsy: A comparison between MEG and simultaneous EEG. *Journal of Clinical Neurophysiology*, 20(4), 227. <https://doi.org/10.1097/00004691-200307000-00001>
- Leszczyński, M., Fell, J., & Axmacher, N. (2015). Rhythmic working memory activation in the human hippocampus. *Cell Reports*, 13(6), 1272–1282. <https://doi.org/10.1016/j.celrep.2015.09.081>
- Lindquist, M. A., Meng Loh, J., Atlas, L. Y., & Wager, T. D. (2009). Modeling the hemodynamic response function in fMRI: Efficiency, bias and mis-modeling. *NeuroImage*, 45(1), S187–S198. <https://doi.org/10.1016/j.neuroimage.2008.10.065>
- Luck, S. J., & Vogel, E. K. (1997). The capacity of visual working memory for features and conjunctions. *Nature*, 390(6657), 279–281. <https://doi.org/10.1038/36846>
- Luria, R., Balaban, H., Awh, E., & Vogel, E. K. (2016). The contralateral delay activity as a neural measure of visual working memory. *Neuroscience and Biobehavioral Reviews*, 62, 100–108. <https://doi.org/10.1016/j.neubiorev.2016.01.003>
- Mainy, N., Kahane, P., Minotti, L., Hoffmann, D., Bertrand, O., & Lachaux, J. P. (2007). Neural correlates of consolidation in working memory. *Human Brain Mapping*, 28(3), 183–193. <https://doi.org/10.1002/hbm.20264>
- Malmivuo, J. (2012). Comparison of the properties of EEG and MEG in detecting the electric activity of the brain. *Brain Topography*, 25(1), 1–19. <https://doi.org/10.1007/s10548-011-0202-1>
- Manza, P., Hau, C. L., & Leung, H. C. (2014). Alpha power gates relevant information during working memory updating. *The Journal of Neuroscience*, 34(17), 5998–6002. <https://doi.org/10.1523/jneurosci.4641-13.2014>
- Mapelli, I., & Ozkurt, T. E. (2019). Brain oscillatory correlates of visual short-term memory errors. *Frontiers in Human Neuroscience*, 13, 33. <https://doi.org/10.3389/fnhum.2019.00033>
- Maurer, U., Brem, S., Liechti, M., Maurizio, S., Michels, L., & Brandeis, D. (2015). Frontal midline theta reflects individual task performance in a working memory task. *Brain Topography*, 28(1), 127–134. <https://doi.org/10.1007/s10548-014-0361-y>
- Medendorp, W. P., Kramer, G. F. I., Jensen, O., Oostenveld, R., Schoffelen, J.-M., & Fries, P. (2007). Oscillatory activity in human parietal and occipital cortex shows hemispheric lateralization and memory effects in a delayed double-step saccade task. *Cerebral Cortex*, 17(10), 2364–2374. <https://doi.org/10.1093/cercor/bhl145>
- Meltzer, J. A., Negishi, M., Mayes, L. C., & Constable, R. T. (2007). Individual differences in EEG theta and alpha dynamics during working memory correlate with fMRI responses across subjects. *Clinical Neurophysiology*, 118(11), 2419–2436. <https://doi.org/10.1016/j.clinph.2007.07.023>
- Meltzer, J. A., Zaveri, H. P., Goncharova, I. I., Distasio, M. M., Papademetris, X., Spencer, S. S., Spencer, D. D., & Constable, R. T. (2008). Effects of working memory load on oscillatory power

- in human intracranial EEG. *Cerebral Cortex*, 18(8), 1843–1855. <https://doi.org/10.1093/cercor/bhm213>
- Michels, L., Bucher, K., Lüchinger, R., Klaver, P., Martin, E., Jeanmonod, D., & Brandeis, D. (2010). Simultaneous EEG-fMRI during a working memory task: Modulations in Low and high frequency bands. *PLoS One*, 5(4), e10298. <https://doi.org/10.1371/journal.pone.0010298>
- Michels, L., Moazami-Goudarzi, M., Jeanmonod, D., & Sarnthein, J. (2008). EEG alpha distinguishes between cuneal and precuneal activation in working memory. *NeuroImage*, 40(3), 1296–1310. <https://doi.org/10.1016/j.neuroimage.2007.12.048>
- Moran, R. J., Campo, P., Maestú, F., Reilly, R. B., Dolan, R. J., & Strange, B. A. (2010). Peak frequency in the theta and alpha bands correlates with human working memory capacity. *Frontiers in Human Neuroscience*, 4. <https://doi.org/10.3389/fnhum.2010.00200>
- Morgan, H. M., Muthukumaraswamy, S. D., Hibbs, C. S., Shapiro, K. L., Bracewell, R. M., Singh, K. D., & Linden, D. E. J. (2011). Feature integration in visual working memory: Parietal gamma activity is related to cognitive coordination. *Journal of Neurophysiology*, 106(6), 3185–3194. <https://doi.org/10.1152/jn.00246.2011>
- Murphy, O. W., Hoy, K. E., Wong, D., Bailey, N. W., Fitzgerald, P. B., & Segrave, R. A. (2019). Individuals with depression display abnormal modulation of neural oscillatory activity during working memory encoding and maintenance. *Biological Psychology*, 148, 107766. <https://doi.org/10.1016/j.biopsycho.2019.107766>
- Muthukumaraswamy, S. (2013). High-frequency brain activity and muscle artifacts in MEG/EEG: A review and recommendations. *Frontiers in Human Neuroscience*, 7. <https://doi.org/10.3389/fnhum.2013.00138>
- Myers, N. E., Walther, L., Wallis, G., Stokes, M. G., & Nobre, A. C. (2015). Temporal dynamics of attention during encoding versus maintenance of working memory: Complementary views from event-related potentials and alpha-band oscillations. *Journal of Cognitive Neuroscience*, 27(3), 492–508. https://doi.org/10.1162/jocn_a_00727
- Nenert, R., Viswanathan, S., Dubuc, D. M., & Visscher, K. M. (2012). Modulations of ongoing alpha oscillations predict successful short-term visual memory encoding. *Frontiers in Human Neuroscience*, 6, 127. <https://doi.org/10.3389/fnhum.2012.00127>
- Oberauer, K., Lewandowsky, S., Awh, E., Brown, G. D. A., Conway, A., Cowan, N., Donkin, C., Farrell, S., Hitch, G. J., Hurlstone, M. J., Ma, W. J., Morey, C. C., Nee, D. E., Schweppe, J., Vergauwe, E., & Ward, G. (2018). Benchmarks for models of short-term and working memory. *Psychological Bulletin*, 144(9), 885. <https://doi.org/10.1037/bul0000153>
- Ocklenburg, S., Friedrich, P., Schmitz, J., Schlüter, C., Genc, E., Güntürkün, O., Peterburs, J., & Grimshaw, G. (2019). Beyond frontal alpha: Investigating hemispheric asymmetries over the EEG frequency spectrum as a function of sex and handedness. *Laterality: Asymmetries of Body, Brain and Cognition*, 24(5), 505–524. <https://doi.org/10.1080/1357650x.2018.1543314>
- Okada, Y. C., & Salenius, S. (1998). Roles of attention, memory and motor preparation in modulating human brain activity in a spatial working memory task. *Cerebral Cortex*, 8(1), 80–96. <https://doi.org/10.1093/cercor/8.1.80>
- Okuhata, S., Kusanagi, T., & Kobayashi, T. (2013). Parietal EEG alpha suppression time of memory retrieval reflects memory load while the alpha power of memory maintenance is a composite of the visual process according to simultaneous and successive Sternberg memory tasks. *Neuroscience Letters*, 555, 79–84. <https://doi.org/10.1016/j.neulet.2013.09.010>
- Olsen, R. K., Rondina, R., Riggs, L., Meltzer, J. A., & Ryan, J. D. (2013). Hippocampal and neocortical oscillatory contributions to visuospatial binding and comparison. *Journal of Experimental Psychology-General*, 142(4), 1335–1345. <https://doi.org/10.1037/a0034043>
- Onton, J., Delorme, A., & Makeig, S. (2005). Frontal midline EEG dynamics during working memory. *NeuroImage*, 27(2), 341–356. <https://doi.org/10.1016/j.neuroimage.2005.04.014>
- Pae, C.-U. (2015). Why systematic review rather than narrative review? *Psychiatry Investigation*, 12(3), 417–419. <https://doi.org/10.4306/pi.2015.12.3.417>
- Pahor, A., & Jausovec, N. (2017). Multifaceted pattern of neural efficiency in working memory capacity. *Intelligence*, 65, 23–34. <https://doi.org/10.1016/j.intell.2017.10.001>
- Palomaki, J., Kivikangas, M., Alafuzoff, A., Hakala, T., & Krause, C. M. (2012). Brain oscillatory 4–35 Hz EEG responses during an n-back task with complex visual stimuli. *Neuroscience Letters*, 516(1), 141–145. <https://doi.org/10.1016/j.neulet.2012.03.076>
- Park, H., Kang, E., Kang, H., Kim, J. S., Jensen, O., Chung, C. K., & Lee, D. S. (2011). Cross-frequency power correlations reveal the right superior temporal gyrus as a hub region during working memory maintenance. *Brain Connectivity*, 1(6), 460–472. <https://doi.org/10.1089/brain.2011.0046>
- Park, J. Y., Min, B. K., Jung, Y. C., Pak, H., Jeong, Y. H., & Kim, E. (2013). Illumination influences working memory: An EEG study. *Neuroscience*, 247, 386–394. <https://doi.org/10.1016/j.neuroscience.2013.05.016>
- Parvizi, J., & Kastner, S. (2018). Promises and limitations of human intracranial electroencephalography. *Nature Neuroscience*, 21(4), 474–483. <https://doi.org/10.1038/s41593-018-0108-2>
- Pavlov, Y. G., & Kotchoubey, B. (2017). EEG correlates of working memory performance in females. *BMC Neuroscience*, 18(1), 26. <https://doi.org/10.1186/s12868-017-0344-5>
- Payne, L., & Sekuler, R. (2014). The importance of ignoring: Alpha oscillations protect selectivity. *Current Directions in Psychological Science*, 23(3), 171–177. <https://doi.org/10.1177/0963721414529145>
- Pesonen, M., Hämäläinen, H., & Krause, C. M. (2007). Brain oscillatory 4–30 Hz responses during a visual n-back memory task with varying memory load. *Brain Research*, 1138, 171–177. <https://doi.org/10.1016/j.brainres.2006.12.076>
- Poch, C., Campo, P., & Barnes, G. R. (2014). Modulation of alpha and gamma oscillations related to retrospectively orienting attention within working memory. *European Journal of Neuroscience*, 40(2), 2399–2405. <https://doi.org/10.1111/ejn.12589>
- Poch, C., Campo, P., Parmentier, F. B. R., Ruiz-Vargas, J. M., Elsley, J. V., Castellanos, N. P., Maestú, F., & del Pozo, F. (2010). Explicit processing of verbal and spatial features during letter-location binding modulates oscillatory activity of a fronto-parietal network. *Neuropsychologia*, 48(13), 3846–3854. <https://doi.org/10.1016/j.neuropsychologia.2010.09.015>
- Poch, C., Valdivia, M., Capilla, A., Hinojosa, J. A., & Campo, P. (2018). Suppression of no-longer relevant information in Working Memory: An alpha-power related mechanism? *Biological Psychology*, 135, 112–116. <https://doi.org/10.1016/j.biopsycho.2018.03.009>
- Postle, B. R. (2006). Working memory as an emergent property of the mind and brain. *Neuroscience*, 139(1), 23–38. <https://doi.org/10.1016/j.neuroscience.2005.06.005>

- Proskovec, A. L., Heinrichs-Graham, E., & Wilson, T. W. (2019). Load modulates the alpha and beta oscillatory dynamics serving verbal working memory. *NeuroImage*, *184*, 256–265. <https://doi.org/10.1016/j.neuroimage.2018.09.022>
- Proskovec, A. L., Wiesman, A. I., Heinrichs-Graham, E., & Wilson, T. W. (2018). Beta oscillatory dynamics in the prefrontal and superior temporal cortices predict spatial working memory performance. *Scientific Reports*, *8*(1), 8488. <https://doi.org/10.1038/s41598-018-26863-x>
- Proskovec, A. L., Wiesman, A. I., Heinrichs-Graham, E., & Wilson, T. W. (2019). Load effects on spatial working memory performance are linked to distributed alpha and beta oscillations. *Human Brain Mapping*, *40*, 3682–3689. <https://doi.org/10.1002/hbm.24625>
- Raghavachari, S., Kahana, M. J., Rizzuto, D. S., Caplan, J. B., Kirschen, M. P., Bourgeois, B., Madsen, J. R., & Lisman, J. E. (2001). Gating of human theta oscillations by a working memory task. *The Journal of Neuroscience*, *21*(9), 3175–3183. <https://doi.org/10.1523/JNEUROSCI.21-09-03175.2001>
- Rawle, C. J., Miall, R. C., & Praamstra, P. (2012). Frontoparietal theta activity supports behavioral decisions in movement-target selection. *Frontiers in Human Neuroscience*, *6*, 138. <https://doi.org/10.3389/fnhum.2012.00138>
- Roberts, B. M., Hsieh, L. T., & Ranganath, C. (2013). Oscillatory activity during maintenance of spatial and temporal information in working memory. *Neuropsychologia*, *51*(2), 349–357. <https://doi.org/10.1016/j.neuropsychologia.2012.10.009>
- Román-López, T. V., Caballero-Sánchez, U., Cisneros-Luna, S., Franco-Rodríguez, J. A., Méndez-Díaz, M., Prospéro-García, O., & Ruiz-Contreras, A. E. (2019). Brain electrical activity from encoding to retrieval while maintaining and manipulating information in working memory. *Memory*, *27*, 1–16. <https://doi.org/10.1080/09658211.2019.1620287>
- Rominger, C., Fink, A., Weiss, E. M., Schuster, G., Perchtold, C. M., & Papousek, I. (2019). The propensity to perceive meaningful coincidences is associated with increased posterior alpha power during retention of information in a modified Sternberg paradigm. *Consciousness and Cognition*, *76*, 102832. <https://doi.org/10.1016/j.concog.2019.102832>
- Roux, F., & Uhlhaas, P. J. (2014). Working memory and neural oscillations: Alpha-gamma versus theta-gamma codes for distinct WM information? *Trends in Cognitive Sciences*, *18*(1), 16–25. <https://doi.org/10.1016/j.tics.2013.10.010>
- Roux, F., Wibral, M., Mohr, H. M., Singer, W., & Uhlhaas, P. J. (2012). Gamma-band activity in human prefrontal cortex codes for the number of relevant items maintained in working memory. *Journal of Neuroscience*, *32*(36), 12411–12420. <https://doi.org/10.1523/jneurosci.0421-12.2012>
- Sauseng, P., Klimesch, W., Doppelmayr, M., Pecherstorfer, T., Freunberger, R., & Hanslmayr, S. (2005). EEG alpha synchronization and functional coupling during top-down processing in a working memory task. *Human Brain Mapping*, *26*(2), 148–155. <https://doi.org/10.1002/hbm.20150>
- Sauseng, P., Klimesch, W., Heise, K. F., Gruber, W. R., Holz, E., Karim, A. A., Glennon, M., Gerloff, C., Birbaumer, N., & Hummel, F. C. (2009). Brain oscillatory substrates of visual short-term memory capacity. *Current Biology*, *19*(21), 1846–1852. <https://doi.org/10.1016/j.cub.2009.08.062>
- Schack, B., & Klimesch, W. (2002). Frequency characteristics of evoked and oscillatory electroencephalic activity in a human memory scanning task. *Neuroscience Letters*, *331*(2), 107–110. [https://doi.org/10.1016/S0304-3940\(02\)00846-7](https://doi.org/10.1016/S0304-3940(02)00846-7)
- Scheeringa, R., Petersson, K. M., Oostenveld, R., Norris, D. G., Hagoort, P., & Bastiaansen, M. C. M. (2009). Trial-by-trial coupling between EEG and BOLD identifies networks related to alpha and theta EEG power increases during working memory maintenance. *NeuroImage*, *44*(3), 1224–1238. <https://doi.org/10.1016/j.neuroimage.2008.08.041>
- Schneider, D., Goddertz, A., Haase, H., Hickey, C., & Wascher, E. (2019). Hemispheric asymmetries in EEG alpha oscillations indicate active inhibition during attentional orienting within working memory. *Behavioural Brain Research*, *359*, 38–46. <https://doi.org/10.1016/j.bbr.2018.10.020>
- Schroeder, S. C. Y., Ball, F., & Busch, N. A. (2018). The role of alpha oscillations in distractor inhibition during memory retention. *European Journal of Neuroscience*, *48*(7), 2516–2526. <https://doi.org/10.1111/ejn.13852>
- Seemuller, A., Muller, E. M., & Rosler, F. (2012). EEG-power and -coherence changes in a unimodal and a crossmodal working memory task with visual and kinesthetic stimuli. *International Journal of Psychophysiology*, *83*(1), 87–95. <https://doi.org/10.1016/j.ijpsycho.2011.10.009>
- Segrave, R. A., Thomson, R. H., Cooper, N. R., Croft, R. J., Sheppard, D. M., & Fitzgerald, P. B. (2010). Upper alpha activity during working memory processing reflects abnormal inhibition in major depression. *Journal of Affective Disorders*, *127*(1–3), 191–198. <https://doi.org/10.1016/j.jad.2010.05.022>
- Sharon, D., Hämäläinen, M. S., Tootell, R. B. H., Halgren, E., & Belliveau, J. W. (2007). The advantage of combining MEG and EEG: Comparison to fMRI in focally stimulated visual cortex. *NeuroImage*, *36*(4), 1225–1235. <https://doi.org/10.1016/j.neuroimage.2007.03.066>
- Smyrnis, N., Protopapa, F., Tsoukas, E., Balogh, A., Siettos, C. I., & Evdokimidis, I. (2014). Amplitude spectrum EEG signal evidence for the dissociation of motor and perceptual spatial working memory in the human brain. *Experimental Brain Research*, *232*(2), 659–673. <https://doi.org/10.1007/s00221-013-3774-z>
- Šoškić, A., Jovanović, V., Styles, S. J., Kappenman, E. S., & Kovic, V. (2020). How to do better N400 studies: Reproducibility, consistency and adherence to research standards in the existing literature. *PsyArXiv* <https://doi.org/10.31234/osf.io/jp6wy>
- Spitzer, B., & Blankenburg, F. (2012). Supramodal parametric working memory processing in humans. *The Journal of Neuroscience*, *32*(10), 3287–3295. <https://doi.org/10.1523/JNEUROSCI.5280-11.2012>
- Spitzer, B., Fleck, S., & Blankenburg, F. (2014). Parametric alpha- and beta-band signatures of supramodal numerosity information in human working memory. *The Journal of Neuroscience*, *34*(12), 4293–4302. <https://doi.org/10.1523/jneurosci.4580-13.2014>
- Stefan, H., Schüler, P., Abraham-Fuchs, K., Schneider, S., Gebhardt, M., Neubauer, U., Hummel, C., Huk, W. J., & Thierauf, P. (1994). Magnetic source localization and morphological changes in temporal lobe epilepsy: Comparison of MEG/EEG, ECoG and volumetric MRI in presurgical evaluation of operated patients. *Acta Neurologica Scandinavica*, *89*(S152), 83–88. <https://doi.org/10.1111/j.1600-0404.1994.tb05192.x>
- Steinbrink, J., Villringer, A., Kempf, F., Haux, D., Boden, S., & Obrig, H. (2006). Illuminating the BOLD signal: Combined fMRI–fNIRS studies. *Magnetic Resonance Imaging*, *24*(4), 495–505. <https://doi.org/10.1016/j.mri.2005.12.034>

- Stephane, M., Ince, N. F., Kuskowski, M., Leuthold, A., Tewfik, A. H., Nelson, K., McClannahan, K., Fletcher, C. R., & Tadipatri, V. A. (2010). Neural oscillations associated with the primacy and recency effects of verbal working memory. *Neuroscience Letters*, *473*(3), 172–177. <https://doi.org/10.1016/j.neulet.2010.02.025>
- Sternberg, S. (1966). High-speed scanning in human memory. *Science*, *153*(3736), 652–654. <https://doi.org/10.1126/science.153.3736.652>
- Stewart, J. L., Coan, J. A., Towers, D. N., & Allen, J. J. B. (2014). Resting and task-elicited prefrontal EEG alpha asymmetry in depression: Support for the capability model: Resting and task-elicited asymmetry. *Psychophysiology*, *51*(5), 446–455. <https://doi.org/10.1111/psyp.12191>
- Takase, R., Boasen, J., & Yokosawa, K. (2019). Different roles for theta- and alpha-band brain rhythms during sequential memory. *2019 41st Annual International Conference of the IEEE Engineering in Medicine and Biology Society (EMBC)*, 1713–1716.
- Tallon-Baudry, C., Kreiter, A., & Bertrand, O. (1999). Sustained and transient oscillatory responses in the gamma and beta bands in a visual short-term memory task in humans. *Visual Neuroscience*, *16*(3), 449–459.
- Todd, J. J., & Marois, R. (2004). Capacity limit of visual short-term memory in human posterior parietal cortex. *Nature*, *428*(6984), 751–754. <https://doi.org/10.1038/nature02466>
- Trubutschek, D., Marti, S., Ojeda, A., King, J. R., Mi, Y., Tsodyks, M., & Dehaene, S. (2017). A theory of working memory without consciousness or sustained activity. *Elife*, *6*. <https://doi.org/10.7554/eLife.23871>
- Trubutschek, D., Marti, S., Ueberschar, H., & Dehaene, S. (2019). Probing the limits of activity-silent non-conscious working memory. *Proceedings of the National Academy of Sciences of the United States of America*, *116*(28), 14358–14367. <https://doi.org/10.1073/pnas.1820730116>
- Tuladhar, A. M., ter Huurne, N., Schoffelen, J.-M., Maris, E., Oostenveld, R., & Jensen, O. (2007). Parieto-occipital sources account for the increase in alpha activity with working memory load. *Human Brain Mapping*, *28*(8), 785–792. <https://doi.org/10.1002/hbm.20306>
- van der Vinne, N., Vollebregt, M. A., van Putten, M. J. A. M., & Arns, M. (2017). Frontal alpha asymmetry as a diagnostic marker in depression: Fact or fiction? *A meta-analysis. Neuroimage: Clinical*, *16*, 79–87. <https://doi.org/10.1016/j.nicl.2017.07.006>
- van Dijk, H., van der Werf, J., Mazaheri, A., Medendorp, W. P., & Jensen, O. (2010). Modulations in oscillatory activity with amplitude asymmetry can produce cognitively relevant event-related responses. *Proceedings of the National Academy of Sciences of the United States of America*, *107*(2), 900–905. <https://doi.org/10.1073/pnas.0908821107>
- van Ede, F. (2018). Mnemonic and attentional roles for states of attenuated alpha oscillations in perceptual working memory: A review. *The European Journal of Neuroscience*, *48*(7), 2509–2515. <https://doi.org/10.1111/ejn.13759>
- van Ede, F., Jensen, O., & Maris, E. (2017). Supramodal theta, gamma, and sustained fields predict modality-specific modulations of alpha and beta oscillations during visual and tactile working memory. *Journal of Cognitive Neuroscience*, *29*(8), 1455–1472. https://doi.org/10.1162/jocn_a_01129
- Vandenbroucke, A. R. E., Sligte, I. G., de Vries, J. G., Cohen, M. X., & Lamme, V. A. F. (2015). Neural correlates of visual short-term memory dissociate between fragile and working memory representations. *Journal of Cognitive Neuroscience*, *27*(12), 2477–2490. https://doi.org/10.1162/jocn_a_00870
- Vissers, M. E., van Driel, J., & Slagter, H. A. (2016). Proactive, but not reactive, distractor filtering relies on local modulation of alpha oscillatory activity. *Journal of Cognitive Neuroscience*, *28*(12), 1964–1979. https://doi.org/10.1162/jocn_a_01017
- Vogel, E. K., & Machizawa, M. G. (2004). Neural activity predicts individual differences in visual working memory capacity. *Nature*, *428*(6984), 748–751. <https://doi.org/10.1038/nature02447>
- Wianda, E., & Ross, B. (2019). The roles of alpha oscillation in working memory retention. *Brain and Behavior*, *9*(4), e01263. <https://doi.org/10.1002/brb3.1263>
- Wilson, G. F., Swain, C. R., & Ullsperger, P. (1999). EEG power changes during a multiple level memory retention task. *International Journal of Psychophysiology*, *32*(2), 107–118. [https://doi.org/10.1016/S0167-8760\(99\)00005-7](https://doi.org/10.1016/S0167-8760(99)00005-7)
- Wolff, M. J., Jochim, J., Akyürek, E. G., & Stokes, M. G. (2017). Dynamic hidden states underlying working-memory-guided behavior. *Nature Neuroscience*, *20*(6), 864. <https://doi.org/10.1038/nn.4546>
- Worden, M. S., Foxe, J. J., Wang, N., & Simpson, G. V. (2000). Anticipatory biasing of visuospatial attention indexed by retinotopically specific-band electroencephalography increases over occipital cortex. *Journal of Neuroscience*, *20*(RC63), 1–6.
- Xie, Y., Feng, Z., Xu, Y., Bian, C., & Li, M. (2016). The different oscillation patterns of alpha band in the early and later stages of working memory maintenance. *Neuroscience Letters*, *633*, 220–226. <https://doi.org/10.1016/j.neulet.2016.09.047>
- Yin, J., Gao, Z., Jin, X., Ding, X., Liang, J., & Shen, M. (2012). The neural mechanisms of percept-memory comparison in visual working memory. *Biological Psychology*, *90*(1), 71–79. <https://doi.org/10.1016/j.biopsycho.2012.02.023>
- Zakrzewska, M. Z., & Brzezicka, A. (2014). Working memory capacity as a moderator of load-related frontal midline theta variability in Sternberg task. *Frontiers in Human Neuroscience*, *8*, 399. <https://doi.org/10.3389/fnhum.2014.00399>
- Zhang, D., Zhao, H., Bai, W., & Tian, X. (2016). Functional connectivity among multi-channel EEGs when working memory load reaches the capacity. *Brain Research*, *1631*, 101–112. <https://doi.org/10.1016/j.brainres.2015.11.036>
- Zuure, M. B., Hinkley, L. B. N., Tiesinga, P. H. E., Nagarajan, S. S., & Cohen, M. X. (2020). Multiple midfrontal thetas revealed by source separation of simultaneous MEG and EEG. *The Journal of Neuroscience*, *40*(40), 7702–7713. <https://doi.org/10.1523/JNEUROSCI.0321-20.2020>

How to cite this article: Pavlov, Y. G., Kotchoubey, B. (2022). Oscillatory brain activity and maintenance of verbal and visual working memory: A systematic review. *Psychophysiology*, *59*, e13735. <https://doi.org/10.1111/psyp.13735>