Title

Methodological Considerations for Studying Neural Oscillations

running title: analysis of neural oscillations

Authors

Thomas Donoghue¹, Natalie Schaworonkow¹, & Bradley Voytek^{1, 2, 3, 4}

Affiliations

¹Department of Cognitive Science, University of California, San Diego

²Neurosciences Graduate Program, University of California, San Diego

³Halicioğlu Data Science Institute, University of California, San Diego

⁴Kavli Institute for Brain and Mind, University of California, San Diego

Corresponding Author

Thomas Donoghue: tdonoghue.research@gmail.com

Contributions

All authors contributed to designing the study. T.D and N.S did the analyses and created the figures. All authors contributed to writing and editing the paper.

ORCIDs

Donoghue 0000-0001-5911-0472

Schaworonkow 0000-0002-2103-1178

Voytek 0000-0003-1640-2525

Conflicts of Interest

The authors declare no competing interests.

Acknowledgements

We would like to thank Ryan Hammonds for his contributions to the methods and software tools used for this report. This work was supported by research funding from the National Institute of General Medical Sciences grant R01GM134363-02, National Science Foundation grant BCS-1736028, and a UC San Diego Halicioğlu Data Science Institute Fellowship.

Materials Descriptions & Availability Statements

Project Repository

This project is also made openly available through an online project repository in which the code and data are made available, with step-by-step guides through the analyses.

Project Repository: <https://github.com/voytekresearch/OscillationMethods>

Datasets

This project uses simulated data. Code to recreate the exact simulations used is included in the project repository.

Software

Code used and written for this project was written in the Python programming language ($\geq v3.7$). All the code used within this project is deposited in the project repository and is made openly available and licensed for reuse.

This project uses the following Python packages for simulating and analyzing neural data:

neurodsp	https://neurodsp-tools.github.io/
fooof	https://fooof-tools.github.io/
bycycle	https://bycycle-tools.github.io/

In addition, third party Python toolboxes including mne, numpy, scipy, matplotlib, and seaborn were used in this project.

methods for analyzing neural oscillations

reviewing data properties & method assumptions

65 **Graphical Abstract.** Neural oscillations are ubiquitous features of neural field data, with great potential for
 66 informing our understanding of neural function and how it relates to cognition. However, there is a great
 67 degree of variability in methods for investigating them, and findings that are reported. In this piece, we
 68 explore methodological considerations for analyzing neural oscillations, that may underlie some potential
 69 misinterpretations, and propose best practice guidelines for addressing them.

Abstract

Neural oscillations are ubiquitous across recording methodologies and species, broadly associated with cognitive tasks, and amenable to computational modeling that investigates neural circuit generating mechanisms and neural population dynamics. Because of this, neural oscillations offer an exciting potential opportunity for linking theory, physiology, and mechanisms of cognition. However, despite their prevalence, there are many concerns—new and old—about how our analysis assumptions are violated by known properties of field potential data. For investigations of neural oscillations to be properly interpreted, and ultimately developed into mechanistic theories, it is necessary to carefully consider the underlying assumptions of the methods we employ. Here, we discuss seven methodological considerations for analyzing neural oscillations. The considerations are to 1) verify the presence of oscillations, as they may be absent; 2) validate oscillation band definitions, to address variable peak frequencies; 3) account for concurrent non-oscillatory aperiodic activity, which might otherwise confound measures; 4) measure and account for 4) temporal variability and 5) waveform shape of neural oscillations, which are often bursty and/or nonsinusoidal, potentially leading to spurious results; 6) separate spatially overlapping rhythms, which may interfere with each other; and 7) consider the required signal-to-noise ratio for obtaining reliable estimates. For each topic, we provide relevant examples, demonstrate potential errors of interpretation, and offer suggestions to address these issues. We primarily focus on univariate measures, such as power and phase estimates, though we discuss how these issues can propagate to multivariate measures. These considerations and recommendations offer a helpful guide for measuring and interpreting neural oscillations.

Keywords

neural field data, digital signal processing, electrophysiology, time series analysis, spectral analysis

Abbreviations

EEG: electroencephalography; MEG: magnetoencephalography; SNR: signal-to-noise ratio

Introduction

Recordings of electrical or magnetic fields in the brain, collectively referred to as neural field recordings, are commonly used for investigating links between physiology and behavior, cognition, and disease. A striking feature of such recordings is the prominent rhythmic activity, termed neural oscillations (Buzsáki & Draguhn, 2004), that stands out in the otherwise seemingly chaotic activity of the brain. Neural oscillations have been a feature of interest since the early days of electrical brain recordings (Brazier, 1958), and are widely observed, being ubiquitously present across species (Buzsáki et al., 2013). Physiologically, field potential recordings largely reflect the aggregate postsynaptic and transmembrane currents of thousands to millions of neurons (Buzsáki et al., 2012), with neural oscillations thought to relate to population synchrony (Wang, 2010). As such, neural oscillations potentially offer insight into the coordination of neural activity at the population level. Theories of the functions of oscillations argue that they facilitate dynamic temporal and spatial organization of neural activity (Fries, 2005; VanRullen, 2016; Varela et al., 2001; Voytek & Knight, 2015). Disruptions of oscillations have also been widely linked to neurological and psychiatric disease, and have been explored as potential biomarkers of disease status, drug efficacy, and other clinical indicators (Başar, 2013; Buzsáki & Watson, 2012; Newson & Thiagarajan, 2019).

Reflecting this broad interest, thousands of investigations conducted across many decades have reported associations between oscillations and just about every aspect of behavior and cognition that can be operationalized (Başar et al., 2001; Lopes da Silva, 2013; Mazaheri et al., 2018). As neural oscillations appear at many different temporal scales (Buzsáki et al., 2013), investigations often focus on predefined canonical frequency band ranges that are thought to capture distinct oscillations. For example, sleep researchers often study delta (1-4 Hz), memory researchers theta (4-8 Hz), visual researchers alpha (8-12 Hz), and cognitive and motor researchers beta (13-30 Hz) frequency bands. In doing so, research in neural oscillations spans across different recording modalities (Buzsáki et al., 2012)—including both non-invasive and invasive methods—and across different brain regions (Frauscher et al., 2018; Mahjoory et al., 2020).

While oscillations provide an exciting possibility to link cognition and disease to theory and physiology, there are often inconsistent reports regarding which oscillations are modulated by which conditions and how. In part, this likely reflects the variety of approaches taken, with limited consistency in terms of experimental design, analysis methods, parameter choices, and theoretical frameworks used across studies. Open challenges include developing more consistent

terminology and interpretations (Cohen & Gulbinaite, 2014), and the need for explicitly considering replicability in electrophysiological investigations (Cohen, 2017a). Accordingly, best practice guidelines for research (Gross et al., 2013; Keil et al., 2014; Pernet et al., 2020; Pesaran et al., 2018) and clinical investigations (Babiloni et al., 2020) have been proposed to improve standards of reporting, and therefore reproducibility, for research using neural field recordings.

As an extension of these general guidelines, here we examine common interpretational considerations in analyzing neural field recordings. Given the advances in both methods development and our understanding of the empirical properties of the data under study, it is critically important to ensure that common analysis methods are appropriately applied, as this is a core requisite for accurate interpretation. There is a large toolkit of analysis methods for studying neural oscillations, across both the spectral and temporal domains, borrowed and adapted from the field of digital signal processing. These methods are described and compared in other work focused on methodological properties of particular estimation techniques (Bruns, 2004; Gross, 2014; van Vugt et al., 2007; Wacker & Witte, 2013).

Here, we focus more explicitly on properties of neural oscillations, and how these properties relate to commonly applied methods, rather than focus on the methods themselves. We address how common analysis approaches can give rise to results that are easy for researchers to misinterpret, due to the misalignment between methodological or experimental assumptions, and properties of the data. As such, these considerations are not restricted to individual estimators (such as using particular filters, or a particular estimate of power), as they reflect more general properties of signal processing methods and neural data. Importantly, these are not failures of the algorithms per se, which do, mathematically, exactly what they should; the potential pitfalls lay in how we interpret their outputs. If and when there is a misalignment between methodological assumptions and data properties, computed measures can lack validity which can lead to inconsistent results. This in turn impedes us from properly grounding oscillation research in physiology and theory.

To address these issues, we examine common interpretational considerations in studying neural oscillations, in order to identify and address possible methodological concerns that may lead to interpretation errors. We consider recurring themes based on our developing understanding of neural field data, and how this understanding relates to the application of analysis methods. For example, a common assumption is that neural field data can be quantified as a series of oscillatory signals, often assumed to be stationary. However, in empirical neurophysiological data, oscillations show large variability in their presence and extent across time, as well as across participants and cortical regions (Donoghue et al., 2020b; Frauscher et

al., 2018; Groppe et al., 2013). Even when oscillations are present, they are highly variable (Jones, 2016; Neymotin, Barczak, et al., 2020), waxing and waning in short bursts and including longer, more tonic rhythms, with rapidly changing amplitude, frequency, and phase dynamics that are not easily captured by common analyses and predefined canonical frequency ranges. This potentially meaningful variation of cycle features across time can be blurred by narrowband filtering (de Cheveigné & Nelken, 2019) and lead to misinterpretations of which features of the oscillation have truly changed (Cole & Voytek, 2019). All of these properties, and more, need to be explicitly considered in order to accurately and reliably measure oscillatory neural activity.

We organize methodological considerations for analyzing neural oscillations into seven areas, each with example demonstrations (see **Box 1**). The primary focus is on univariate measures of oscillatory power, frequency, and phase, including potential pitfalls and considerations for ensuring accurate measurement and interpretation of these aspects, as well as discussions of how these issues can propagate to multivariate analyses, such as cross-frequency coupling. These demonstrations make use of simulated data, which is created to match known properties of neural field recordings whereby key features of the simulated neural field activity were chosen and manipulated to reflect experimentally observed variations in empirical data. We analyze the simulated data using common spectral and time-domain analysis methods in order to evaluate their performance in relation to the interplay of data properties and method assumptions. Each consideration is then contextualized within the broader literature, and specific practical recommendations are made to help guide the analysis of neural oscillations. The simulated data and analysis methods were created and used from the NeuroDSP module (Cole et al., 2019), with all associated code for recreating and further exploring the illustrations openly available in the project repository (<https://github.com/voytekresearch/oscillationmethods>).

191 **Box 1:** Overview of methodological considerations for measuring neural oscillations

Topic	Data Properties	Methodological Issues	Recommendation
#1 Oscillation Presence	neural oscillations are variably present, and may not be present in the recording	if there are no oscillations, applied measures won't reflect oscillatory activity, but will return a value reflecting aperiodic activity	verify the presence of an oscillation, such as with spectral peak detection or with burst detection in the time domain
#2 Frequency Variation	neural oscillations have variable peak frequencies	measures applied using canonically defined frequency bands may fail to accurately capture oscillatory activity	verify frequency ranges and individualize as needed
#3 Aperiodic Activity	neural oscillations co-exist with dynamic aperiodic activity	measured variation may arise due to changes in aperiodic activity, rather than changes in oscillations	measure and control for changes in aperiodic neural activity, evaluating whether it explains measured changes
#4 Temporal Variability	neural oscillations are variable across time, exhibiting burst-like properties	burst properties may be conflated when analyzing spectral power, and trial averages may suggest illusory sustained activity	examine single trial data for temporal variation, and use burst detection to evaluate burst properties
#5 Waveform Shape	neural oscillations have non-sinusoidal waveform shape	analysis methods often assume sinusoidal structure, and may return spurious results in the case of non-sinusoidal oscillations	examine waveform shape measures to evaluate if waveform shape may underlie the results
#6 Overlapping Oscillations	multiple neural oscillations co-exist across the brain, and may overlap across space	multiple distinct sources may create destructive interference, in which case measures won't accurately reflect underlying activity	apply source separation techniques in order to reduce overlap of different types of oscillations
#7: Signal-to- Noise Ratio	neural oscillations have variable signal-to-noise ratio	without adequate signal to noise ratio, measures may be unreliable or inaccurate	evaluate the required signal-to-noise ratio, and potential ways to optimize it for all applied measures

#1 Neural oscillations are not always present

Why this matters

Neural field recordings are characterized not only by oscillatory activity, but also aperiodic ‘1/f’ or ‘1/f-like’ activity, in which signal power decreases exponentially as a function of frequency (Freeman et al., 2003; B. J. He, 2014). This is usually formalized as $1/f^\chi$ where χ represents the decay of power across frequencies. In neural data, χ often ranges between 0 and 4, where a signal with $\chi=0$ is white noise, with equal power across all frequencies, and higher values of χ indicate increasingly ‘steeper’ spectra. Aperiodic neural activity has been linked to the underlying activity of postsynaptic potentials and is a ubiquitous and sometimes dominant feature of neural field data (Gao et al., 2017; K. J. Miller et al., 2009).

The fact that aperiodic activity is omnipresent together with the large observable variability of neural oscillations (Donoghue et al., 2020b; Frauscher et al., 2018; Groppe et al., 2013) requires care in how band-limited power obtained by spectral analysis is measured and interpreted. Due to the presence of aperiodic activity, there is always non-zero power at all frequency bands. This means that any spectral measure—including computing a power spectrum, narrowband filtering, and average band-power measures—will always return a numerical value for power for a given frequency band, even if there is no oscillatory activity present. That is, just because there is power in a *frequency band* does not imply that there is an *oscillation* in that same frequency band (Bullock et al., 2003). It is a fallacy to presume that an analysis of a predefined narrowband frequency range necessarily reflects physiological oscillatory activity.

To introduce how transient and aperiodic signals are represented in the spectral domain, the Dirac delta can be used, whereby a single non-zero value in the time domain is represented by constant power across all frequencies in the frequency domain (Fig. 1A). This illustrates that power in a specific frequency band does not generally correspond to a present oscillation in the time domain. Similarly, 1/f-like aperiodic activity, which is common in neural data, shows power across all frequencies, with decreasing power for higher frequencies (Fig. 1B). Despite the lack of periodic activity in aperiodic time series, narrowband filtering, which imposes a sinusoidal basis, extracts components that appear to be oscillatory, when filtered into canonical band ranges (Fig. 1C). By comparison, rhythmic signals, such as a pure sinusoid, exhibit as a frequency specific peak in the power spectrum (Fig. 1D). Neural field recordings can be simulated as a summation of oscillatory and aperiodic components, resulting in a power spectrum that exhibits a spectral peak exceeding the aperiodic component, reflecting a high amount of band-specific power (Fig.

1E). In this case, the presence of the spectral peak is indicative of oscillatory power. In general, since different signal components can contribute to spectral power across different frequency ranges, power in a frequency band may not reflect oscillatory activity.

Recommendations

Investigations of oscillations should start with a *detection* step, verifying the presence of oscillations of interest. This verification step can be done in both the frequency and time domains. In the time domain, visualizing the data allows for examining if there are clear rhythmic segments in the data. In the frequency domain, oscillations manifest as peaks of power over and above the aperiodic signal (Buzsáki et al., 2013). As an initial check, visually inspecting power spectra can help to verify the presence of prominent oscillations. Including figures of power spectra in manuscripts is recommended, as it provides supporting evidence to the reader that there is oscillatory activity in the data under study.

Numerous quantitative methods also exist to detect oscillatory activity in neural field data, such as automated methods that detect narrowband spectral peaks (Pascual-Marqui et al., 1988). This can be systematically done by parameterizing the power spectrum, in which a mathematical model that quantifies periodic and aperiodic activity is applied to detect any putative oscillatory peaks above the measured aperiodic component (Donoghue et al., 2020b) (see Fig. 1F). Similarly, both the ‘multiple oscillation detection algorithm’ (MODAL) method (Watrous et al., 2018) and the ‘extended better oscillation detection’ (eBOSC) method (Kosciessa et al., 2020), which is itself an extension of prior methods (Caplan et al., 2015; Whitten et al., 2011), use a fit of the aperiodic activity to detect frequency specific activity.

It may also be useful to examine rhythmic properties of the data, to search for putative oscillatory activity in situations in which a spectral peak may be difficult to observe (Pesaran et al., 2018). For example, oscillations may be present in the form of rare or infrequent bursts, which will not appear as clear spectral peaks when the spectrum is calculated across the whole time interval. In such situations, examining shorter time ranges, and selecting time windows with higher band power and/or around events of interest may be required to resolve peaks in the frequency domain. Alternatively, time domain and burst detection methods, further described in sections 4 and 5, may be more applicable. Another potential approach for addressing this is lagged coherence (Fransen et al., 2015), which explicitly quantifies the rhythmicity in time series, in contrast to measuring solely spectral power, and can also be used to differentiate between oscillatory signals and transients (see Fig. 1A).

Because oscillations can vary in their presence within and between participants, and across different frequency bands (Donoghue et al., 2020b; Frauscher et al., 2018) oscillation detection should be performed for each frequency band of interest, participant and analyzed region. If oscillations are not detected, this may preclude further analyses. Group-level analyses may obscure variation in oscillatory presence in individual participants. For example, if not all participants display a clear rhythm, effect size estimates of oscillatory changes at the group level may be confounded by including the subset of participants without any clear oscillatory activity. Alternatively, a comparison of oscillatory power between regions without doing oscillation detection may conflate a change in oscillatory power with a difference in oscillatory presence. Analyses that include filtering or band-specific measures without first examining if an oscillation is present can provide ambiguous results that may reflect aperiodic activity, in which case it is a misinterpretation to describe physiological *oscillatory* activity. Applying analyses to detect oscillatory presence can assure that measures reflect oscillatory activity.

Figure 1: Without verified oscillatory activity, applied measures may reflect aperiodic activity. **A)** Non-oscillatory signals such as the dirac delta function exhibit power across all frequencies. **B)** Similarly, a non-oscillatory $1/f$ signal also has power across all frequencies, including canonical narrowband regions: delta (yellow), theta (green), alpha (blue), and beta (purple). This power spectrum illustrates the fact that just because there is power at a frequency, that does not imply there is a dominant oscillation at that frequency. **C)** Narrowband filtered traces of the signal shown in B, that appear to be rhythmic. Note that this reflects band-power from the aperiodic activity, rather than any narrowband oscillation. **D)** Rhythmic signals, such as a pure sinusoid, exhibit as a dominant peak in the power spectrum. **E)** A combined signal that contains aperiodic activity and a narrowband alpha oscillation. In this case, the oscillation is visible as a peak in the power spectrum above the spectral contribution from the aperiodic $1/f$ -like signal. **F)** Spectral peaks can be detected in order to identify putative oscillations in the data, as shown by the identified peak, in green. Spectral peak detection can be used to select frequency bands for further analysis, for example selecting the alpha range to be filtered for subsequent analysis (bottom).

#2 Neural oscillations vary in their peak frequencies

Why this matters

Neural oscillations display significant variations in their peak frequencies, including variation across age (Lindsley, 1939), within and between participants (Haegens et al., 2014), and across cortical locations (Mahjoory et al., 2020). Alpha peak frequency, for example, is considered a stable trait marker (Grandy et al., 2013), and is also associated with some clinical disorders, displaying, for example, a slower frequency in attention-deficit hyperactivity disorder (ADHD) (Lansbergen et al., 2011). The frequency of neural oscillations can also vary within participants within a task (Benwell et al., 2019), including in task relevant ways (Wutz et al., 2018).

Due to frequency variation, even if the presence of oscillations is verified, the use of canonically defined frequency ranges may still fail to accurately reflect the data, as this may misestimate power of an oscillation if the spectral peak is not well captured in the canonical range. For example, in Figure 2, a canonically defined alpha range of 8-12 Hz captures the peak of a 10 Hz oscillation (Fig. 2A), but fails to accurately capture a 8 Hz peak (Fig. 2B). Despite the signals being simulated with the same amount of oscillatory power, estimated alpha power using a canonical frequency range differs between the signals (Fig. 2C), due to an underestimate of the power in the signal with an idiosyncratic peak frequency. This issue also impacts the result of band-pass filtering, as a canonical filter range underestimates the amount of alpha power present, as compared to an individualized band in which the filter range is made to reflect the oscillation in the data (Fig. 2D). Using individualized frequency band ranges to control for frequency differences accurately captures the alpha power in each signal (Fig. 2E). Overall, predefined frequency band definitions may fail to address variation in peak frequencies, and lead to misestimations.

Potential differences in peak frequency are important for analyses that compute an estimate within a specific frequency range, such as calculating band power, or narrowband filtering to a frequency range of interest. Applying a fixed frequency range may lead to information loss when the individual peak frequency lies near the border or outside of the defined range; it can also be non-specific if the range captures an adjacent oscillation or aperiodic activity. These issues apply both to analyses of individual frequency bands, as well as to composite measures such as ratios computed between the power of different frequency bands, since variation in the peak frequency or bandwidth of peaks can impact measured results (Donoghue et al., 2020a). For example, what had previously been reported as a difference in the theta / beta ratio of

participants with ADHD was found to be partially driven by a slowed alpha peak in the ADHD group, changing the interpretation of the data (Lansbergen et al., 2011).

Recommendations

In order to address the variability of peak frequencies, any analyses that employ narrowband frequency ranges should assess how well the chosen ranges match the data. Visual inspection can help determine how well the defined frequency boundaries reflect actual peaks in the power spectra. This should be done for all analyzed frequency bands at the level of individual participants, because individual participants may have idiosyncratic peak frequencies that could influence group level results if they are misestimated. For within-subject analyses, changes in peak frequency over time or between tasks should also be considered in order to address whether a measured change in power could reflect a change in peak frequency, in which frequencies may 'drift' outside defined ranges of interest. Including power spectra in manuscripts also enables readers to observe that applied band ranges match the peaks observed in the data.

If canonically defined frequency ranges do appropriately match the data, then they can safely be used for subsequent analyses. However, if chosen band ranges of interest do not appropriately reflect the data, then individualized frequency bands may be applied (Klimesch, 1999). Methods for computing individualized frequency bands often do so by measuring spectral peaks (Haegens et al., 2014; Pascual-Marqui et al., 1988). Automated approaches have also been developed, that include spectral smoothing to improve performance (Corcoran et al., 2018). Such approaches don't always generalize to multiple peaks or bands, though some approaches use 'anchor frequencies' (Klimesch, 1999), defining, for example, theta as a range below the identified range of alpha. This approach has the limitation of not considering the oscillation detection step. Peak detection for multiple putative peaks, without predefining frequency ranges, can also be done with spectral parameterization (Donoghue et al., 2020b), after which peaks can be grouped into observed bands of interest.

Beyond spectral peak detection, methods for detecting oscillations can be used to detect frequencies with peak rhythmicity, for example, by applying lagged coherence across frequencies (Fransen et al., 2015). Some methods also allow for jointly learning multiple band definitions. For example, the Oscillation ReConstruction Algorithm (ORCA) evaluates multiple band definitions in terms of how well each definition is able to reconstruct the data (Watrous & Buchanan, 2020), and the gedBounds method identifies frequency boundaries by clustering similarities across frequencies (Cohen, 2021). These methods, which examine all analyzed frequencies together, may help to obtain more consistent groups of frequency ranges within and across participants.

350 Collectively, some form of evaluation needs to be done to verify frequency bands, in order to
351 ensure that applied measures accurately capture the intended oscillatory activity.
352

Figure 2: Canonical frequency band ranges may fail to capture narrowband peaks. **A)** A simulated signal, and corresponding power spectrum, with a prominent 10 Hz alpha oscillation. Shaded in blue is the canonical alpha range (8-12 Hz). **B)** Another signal with a prominent alpha oscillation, with a peak frequency of 8 Hz. **C)** Using canonical band ranges, the amount of alpha power is found to be significantly different between the signals from A & B. When examining adjacent frequency bands, (right), there is also a measured difference in theta power, due to the alpha peak drifting into the canonical theta range. These results suggest differences in oscillatory power between signals, however this is actually driven by a difference in alpha peak frequency. **D)** The time series from B, filtered into the alpha frequency range, using both the canonical range (blue) and an individualized range (green). The individualized range is tuned to the peak frequency of the time series (see inset power spectra). Note that the individualized filter captures more narrowband activity. **E)** Using individualized frequency bands, a difference in measured alpha power is no longer seen, consistent with the measured difference in C being due to a mismatch in peak frequency.

#3 Neural oscillations coexist with aperiodic activity

Why this matters

As previously introduced, neural field recordings contain aperiodic activity (B. J. He, 2014). This activity is not only ubiquitously present, but is also variable and dynamic within and between subjects (Freeman & Zhai, 2009; Podvalny et al., 2015). Between subject variability of aperiodic activity can relate to age (W. He et al., 2019; Voytek et al., 2015), and clinical diagnoses (Robertson et al., 2019), whereas within subjects, aperiodic activity varies with state, such as sleep (Lendner et al., 2020), relates to behavioral tasks (Podvalny et al., 2015) and can be influenced by exogenous stimuli and cognitive demands (Waschke et al., 2021). This dynamic aperiodic activity has different putative generators, physiological interpretations, and task related dynamics (Gao et al., 2017, 2020; K. J. Miller et al., 2009, 2014), as compared to oscillations, making it an interesting feature of interest in itself. Altogether, aperiodic neural activity is dynamic in many contexts in which neural oscillations are usually the focus of inquiry.

This dynamic quality of aperiodic activity is an important consideration for detecting neural oscillations, as previously discussed (see #1), as well as for interpreting measured changes in the data. With multiple dynamic components, analyses must adjudicate which aspects of the data are changing, and how, in order to allow for appropriate interpretations. Since aperiodic activity has power at all frequencies, changes or differences in aperiodic activity can induce patterns of differential activity across all frequencies. This can be seen by comparing white ($\chi = 0$) and pink ($\chi = 1$) noise $1/f^\chi$ signals, which have different amounts of power in a canonically defined alpha-band (Fig. 3A). Even with a validated spectral peak and frequency range, a difference in band-power between two conditions within a given frequency range may not be specific to oscillatory changes, as it may reflect a global change in aperiodic activity. For example, in Fig. 3B, a measured difference in alpha-band power between two conditions reflects a change in the aperiodic exponent, not changes relating to a spectral peak in the alpha-band, since the periodic activity is the same in the two signals.

Considering aperiodic activity is particularly important for analyses that investigate band-power across a series of frequency bands, since systematic patterns of measured changes across bands may not reflect any changes in oscillatory activity. For example, in Fig. 3C, the band-power of two conditions is compared across five different frequency bands. Despite this analysis suggesting a pattern of changes in band power across a series of canonically defined frequency bands (Fig. 3D), the changes are actually driven by a change in aperiodic activity. Patterns of

correlated changes across frequency bands can therefore sometimes be more parsimoniously explained by a change in aperiodic activity, rather than as multiple distinct oscillatory changes, as has been shown to be the case in development (W. He et al., 2019).

Changes in global power, due to aperiodic changes, can also impact relative or normalized measures of oscillatory activity. In the spectra in Fig. 3C, there is a visible spectral peak in the alpha-band. Even though there is no change in peak power, a relative power measure suggests a change in alpha power, due to a change in the power across all frequencies, that is driven by a change in aperiodic activity (Fig. 3E). This issue also impacts other compound measures, such as ratios of band-power, including the theta/beta-ratio, often investigated as a potential biomarker for ADHD (Lansbergen et al., 2011; Robertson et al., 2019), as it has been shown that band ratio measures often reflect a change of the aperiodic activity (Donoghue et al., 2020a), and that the putative relationship between ADHD and theta/beta-ratio appears to be driven by aperiodic activity (Robertson et al., 2019).

Recommendations

As both oscillatory and aperiodic components are dynamic, it is important for analyses to validate which elements of the data are specifically changing, in order to appropriately interpret results. This is relevant for any analysis investigating putative narrowband power, including investigations that examine multiple oscillation bands. Aperiodic activity should be explicitly measured to evaluate whether it explains the band-specific changes, including whether correlated patterns of changes across frequency bands may be more parsimoniously explained as a change in the broadband aperiodic activity. Approaches that assume oscillations exist upon a stationary 'background', such as relative power measures that divide by power across all frequencies, or band ratio measures, should be avoided, as they conflate changes in oscillatory and aperiodic components (Donoghue et al., 2020a). For example, a change in a relative power measure could arise from a change in band-specific power of interest, or be due to a change in aperiodic component that changes the measured power across all frequencies that is used as the denominator.

Explicitly measuring aperiodic activity requires methods that explicitly conceptualize both aperiodic and periodic activity, to avoid erroneously attributing aperiodic activity as oscillatory changes. Methods that define and measure oscillatory activity relative to aperiodic activity, including previously introduced methods such as spectral parameterization (Donoghue et al., 2020b) and eBOSC (Kosciessa et al., 2020), are designed to measure and control for aperiodic activity, and so address this issue. There are also dedicated methods for measuring aperiodic

activity. For example, the irregular-resampling auto-spectral analysis (IRASA) method leverages the scale-free nature of aperiodic activity by proposing a resampling procedure to isolate aperiodic activity (Wen & Liu, 2016). IRASA can be used to separate and measure aperiodic neural activity, after which analyses can evaluate each component to examine whether measures of interest specifically reflect the intended component. Overall, controlling for aperiodic activity requires employing an oscillation detection step and evaluating oscillatory power relative to the aperiodic component in order to assess whether measured changes are capturing oscillatory or aperiodic activity.

Figure 3: Variations in aperiodic activity influence band-power measures. **A)** Examples of aperiodic white (black) and pink (red) noise signals that display different patterns of power across frequencies, as seen in their power spectra. Shaded in blue is the canonical alpha range, with time-series filtered in the alpha-range shown in the inset. Note that the pink noise signal appears to have increased ‘alpha’ power. **B)** Simulated combined signals containing both aperiodic and oscillatory power (black), and a transformed version of the signal with the same periodic component with a change in the aperiodic component (red), after being rotated in the spectral domain. Note that in A & B, what appears to be band-specific changes actually reflect differences in aperiodic activity. **C)** A comparison between power spectra for combined signals simulated with the same oscillatory component with different aperiodic activity. Shading reflects different frequency bands, including delta (yellow), theta (green), alpha (blue), beta (purple) and gamma (red). **D)** Absolute differences in power, calculated separately for each frequency band, for the spectra in C. Note that despite the difference in the data being simulated as a change in the aperiodic component, a band-by-band analysis suggests a pattern of changes across distinct frequency bands. **E)** Relative alpha power (bottom) is calculated as absolute band power (top left), divided by the power across all frequencies (top right). Note that despite no difference in the amount of alpha power, there is measured change in relative power, due to systematically different aperiodic activity between the signals.

#4 Neural oscillations are variable across time

Why this matters

Neural oscillations often display burst-like temporal dynamics (Lundqvist et al., 2016; Sherman et al., 2016) and are rarely, if ever, completely consistent and continuous. These temporal dynamics of neural oscillations are a potentially important feature; for example, the rate of burst events has been found to be predictive of behavior across tasks and species (Shin et al., 2017), including in investigations of working memory (Lundqvist et al., 2016) and motor activity (Wessel, 2020). Some generative models of oscillations predict non-continuous events in a way that is consistent with what is seen in empirical data (Sherman et al., 2016).

Despite this, many methods implicitly assume stationarity of the signal under study, when analyzing, for example, average band power across time or trials. In such cases, variability of oscillation presence or temporal dynamics can be misinterpreted as differences in power. For example, in simulations with stochastic onset and offset of oscillatory activity, signals can display different proportions of the data with oscillatory activity present, with the oscillatory power when present is equivalent (Fig. 4A). Measured power in such cases reports a different amount of band specific power, typically interpreted as reflecting a change in the overall amplitude of the oscillation, however, measured differences can be due to temporal variability (Fig. 4B). These kinds of averaging effects are also important in scenarios such as time-frequency analyses that average across trials, which may create an illusion of sustained activity in averaged data (Feingold et al., 2015; Jones, 2016). This can happen if individual trials have burst-like temporal dynamics that occur at different times across different trials, which can average together in a way as to suggest a sustained response in average data, despite such continuity not occurring in any individual trial (Fig. 4C). The temporal variability of neural oscillations motivates the importance of considering single trial dynamics (Kosciessa et al., 2020; Rey et al., 2015; Stokes & Spaak, 2016).

Oscillatory bursts can vary in multiple ways that can lead to similar measured changes in band power, which may be misinterpreted as reflecting changes in tonic band power. This includes changes in burst duration (Fig. 4D), burst occurrence (Fig. 4E), or burst amplitude (4F), each of which can vary within or between analyzed time periods (Quinn et al., 2019; Zich et al., 2020). Understanding the different sources of variability has implications on how these signals should be interpreted, as a change in the length, number, or size of bursts each likely reflect different circuit mechanisms and putative relationships to neural function. However, this can not

be appropriately evaluated unless methods acknowledge oscillations as potentially transient, with potential variability in rate, timing, and duration as well as amplitude (van Ede et al., 2018).

Recommendations

Analyses of neural oscillations must therefore evaluate whether temporal variability, rather than overall power, may be driving measured changes. In order to address temporal variability, both the spectral and temporal domain have to be considered together (Zich et al., 2020). Time-frequency analyses, such as spectrograms, can be used to examine spectral properties across time in order to adjudicate between changes in the average power of oscillations and changes in their temporal dynamics. In doing so, it is important to analyze single-trials (Rey et al., 2015; Stokes & Spaak, 2016), to avoid misinterpreting averaged power. If reporting spectrograms, single-trial examples should be included in order to evaluate whether apparent sustained activity is truly sustained, or arises as a result of averaging many short bursts.

Burst detection methods can also be applied to identify segments of the signal in which oscillations are present, which can then be characterized in terms of the durations of the bursts, the number of bursts, and/or the amplitude of the bursts. A common approach for burst detection is to use an amplitude threshold, detecting segments of power in which frequency specific power is greater than a chosen threshold level (Feingold et al., 2015). The previously described eBOSC algorithm (Kosciessa et al., 2020) can be considered to be a threshold based burst detection, in which the threshold is based on the aperiodic component, and can be used for burst detection.

Other algorithms for burst detection include matching pursuit, in which a dictionary of atoms, which can include oscillatory bursts, is fit to the data, providing potentially more accurate estimates of burst onset and duration (Chandran KS et al., 2018). Alternatively, methods such as hidden markov modelling can be used, which seek to characterize state transitions, and can be used to model transitions into and out of oscillatory states in a probabilistic way (Quinn et al., 2019; Vidaurre et al., 2016). Time-domain measures that identify oscillations by characterizing individual cycles, further described in #5, can also be used to detect and analyze the number and duration of bursts, and their cycle-by-cycle properties (Cole & Voytek, 2019; Schaworonkow & Nikulin, 2019). After detection, analyses of burst-like neural activity typically involve subsequent analysis of the identified bursts, in order to evaluate whether they are changing in their duration, occurrence, and/or amplitude.

Figure 4: Temporal dynamics of neural oscillations influence spectral measures. **A)** Two simulated signals with lower (top; blue) and higher (bottom; green) levels of bursting activity in the alpha band, simulated with probabilistic burst onset and offset. Segments identified as bursts are shaded in red. Note that oscillation power, when present, is the same in both signals. **B)** Power spectra for the signals in A. Note the difference in size of the alpha peak, suggesting a difference in alpha power between the signals. However, when quantifying the power within the bursts (inset bar plot), the power is found to be approximately the same. The apparent difference in power is due to differences in temporal variability. **C)** Temporal variability can lead to spurious sustained power in averaged results. Spectrograms for individual trials (top) show short bursts of oscillatory power, which average to create what appears to be a sustained response (bottom). **D-F)** Measured differences in power can arise due to multiple features of bursting oscillations, including changes in the duration (**D**), occurrence (**E**), and/or amplitude (**F**) of the bursts. In these simulations, one feature differed between the two time series, while all others were held constant. Each feature creates a similar difference in the resultant alpha peaks, demonstrating that measured power can reflect multiple facets of temporal variability of analyzed time series.

#5 Neural oscillations are non-sinusoidal

Why this matters

The waveform shape of neural oscillations is often non-sinusoidal (Cole & Voytek, 2017; Jones, 2016), as seen, for example, in the arc-shaped sensorimotor mu-rhythm, visual alpha, which can be triangular, and the sawtooth-shaped hippocampal theta-rhythm. These waveform properties of neural oscillations may reflect physiological properties, for example the synchronization of neural activity (Schaworonkow & Nikulin, 2019), spiking patterns of underlying neurons (Cole & Voytek, 2018), or behavioral correlates such as running speed (Ghosh et al., 2020). Waveform shape can therefore be an important feature of interest, with potential to impose constraints on generative circuit models of oscillations (Sherman et al., 2016) as well as time constants of involved synaptic currents.

The variable waveform shape of oscillations also creates substantial methodological and interpretation hurdles, due to the assumed sinusoidal basis underlying most methods. For instance, estimating instantaneous phase typically involves narrowband filtering the signal before applying a Hilbert transform. Applying a narrowband filter on data with variations in waveform shape can be problematic, as the phases of sinusoidal outputs of narrowband filtering will not correspond to phases of a non-sinusoidal signal (Fig. 5A). This occurs because in the spectral domain, nonsinusoidal shapes are represented by power across multiple frequencies, and if the signal content in the harmonic frequencies is removed, the resulting filtered signal will have shifted peaks and troughs compared to the original non-sinusoidal signal (Fig. 5A). This is an important consideration for any analyses that examine cycle properties, such as the location of signal peaks and troughs, as putatively corresponding to specific physiological states. For analyses that rely on exact temporal characteristics (e.g. investigating the effects of pre-stimulus phase on behavioral measures), controlling for waveform shape may be beneficial.

In spectral analysis, non-sinusoidal waveforms are reflected in the power spectrum as harmonics occurring at multiples of the dominant frequency, as illustrated in Fig. 5B. This can result in interpreting these separate peaks as independent physiological rhythms. In the case of an arc-shaped mu-rhythm, for example, the waveform shape of the oscillation will create peaks in both the alpha- and beta-frequency ranges. This may be interpreted as separate alpha- and beta-rhythms with an assumed phase- and amplitude-coupled relationship, when in reality only one non-sinusoidal rhythm is present. Differentiating between those situations is complicated by the fact that several types of rhythms can be found across the cortex (see section #6). Fig. 5C

shows how the degree of non-sinusoidality is reflected in the power of harmonic frequencies, with higher power in the harmonic frequency range for increasing non-sinusoidality. This should be considered when evaluating differences in spectral power between conditions, to control for potential changes in waveform shape.

The spurious coupling that waveform shape can induce between frequencies (Kramer et al., 2008) is especially important when considering measures such as phase-amplitude coupling that are greatly influenced by waveform shape (Cole et al., 2017; Lozano-Soldevilla et al., 2016). Waveform shape can result in systematic changes in the amplitude at harmonic frequencies, as seen in Fig. 5D, which can depend on the phase of the base oscillation, as quantified in Fig. 5E. This results in significant measures of cross-frequency phase-amplitude coupling. Numerically, these values are not objectionable, as they reflect a relationship between frequencies in the spectral domain. However, there is possible fallacy in the interpretation, if this relationship is taken to reflect significant coupling between *independent* rhythms, when in fact no such interaction between multiple rhythms need exist. Because of these methodological limitations, careful work needs to be done to adjudicate between phase amplitude coupling measures that reflect waveform shape versus those that truly reflect nested oscillations (Giehl et al., 2021; Vaz et al., 2017).

Recommendations

In order to evaluate and control for waveform shape, explicit measurement of waveform and cycle properties should be done. Time domain measures of individual cycles can be used to characterize waveform shape by, for example, calculating measures such as the rise/decay symmetry or peak sharpness (Cole & Voytek, 2019; Schaworonkow & Nikulin, 2019). Other methods aim at learning and grouping waveforms into observed categories, for example through attempting to learn recurring patterns in the data by sliding-window matching (Gips et al., 2017) or by attempting to learn a dictionary of observed shapes in the data and finding occurrences of particular waveforms in the data based on templates (Barthélemy et al., 2013; Brockmeier & Principe, 2016; Jas & Dupré, 2017).

In the frequency domain, specific waveforms can create stereotypical patterns in power spectra and time-frequency representations, which can complicate the detection of oscillations (see #1). If spectral peaks are present at exact multiples of slower frequencies, quantifying waveform shape may help to distinguish between an independent oscillation at that particular frequency or harmonic spectral peaks induced by waveform shape. Since different waveform shapes may exhibit similar time-frequency representations (Jones, 2016), time-domain analyses

may be required to evaluate if and how waveform shape is contributing to spectral representations.

For cross-frequency coupling analysis, the frequency extent of local coupling within a region (e.g., for phase amplitude coupling, the range of higher frequencies that are coupled to the low frequency phase) can suggest whether it is likely to be genuine oscillatory coupling or a shape effect (Cole et al., 2017; Vaz et al., 2017), with narrow ranges at exactly multiples of the base frequencies indicative of possible coupling caused by waveform shape. Applying and comparing multiple measures of cross-frequency coupling can dissociate harmonic and non-harmonic phase-amplitude coupling (Giehl et al., 2021). More generally, frequency domain methods such as bicoherence, a measure of non-linear interactions between frequencies, can also be used to investigate waveform shape in the frequency domain (Bartz et al., 2019).

Figure 5: Waveform shape of neural oscillations influences power and coupling measures. **A)** Four different time domain signals with varying rise-decay asymmetry (colored traces) and their narrowband filtered versions (black traces). Narrowband filtering of asymmetric oscillations shifts the peak times of the signals as indicated by the shaded regions marking the distance between the peaks of original signal and the filtered version. **B)** In the corresponding power spectrum, there are emerging spectral peaks at harmonic frequencies (exactly two and three times the frequency) as a result of the asymmetry. **C)** The scale of these harmonic peaks relates to the asymmetry, such that increasing waveform asymmetry can exhibit as increased power in the beta-frequency range. **D)** Non-sinusoidal rhythms can also create spurious phase amplitude coupling. A 10 Hz non-sinusoidal alpha signal is band-pass filtered around the beta peak frequency (15 - 25 Hz). The beta signal shows deviations in amplitude depending on alpha phase driven by the non-sinusoidal waveform shape (inset shows power spectra for each signal). **E)** Phase amplitude coupling is quantified by calculating beta envelope as a function of alpha phase. In contrast to a pure beta-sinusoid, the beta envelope from the non-sinusoidal signal shows a minimum for a specific alpha phase, indicating phase-amplitude coupling, which is driven by the waveform shape of the alpha rhythm.

#6 Multiple oscillations coexist across the brain

Why this matters

Non-invasive recordings of neural oscillations reflect aggregate activity across relatively large cortical areas. Through volume conduction, a term used to describe the propagation of electrical fields from their original source across tissues to recording sensors, recording electrodes can reflect activity from multiple local sources, as well contributions from more distant sources that overlap both spatially as well as temporally (Buzsáki et al., 2012; Nunez & Srinivasan, 2006). For instance, in the context of MEG/EEG, there are several alpha-rhythm sources, with locations in somatosensory, occipital, parietal and temporal cortex (Hindriks et al., 2017), which can be co-active at the same time. In many studies, recordings are analyzed in sensor space, by directly analyzing activity from recording electrodes. In such cases, the aggregate signal may appear markedly different from the underlying sources of interest due to the spatial and temporal overlap of multiple distinct sources. Measures applied to these combined signals may therefore not accurately reflect the underlying sources, with distortions in measures of temporal dynamics or waveform shape (Schaworonkow & Nikulin, 2019).

Examining how spectral and time domain measures can be affected by overlapping sources is shown in an example in which sensor space activity from a single electrode is composed of activity from two underlying sources in the parietal and visual cortices (Fig. 6A). In the spectral domain, this configuration can result in two peaks in the alpha-frequency range (Fig. 6B), when the two sources have slightly different peak frequencies. This has been observed in empirical data as ‘double alpha’ or ‘split alpha’ peaks (Chiang et al., 2008). Analyses in sensor space may lead to the interpretation that a specific circuit generates signals with two simultaneously present peak frequencies, which in turn will influence theories of generating mechanisms. Spatial summation of multiple underlying rhythms of similar peak frequencies can also mask temporal features of interest of the underlying rhythms, as seen in Fig. 6C, due to constructive and destructive interference effects (Schaworonkow & Nikulin, 2019). Phase differences between sources of similar frequencies can attenuate the oscillation in sensor space, due to interference, even though oscillatory power has not changed in the underlying sources. This may lead to erroneous interpretations regarding changing oscillatory power of the sources, when it may be that only their relative temporal relationship has changed.

Inter-regional connectivity measures are also impacted by the simultaneous presence of multiple sources. Computing connectivity measures using sensor space signals can lead to

spurious findings, because volume conduction influences these measures (Haufe et al., 2013; J. M. Palva et al., 2018; S. Palva & Palva, 2012; Schoffelen & Gross, 2009). Because individual sources propagate to multiple sensors, regularities in amplitude and phase will be present across multiple sensors. This can yield highly significant statistical relationships between electrodes, reflecting signal content that is present due to a common source rather than genuine interregional coupling, which may lead to erroneous interpretation of connectivity between oscillatory sources.

Recommendations

Due to overlapping sources, analyzing sensor level time series or power spectra can be misleading regarding which aspects of the oscillation are present and/or are changing. Whenever possible, sensor space analysis should be complemented by source-level analysis. Source separation methods can be applied to attempt to separate different narrowband periodic components in the signal, which can help to reveal features that are not visible in sensor space data, as well as helping to localize sources. There are many possible approaches for source separation. Because inferring the activity of many more sources than channels is not possible, constraints are needed to arrive at a specific decomposition. The choice of the appropriate method also depends on the specific goals of source separation, including, for example, localizing activity to specific regions and/or decomposing time series into components based on statistical properties.

Based on these goals, two main approaches with different optimization criteria can be used for estimating source activity from sensor space activity. The first main type of methods use anatomical information to constrain the inverse solution based on individual or template structural MRI, in combination with methods such as beamformer or minimum norm estimation techniques (Hauk et al., 2019). The second main type of methods are agnostic to anatomical information and rely solely on the statistical structure of signals across channels. In this approach, channel activity is assumed to be a linear mixture of multiple underlying sources, defined by a leadfield matrix, which describes how individual sources map onto sensors (Parra et al., 2005). By assuming specific statistical properties of the source time series as well as mixing properties, demixing can be attempted. Methods in this realm include joint decorrelation (de Cheveigné & Parra, 2014) or independent component analysis (Hyvärinen & Oja, 2000). In the context of investigating neural oscillations, there are variants that specifically maximize SNR of narrowband oscillatory components, while minimizing SNR in flanking bands or in comparison to broadband activity. For enhancing oscillatory SNR, spatial-spectral decomposition (Nikulin et al., 2011) or generalized eigendecomposition (Cohen, 2017b) can be used. The Common Spatial Patterns algorithm

(Koles, 1991) and its variants (Lotte & Cuntai Guan, 2011) can be used for maximizing differences in narrowband activity between task conditions. For investigating relationships between narrowband activity and a continuous variable, Source Power Correlation analysis (Dähne et al., 2014) may be of interest. Spatial filtering methods can also be used as a preprocessing step for dimensionality reduction (Haufe, Dähne, et al., 2014), easing statistical comparisons and computational needs.

Components that result from source separation need validation, since different methods or parameter settings can yield highly different results, and solutions are not guaranteed to reflect physiologically meaningful activity. As such, source separation can be non-trivial and has its own set of methodological considerations as well as reporting guidelines (Cohen & Gulbinaite, 2014; Haufe, Meinecke, et al., 2014; Mahjoory et al., 2017). These guidelines can be used to evaluate robustness of the solution, such as with goodness of fit and/or localization error metrics, and to adequately convey reconstruction quality and method details to the reader.

Figure 6: Multiple simultaneous rhythms can interfere and impact sensor level data. **A)** A realistic head model with two oscillatory sources (red and blue) placed in the posterior cortex which project on the highlighted electrode (green). Underneath are the topographies of the two sources that contribute to the recording electrode. The leadfield coefficients for the two sources have approximately equal values, indicating equal contribution to the activity recorded at the green electrode. **B)** In this simulation, the electrode signal (green; bottom) reflects multiple underlying sources, including two distinct rhythmic components, with slightly different peak frequencies. These sources can be seen as two spectral peaks in the power spectrum. **C)** A separate simulation of two oscillatory sources with the same peak frequency, with a phase difference. Due to a phase difference of π , the two sources sum together destructively. In this scenario, interference of the sources cancel each other out at the electrode level, even though the oscillatory power of the individual sources is stable and consistent.

718 #7 Measures of neural oscillations require sufficient signal-to-noise ratio

719 Why this matters

720 Neural oscillations are embedded in complex recordings containing multiple rhythmic
 721 signals, aperiodic activity, and transient events. Analyzing oscillatory signals of interest requires
 722 defining features of interest (signal), and extracting this signal from the rest of the data (noise).
 723 As with all measures, methods for analyzing oscillations require an adequate signal to noise ratio
 724 (SNR). Indeed, ubiquitous processing steps such as filtering are used largely in order to increase
 725 the SNR (Widmann et al., 2015). Many of the considerations thus far (detecting oscillations,
 726 adjusting frequency ranges, controlling for aperiodic activity, burst detection, and source
 727 separation) can all be conceptualized as aiming to increase SNR by tuning analyses to specific
 728 properties of the data. Beyond these specific properties, applied measures can still be
 729 inaccurately estimated if SNR is low or variable.

730 The SNR of oscillatory activity relates to the ratio of oscillatory power to noise, typically
 731 the aperiodic background. Oscillatory power is a dynamic property, which can be seen by the
 732 variable height of oscillatory peaks over and above the aperiodic component (Fig. 7A). Many
 733 experimental paradigms will change oscillatory power, as presentation of stimuli may result in
 734 event-related attenuation of oscillations (Pfurtscheller & Lopes da Silva, 1999). This change in
 735 oscillatory power changes SNR, which in turn may influence accuracy and stability of other
 736 oscillatory measures such as instantaneous phase and frequency. When SNR is high, estimations
 737 of phase and frequency can be reliably estimated (Fig. 7B). However, when SNR is low,
 738 estimation can be very noisy (Sameni & Seraj, 2017) as can be seen in Fig. 7C, leading to
 739 artifactual large variations, often referred to as phase slips.

740 Changes in oscillatory power which change SNR and corrupt phase estimations can lead
 741 to inaccurate estimates of derived measures, such as the phase-locking value
 742 (Muthukumaraswamy & Singh, 2011) or inter-trial coherence (van Diepen & Mazaheri, 2018). Low
 743 SNR makes it difficult to reliably extract oscillations of interest (Fig. 7D), leading to variable phase
 744 estimates (Fig. 7E). When computing coupling measures on such estimates, differences in SNR,
 745 absent any true changes in phase alignment, can erode the detection of phase-locking between
 746 two signals (Fig. 7F). Unstable estimation of oscillatory measures can also propagate to
 747 multivariate analysis, such as cross-frequency coupling, whereby oscillatory power changes that
 748 influence SNR can lead to a change in measured cross-frequency coupling (Aru et al., 2015).

Time domain analyses, such as those designed for analyzing waveform shape, are also strongly dependent on their being adequate SNR to meaningfully measure the properties of interest.

In cases of low SNR, unreliable estimates could, for example, lead to false-negatives due to noisy estimations that are not able to adequately capture measures of interest. Conversely, certain analyses may return false positive results, if the measured variability of the signal is misinterpreted as a feature of interest, and/or leads to an artifactual measured change between conditions due to variable SNR. This may be an issue when comparing between groups who are known to have differences in relative power of oscillations, and/or when comparing within participants across conditions that may have different SNR.

Recommendations

Considering the SNR required for stable estimation of measures of oscillations starts by choosing appropriate experimental designs. When designing the protocol and tasks, experimenters should consider what is known about the reliability and effect size of effects of interest, and consider doing a power analysis to design well powered studies. This includes considering recording modalities, as different modalities have different sensitivities to different source locations (Piastra et al., 2020), as well as the different temporal, spatial, and frequency resolutions they offer. When recording the data, best practices should be employed to minimize non-neuronal noise, and use appropriate preprocessing in order to increase the quality of the data with the respect to desired analyses (Keil et al., 2014; Pernet et al., 2020).

Once recordings have been collected, or if considering existing datasets for potential re-analysis, signal-to-noise ratio has to be considered to validate if the dataset is appropriate for the desired analyses. This requires explicitly measuring SNR to verify that applied measures are robust in the SNR regime of the data. If the SNR is too low to provide accurate measurements, the analyses may be non-viable, as any measurements will be uninterpretable. If the analysis can be run, then SNR should still be continuously verified, to evaluate whether potential changes of SNR across time or between conditions may explain measured changes in results (van Diepen & Mazaheri, 2018).

General approaches for optimizing SNR include good filter design (de Cheveigné & Nelken, 2019; Widmann et al., 2015), and using information about spectral estimators and signals of interest to select the most appropriate methods to improve the accuracy and stability of estimates (Chavez et al., 2006; Lepage et al., 2013). There are also specific methods for more robust estimations of phase in low power situations, including Monte Carlo estimation (Sameni & Seraj, 2017) and applying a Kalman smoother (Mortezapouraghdam et al., 2018). Many of the

782 previously described methods such as detecting oscillatory peaks, using individualized frequency
783 ranges, and using burst detection can all improve SNR. Source separation techniques, including
784 those that explicitly optimize SNR (Cheveigné & Arzounian, 2015; Nikulin et al., 2011) can be
785 used to extract oscillatory components with higher SNR.
786

Figure 7: Low oscillatory signal-to-noise ratio impacts measures. **A)** Power spectra for simulated signals with variable SNR for an alpha oscillation, as seen in the different peak heights. **B)** One of the simulated signals, with a high SNR, with the alpha filtered signal (top; blue), from which the instantaneous phase (middle; red) and frequency (bottom; green) are computed. Note that the simulated signal has consistent phase and frequency. **C)** The same as B, for a signal with low SNR. Note that in this case, the estimates of phase and frequency are variable, due to misestimations because of the low SNR. This leads to phase slips, indicated by the arrows, in instantaneous phase, which also leads to erratic estimates of instantaneous frequency. **D)** Filtered versions of high and low SNR signals. In the simulated signals, the underlying signals (grey) are the same, other than a power difference, and have uniform phase. The filtered traces (blue) diverge from the underlying signal, especially in the low SNR signal. **E)** Phase estimates of the signals in D, in which the solid red is the true phase of the simulated oscillation, and the shading reflects the standard deviation of estimated phase across multiple iterations of phase estimation within each SNR regime. This shows that there is higher variance of phase estimates with lower SNR. These unstable phase estimates will impact subsequent measures, such as phase coupling. **F)** The phase locking value computed between a high powered oscillation, and simulated signals with decreasing power, as shown in A. Note that the simulated oscillations all have the same simulated phase time course, such that there is an expected phase locking value of 1, and any estimates below this are misestimations due to low power.

Discussion

How, and to what extent, neural oscillations are mechanistically involved in cognition remains undetermined. This lack of clarity likely arises in part from imprecisions in our methodological approaches for analyzing oscillations that, in turn, give rise to inconsistent results. Here, we highlight specific methodological considerations for analyzing and interpreting neural oscillations, providing explicit recommendations regarding each topic. These considerations acknowledge the heterogeneity of neural oscillations and embrace this complexity as an opportunity to consider ideas and interpretations that may help us to further understand our data. Oscillations vary in their presence and frequency, co-exist with dynamic aperiodic activity, have idiosyncratic temporal and waveform shape properties, overlap with one another, and require sufficient SNR to appropriately analyze. These topics also demonstrate that there is an increasing set of features that can be defined for neural oscillations, with an increasing toolkit of estimation methods. Hopefully, these recommendations can serve as guidelines for potentially reducing misinterpretations and conflicting results, and can increase clarity in our understanding of neural oscillations.

These considerations relate broadly to studies investigating neural oscillations, including investigations of endogenous activity, and/or of rhythmic neural activity that may be induced by stimulus presentation (Doelling et al., 2019; Lakatos et al., 2008). The potential impact of the considerations may vary across different studies. In many cases, these considerations may not change the analyses or interpretations, but may still offer potential avenues for further analyses, and deeper understanding of the data. In some situations, these considerations may greatly impact results and interpretations, potentially reflecting fundamental confounds that do need to be addressed, or even reflect issues that cannot be addressed by current methods, such that it precludes particular analyses from being appropriately applied. Overall, with a range of possible impacts, the general recommendation is to check for all of these possible issues, to identify which topics may matter in each scenario, and proceed accordingly.

Though we present the considerations as seven distinct points, it is important to note that these considerations do not manifest in isolation from one another and can interact. For example, variable aperiodic activity (#3) can interfere with spectral peak (#1) and/or burst (#4) detection, as it complicates approaches that use a threshold criterion to define bursts or spectral peaks. Oscillations may also be difficult to detect (#1) and/or to individualize frequencies for (#2) if they are temporally rare (#4), and/or have low SNR (#7). Further, waveform shape (#5) may systematically vary in relation to underlying sources (#6) (Schaworonkow & Nikulin, 2019) and/or

detected peaks (#1) may be volume conducted from remote sources (#6), resulting, for example, in ‘double alpha’ peaks due to the overlap of occipital and sensorimotor rhythms in the alpha-band (Chiang et al., 2008). Multiple oscillatory features, such as power, waveform shape, burst rate, etc., can covary. These potential multicollinearities need to be explicitly considered and tested by robust analyses that control for multiple potentially confounding features by, for example, addressing overlapping periodic and aperiodic activity (Donoghue et al., 2020b; Kosciessa et al., 2020), controlling for waveform shape, which may result in spurious power- and/or phase-coupling (Cole & Voytek, 2019; Schaworonkow & Nikulin, 2019), and examining trial-by-trial dynamics that may be masked or conflated in average measures (Jones, 2016; Stokes & Spaak, 2016; Zich et al., 2020).

This investigation used a simulation approach that attempts to mimic the properties seen in empirical data, including dynamic aperiodic activity, and oscillatory components that can vary across multiple features (Cole et al., 2019). Because ground-truth properties of physiological data are not known in a way that can be used to evaluate the accuracy of applied measures, simulated data are an important tool for diagnosing available methods. In using simulated data, we must endeavor to reflect on our empirical data—simulating heterogeneous oscillatory features embedded within dynamic aperiodic activity—in order to be representative of empirical data and realistic use cases. As well as the tool used here, there are other approaches for simulating data, including for specific modalities such as EEG (Krol et al., 2018), or that emulate neural circuits (Neymotin, Daniels, et al., 2020), or whole brain recordings (Sanz Leon et al., 2013). Simulation analyses should be employed when developing new analysis approaches, as novel methods require validation and comparison to existing methods, such that best practice guidelines can be continuously developed and updated. All time-frequency methods include settings that should also be validated and explored. Sensitivity analyses, in which one repeats the analyses across mild perturbations of method settings to evaluate the robustness of the measured results, should be used to ensure that results are not overly dependent on specific parameter regimes.

Estimates of oscillatory features of interest are typically further analyzed and compared using statistical methods. Notably, many neuroscientific parameters exhibit skewed distributions (Buzsáki & Mizuseki, 2014), including oscillatory power (Kiebel et al., 2005). Therefore, distributional properties of data should be carefully considered such that appropriate statistical tests can be chosen (Maris, 2012; Maris & Oostenveld, 2007). This is especially important when considering that power-law distributed variables can result in spurious correlations when using methods that assume normality (Schaworonkow et al., 2015). Statistical analyses, in particular in the context of new methods and measures, should also evaluate consistency across participants

(Grice et al., 2020), reliability within participants, and effect size measures, which can be computed using estimation statistics (Calin-Jageman & Cumming, 2019). Considering effect sizes can also aid in designing studies that are sufficiently powered (Button et al., 2013). Adopting the best practices proposed here may also help to increase statistical power, insofar as they help to better and more specifically characterize features of interest, improving SNR.

In our examples, we focused primarily on univariate measures, such as estimating oscillatory power or phase. Issues that affect these estimates also propagate to derived measures, such as correlations between amplitude or phase, as is done in functional connectivity (Haufe et al., 2013) and cross-frequency coupling analyses (Aru et al., 2015). If phase estimates are unreliable due to low oscillatory SNR (Sameni & Seraj, 2017), or if amplitude estimates are biased by changes in aperiodic activity (Donoghue et al., 2020b), or if burst properties vary between analyzed signals (Jones, 2016), then derived measures may fail to reflect the intended oscillatory properties. Methodological limitations are likely to propagate and compound in multivariate or mass univariate analyses, and must therefore be considered for any analyses including, or built on top, of the univariate methods demonstrated here.

Though beyond the scope of this article, investigations of neural oscillations also require employing best-practices for designing, collecting, and preprocessing data in order to ensure sound research design, high quality data, and methodological validity. These considerations are covered in available textbooks (Cohen, 2014; Hari & Puce, 2017), as well as individual reports that discuss topics such as including best practices for reporting and conducting MEG/EEG research (Gross et al., 2013; Keil et al., 2014; Pernet et al., 2020), pre-processing (de Cheveigné & Arzounian, 2018), artifact rejection and data cleaning (Jas et al., 2017; Urigüen & Garcia-Zapirain, 2015), and guides to using common software tools such as MNE (Gramfort, 2013; Jas et al., 2018) and FieldTrip (Oostenveld et al., 2011; Popov et al., 2018). Other work also features dedicated discussion for specific methods such as filtering (de Cheveigné & Nelken, 2019; Widmann et al., 2015), phase estimations (Chavez et al., 2006; Lepage et al., 2013), functional connectivity (O'Neill et al., 2018), and cross-frequency coupling analyses (Aru et al., 2015).

Broader strategies are also required for addressing reproducibility in the field of neural oscillations, including pursuing replication studies, providing clear descriptions of methods and results, and publishing null results (Cohen, 2017a). Open-science practices, including making data and analysis code available, can help foster reproducibility and develop transparency (Gleeson et al., 2017; Kathawalla et al., 2020; Voytek, 2016). Due to their computational nature, investigations of neural oscillations also benefit from good code practice (Wilson et al., 2017). Standardized procedures for organizing datasets also increase shareability, organization, and can

assist in standardized pipelines, making it easier to apply novel methods (Holdgraf et al., 2019; Niso et al., 2018; Pernet et al., 2019). Adopting open science practices provides opportunities for using open tools and datasets that can foster transparency and efficiently allow for revisiting the evidence for how neural oscillations relate to cognition and disease.

Importantly, these considerations also reflect opportunities for developing new theory and understanding of neural field data, which is still in many ways a mystery (Cohen, 2017c). Aperiodic activity is itself a physiologically informative feature (Gao et al., 2017, 2020), reflecting processes distinct from neural oscillations (Donoghue et al., 2020b; B. J. He, 2014). New methods provide new opportunities, for example, the ability to jointly analyze multiple components of the data, such as how oscillations and aperiodic activity jointly contribute to cognitive processing (Cross et al., 2020). New features of interest offer the potential for better understanding underlying physiology and putative computational roles of neural oscillations. For example, modelling that explicitly considers waveform shape and/or burst properties has contributed to physiological models of neocortical beta generation (Sherman et al., 2016), and models proposing mechanisms of beta and gamma activity in working memory (E. K. Miller et al., 2018).

Our emerging understanding of the data under study and how to measure it provides new vistas of opportunity for continuing to understand neural field data, and how it relates to cognition and disease. These methods and topics reflect the current status of methodological considerations for research related to neural oscillations. As our understanding of the many complexities of neural data continues to evolve, future investigations of neural oscillations must continue a consistent process of interrogating the assumptions of our methods and how they relate to current knowledge of the data to validate measures of the data, and develop evolving best practices.

Conclusion

Productively investigating neural oscillations requires dedicated and carefully applied methods that reflect our current understanding of the data. As methodological validity is a prerequisite for appropriate interpretation, analysis methods must reflect that neural field data consists of a complex combination of multiple oscillatory components, variable aperiodic activity, and transient events, within which oscillations vary across multiple dimensions. Here, we have proposed a checklist of methodological considerations for neural oscillations, with recommendations to 1) validate that oscillations are present; 2) verify that used frequency ranges are appropriate; 3) control for potential confounds due to aperiodic activity; consider the 4) temporal variation and 5) waveform shape of neural oscillations; 6) apply source separation, as needed, to separate multiple oscillatory processes; and 7) evaluate that the SNR is adequate for the analyses at hand. These considerations, and new methods that have been developed to address them, reflect our emerging understanding of neural field data and offer new possibilities for investigating, and ultimately, understanding, neural oscillations.

Materials and Methods

A simulation-based approach was used to create the demonstrations in this manuscript. Simulated time series were created with the NeuroDSP toolbox (Cole et al., 2019), version 2.2.0. In most cases, the time series were created as a combination of oscillatory and aperiodic activity, sampled at 1000 Hz. Oscillatory activity was simulated as sine waves unless otherwise noted. Each oscillation was simulated at a specific frequency, typically in the alpha band, unless otherwise specified. Aperiodic activity was simulated by spectrally rotating white noise to the desired 1/f exponent (Timmer & Konig, 1995). Aperiodic and oscillatory signal components were weighted according to a specified variance and combined together in an additive manner. Across all analyses, power spectra were estimated using Welch's method (Welch, 1967), using Hanning windowed 1 second segments with 12.5% overlap. Filtering was done with finite impulse response bandpass filters, with linear phase and filter lengths set to a default of 3 cycles of the highpass frequency, and enforced to be odd (Type I). Canonical band ranges were defined as delta (2-4 Hz), theta (4-8 Hz), alpha (8-13 Hz), and beta (13-30 Hz), unless otherwise specified. Analysis methods were also used as available in the NeuroDSP toolbox, or with custom code included in the project repository (<https://github.com/voytekresearch/oscillationmethods>).

Several of the figure demonstrations used additional processing. For the peak detection in Figure 1, the spectral peak was detected and quantified using spectral parameterization, which models the power spectrum as a combination of aperiodic and oscillatory components, and can be used to detect peaks of putative oscillatory power over and above the measured aperiodic component (Donoghue et al., 2020b). For the individual frequency example in Figure 2, canonical alpha was defined as ± 2 Hz around 10 Hz, and individualized alpha bands were defined as ± 2 Hz around the individual peak frequency. For the demonstrations of varying aperiodic activity in Figure 3, generated time series were spectrally rotated, in the same manner as done to simulate the aperiodic activity (Timmer & Konig, 1995). Relative power was computed as the sum of power in a frequency band of interest, divided by the sum of power across all frequencies in the frequency range of 2-50 Hz.

For the temporal variation demonstrations in Figure 4, bursty oscillations were simulated by specifying time segments that should include an oscillation, optionally controlling the duration, occurrence, and amplitude of the bursts. Burst specific power was calculated by sub-selecting segments of the data with an oscillation present. For the examinations of waveform shape in Figure 5, oscillations were simulated as asymmetric sine waves, and the bycycle toolbox (version 1.0.0) was used to quantify waveform shape in the time domain (Cole & Voytek, 2019). For this,

977 signals were band-pass filtered around the frequency of interest (here: 10 Hz) to extract the time
978 points of zero-crossings of the signal. The time points were used to segment the broadband data
979 into cycles, determining several cycle parameters. For this example, simulated time series were
980 created with varying rise-decay symmetry, which is the ratio of time in the rising and decaying
981 segments of the oscillation, which creates asymmetric oscillations.

982 For the spatial mixing demonstration in Figure 6, the New York Head (ICBM-NY) was used
983 (Huang et al., 2016) as a head model. Two sources are placed in the posterior cortex, and the
984 corresponding sensor signals are calculated using the leadfield. Oscillations were simulated as
985 asymmetric waves, created as the sum of two sines waves with a fixed phase lag. Topographies
986 were visualized using MNE-python (Gramfort, 2013). In Figure 7, instantaneous measures were
987 computed by applying the Hilbert transform to signals that had been bandpass filtered into the
988 alpha range (8-12 Hz), taking the angle as the phase estimate, and using the derivative of the
989 instantaneous phase as a measure of instantaneous frequency. Phase synchrony was measured
990 using the phase locking value (Lachaux et al., 1999).

991

References

- Aru, J., Aru, J., Priesemann, V., Wibral, M., Lana, L., Pipa, G., Singer, W., & Vicente, R. (2015). Untangling cross-frequency coupling in neuroscience. *Current Opinion in Neurobiology*, 31, 51–61. <https://doi.org/10.1016/j.conb.2014.08.002>
- Babiloni, C., Barry, R. J., Başar, E., Blinowska, K. J., Cichocki, A., Drinkenburg, W. H. I. M., Klimesch, W., Knight, R. T., Lopes da Silva, F., Nunez, P., Oostenveld, R., Jeong, J., Pascual-Marqui, R., Valdes-Sosa, P., & Hallett, M. (2020). International Federation of Clinical Neurophysiology (IFCN) – EEG research workgroup: Recommendations on frequency and topographic analysis of resting state EEG rhythms. Part 1: Applications in clinical research studies. *Clinical Neurophysiology*, 131(1), 285–307. <https://doi.org/10.1016/j.clinph.2019.06.234>
- Barthélemy, Q., Gouy-Pailler, C., Isaac, Y., Souiloumiac, A., Larue, A., & Mars, J. I. (2013). Multivariate temporal dictionary learning for EEG. *Journal of Neuroscience Methods*, 215(1), 19–28. <https://doi.org/10.1016/j.jneumeth.2013.02.001>
- Bartz, S., Avarvand, F. S., Leicht, G., & Nolte, G. (2019). Analyzing the waveshape of brain oscillations with bicoherence. *NeuroImage*, 188, 145–160. <https://doi.org/10.1016/j.neuroimage.2018.11.045>
- Başar, E. (2013). Brain oscillations in neuropsychiatric disease. *Dialogues in Clinical Neuroscience*, 15(3), 291–300. <https://doi.org/10.31887/DCNS.2013.15.3/ebasar>
- Başar, E., Başar-Eroğlu, C., Karakaş, S., & Schürmann, M. (2001). Gamma, alpha, delta, and theta oscillations govern cognitive processes. *International Journal of Psychophysiology*, 39(2), 241–248. [https://doi.org/10.1016/S0167-8760\(00\)00145-8](https://doi.org/10.1016/S0167-8760(00)00145-8)
- Benwell, C. S. Y., London, R. E., Tagliabue, C. F., Veniero, D., Gross, J., Keitel, C., & Thut, G. (2019). Frequency and power of human alpha oscillations drift systematically with time-on-task. *NeuroImage*, 192, 101–114. <https://doi.org/10.1016/j.neuroimage.2019.02.067>
- Brazier, M. A. B. (1958). The Development of Concepts Relating to the Electrical Activity of the Brain. *The Journal of Nervous and Mental Disease*, 126(4), 303–321.
- Brockmeier, A. J., & Principe, J. C. (2016). Learning Recurrent Waveforms Within EEGs. *IEEE Transactions on Biomedical Engineering*, 63(1), 43–54. <https://doi.org/10.1109/TBME.2015.2499241>
- Bruns, A. (2004). Fourier-, Hilbert- and wavelet-based signal analysis: Are they really different approaches? *Journal of Neuroscience Methods*, 137(2), 321–332. <https://doi.org/10.1016/j.jneumeth.2004.03.002>
- Bullock, T. H., McClune, M. C., & Enright, J. T. (2003). Are the electroencephalograms mainly rhythmic? Assessment of periodicity in wide-band time series. *Neuroscience*, 121(1), 233–252. [https://doi.org/10.1016/S0306-4522\(03\)00208-2](https://doi.org/10.1016/S0306-4522(03)00208-2)
- Button, K. S., Ioannidis, J. P. A., Mokrysz, C., Nosek, B. A., Flint, J., Robinson, E. S. J., & Munafò, M. R. (2013). Power failure: Why small sample size undermines the reliability of neuroscience. *Nature Reviews Neuroscience*, 14(5), 365–376. <https://doi.org/10.1038/nrn3475>
- Buzsáki, G., Anastassiou, C. A., & Koch, C. (2012). The origin of extracellular fields and currents—EEG, ECoG, LFP and spikes. *Nature Reviews Neuroscience*, 13(6), 407–420. <https://doi.org/10.1038/nrn3241>

- Buzsáki, G., & Draguhn, A. (2004). Neural oscillations in cortical networks. *Science*, 304(5679), 1926–1929. <https://doi.org/10.1126/science.1099745>
- Buzsáki, G., Logothetis, N., & Singer, W. (2013). Scaling Brain Size, Keeping Timing: Evolutionary Preservation of Brain Rhythms. *Neuron*, 80(3), 751–764. <https://doi.org/10.1016/j.neuron.2013.10.002>
- Buzsáki, G., & Mizuseki, K. (2014). The log-dynamic brain: How skewed distributions affect network operations. *Nature Reviews Neuroscience*, 15(4), 264–278. <https://doi.org/10.1038/nrn3687>
- Buzsáki, G., & Watson, B. O. (2012). Brain Rhythms and Neural Syntax: Implications for Efficient Coding of Cognitive Content and Neuropsychiatric Disease. *Dialogues in Clinical Neuroscience*, 14(4), 345–367. <https://doi.org/10.31887/DCNS.2012.14.4/gbuzsaki>
- Calin-Jageman, R. J., & Cumming, G. (2019). Estimation for Better Inference in Neuroscience. *ENEURO*, 6(4), ENEURO.0205-19.2019. <https://doi.org/10.1523/ENEURO.0205-19.2019>
- Caplan, J. B., Bottomley, M., Kang, P., & Dixon, R. A. (2015). Distinguishing rhythmic from non-rhythmic brain activity during rest in healthy neurocognitive aging. *NeuroImage*, 112, 341–352. <https://doi.org/10.1016/j.neuroimage.2015.03.001>
- Chandran KS, S., Seelamantula, C. S., & Ray, S. (2018). Duration analysis using matching pursuit algorithm reveals longer bouts of gamma rhythm. *Journal of Neurophysiology*, 119(3), 808–821. <https://doi.org/10.1152/jn.00154.2017>
- Chavez, M., Besserve, M., Adam, C., & Martinerie, J. (2006). Towards a proper estimation of phase synchronization from time series. *Journal of Neuroscience Methods*, 154(1–2), 149–160. <https://doi.org/10.1016/j.jneumeth.2005.12.009>
- Cheveigné, A. de, & Arzounian, D. (2015). Scanning for oscillations. *Journal of Neural Engineering*, 12(6), 066020. <https://doi.org/10.1088/1741-2560/12/6/066020>
- Chiang, A. K. I., Rennie, C. J., Robinson, P. A., Roberts, J. A., Rigozzi, M. K., Whitehouse, R. W., Hamilton, R. J., & Gordon, E. (2008). Automated characterization of multiple alpha peaks in multi-site electroencephalograms. *Journal of Neuroscience Methods*, 168(2), 396–411. <https://doi.org/10.1016/j.jneumeth.2007.11.001>
- Cohen, M. X. (2014). *Analyzing neural time series data: Theory and practice*. MIT Press.
- Cohen, M. X. (2017a). Rigor and replication in time-frequency analyses of cognitive electrophysiology data. *International Journal of Psychophysiology*, 111, 80–87. <https://doi.org/10.1016/j.ijpsycho.2016.02.001>
- Cohen, M. X. (2017b). Comparison of linear spatial filters for identifying oscillatory activity in multichannel data. *Journal of Neuroscience Methods*, 278, 1–12. <https://doi.org/10.1016/j.jneumeth.2016.12.016>
- Cohen, M. X. (2017c). Where Does EEG Come From and What Does It Mean? *Trends in Neurosciences*, 40(4), 208–218. <https://doi.org/10.1016/j.tins.2017.02.004>
- Cohen, M. X. (2021). A data-driven method to identify frequency boundaries in multichannel electrophysiology data. *Journal of Neuroscience Methods*, 347, 108949. <https://doi.org/10.1016/j.jneumeth.2020.108949>
- Cohen, M. X., & Gulbinaite, R. (2014). Five methodological challenges in cognitive electrophysiology. *NeuroImage*, 85, 702–710. <https://doi.org/10.1016/j.neuroimage.2013.08.010>

- Cole, S. R., Donoghue, T., Gao, R., & Voytek, B. (2019). NeuroDSP: A package for neural digital signal processing. *Journal of Open Source Software*, 4(36), 1272. <https://doi.org/10.21105/joss.01272>
- Cole, S. R., van der Meij, R., Peterson, E. J., de Hemptinne, C., Starr, P. A., & Voytek, B. (2017). Nonsinusoidal Beta Oscillations Reflect Cortical Pathophysiology in Parkinson's Disease. *The Journal of Neuroscience*, 37(18), 4830–4840. <https://doi.org/10.1523/jneurosci.2208-16.2017>
- Cole, S. R., & Voytek, B. (2017). Brain Oscillations and the Importance of Waveform Shape. *Trends in Cognitive Sciences*, 21(2), 137–149. <https://doi.org/10.1016/j.tics.2016.12.008>
- Cole, S. R., & Voytek, B. (2018). *Hippocampal theta bursting and waveform shape reflect CA1 spiking patterns* [Preprint]. bioRxiv. <https://doi.org/10.1101/452987>
- Cole, S. R., & 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>
- Corcoran, A. W., Alday, P. M., Schlesewsky, M., & Bornkessel-Schlesewsky, I. (2018). Toward a reliable, automated method of individual alpha frequency (IAF) quantification. *Psychophysiology*, 55(7), e13064. <https://doi.org/10.1111/psyp.13064>
- Cross, Z. R., Corcoran, A. W., Schlesewsky, M., Kohler, Mark. J., & Bornkessel-Schlesewsky, I. (2020). *Oscillatory and aperiodic neural activity jointly predict grammar learning* [Preprint]. bioRxiv. <https://doi.org/10.1101/2020.03.10.984971>
- Dähne, S., Meinecke, F. C., Haufe, S., Höhne, J., Tangermann, M., Müller, K.-R., & Nikulin, V. V. (2014). SPoC: A novel framework for relating the amplitude of neuronal oscillations to behaviorally relevant parameters. *NeuroImage*, 86, 111–122. <https://doi.org/10.1016/j.neuroimage.2013.07.079>
- de Cheveigné, A., & Arzounian, D. (2018). Robust detrending, rereferencing, outlier detection, and inpainting for multichannel data. *NeuroImage*, 172, 903–912. <https://doi.org/10.1016/j.neuroimage.2018.01.035>
- de Cheveigné, A., & Nelken, I. (2019). Filters: When, Why, and How (Not) to Use Them. *Neuron*, 102(2), 280–293. <https://doi.org/10.1016/j.neuron.2019.02.039>
- de Cheveigné, A., & Parra, L. C. (2014). Joint decorrelation, a versatile tool for multichannel data analysis. *NeuroImage*, 98, 487–505. <https://doi.org/10.1016/j.neuroimage.2014.05.068>
- Doelling, K. B., Assaneo, M. F., Bevilacqua, D., Pesaran, B., & Poeppel, D. (2019). An oscillator model better predicts cortical entrainment to music. *Proceedings of the National Academy of Sciences*, 201816414. <https://doi.org/10.1073/pnas.1816414116>
- Donoghue, T., Dominguez, J., & Voytek, B. (2020a). Electrophysiological Frequency Band Ratio Measures Conflate Periodic and Aperiodic Neural Activity. *Eneuro*, ENEURO.0192-20.2020. <https://doi.org/10.1523/ENEURO.0192-20.2020>
- Donoghue, T., Haller, M., Peterson, E. J., Varma, P., Sebastian, P., Gao, R., Noto, T., Lara, A. H., Wallis, J. D., Knight, R. T., Shestyuk, A., & Voytek, B. (2020b). Parameterizing neural power spectra into periodic and aperiodic components. *Nature Neuroscience*, 23(12), 1655–1665. <https://doi.org/10.1038/s41593-020-00744-x>
- Feingold, J., Gibson, D. J., DePasquale, B., & Graybiel, A. M. (2015). Bursts of beta oscillation differentiate postperformance activity in the striatum and motor cortex of monkeys performing movement tasks. *Proceedings of the National Academy of Sciences*,

- 1124 112(44), 13687–13692. <https://doi.org/10.1073/pnas.1517629112>
- 1125 Fransen, A. M. M., van Ede, F., & Maris, E. (2015). Identifying neuronal oscillations using
1126 rhythmicity. *NeuroImage*, 118, 256–267.
1127 <https://doi.org/10.1016/j.neuroimage.2015.06.003>
- 1128 Frauscher, B., von Ellenrieder, N., Zemann, R., Doležalová, I., Minotti, L., Olivier, A., Hall, J.,
1129 Hoffmann, D., Nguyen, D. K., Kahane, P., Dubeau, F., & Gotman, J. (2018). Atlas of the
1130 normal intracranial electroencephalogram: Neurophysiological awake activity in different
1131 cortical areas. *Brain*, 141(4), 1130–1144. <https://doi.org/10.1093/brain/awy035>
- 1132 Freeman, W. J., Holmes, M. D., Burke, B. C., & Vanhatalo, S. (2003). Spatial spectra of scalp
1133 EEG and EMG from awake humans. *Clinical Neurophysiology*, 114(6), 1053–1068.
1134 [https://doi.org/10.1016/S1388-2457\(03\)00045-2](https://doi.org/10.1016/S1388-2457(03)00045-2)
- 1135 Freeman, W. J., & Zhai, J. (2009). Simulated power spectral density (PSD) of background
1136 electrocorticogram (ECoG). *Cognitive Neurodynamics*, 3(1), 97–103.
1137 <https://doi.org/10.1007/s11571-008-9064-y>
- 1138 Fries, P. (2005). A mechanism for cognitive dynamics: Neuronal communication through
1139 neuronal coherence. *Trends in Cognitive Sciences*, 9(10), 474–480.
1140 <https://doi.org/10.1016/j.tics.2005.08.011>
- 1141 Gao, R., Peterson, E. J., & Voytek, B. (2017). Inferring synaptic excitation/inhibition balance
1142 from field potentials. *NeuroImage*, 158, 70–78.
1143 <https://doi.org/10.1016/j.neuroimage.2017.06.078>
- 1144 Gao, R., van den Brink, R. L., Pfeffer, T., & Voytek, B. (2020). Neuronal timescales are
1145 functionally dynamic and shaped by cortical microarchitecture. *ELife*, 9, e61277.
1146 <https://doi.org/10.7554/eLife.61277>
- 1147 Ghosh, M., Shanahan, B. E., Furtak, S. C., Mashour, G. A., Burwell, R. D., & Ahmed, O. J.
1148 (2020). Instantaneous Amplitude and Shape of Postrhinal Theta Oscillations
1149 Differentially Encode Running Speed. *Behavioral Neuroscience*, 134(6), 516–528.
- 1150 Giehl, J., Noury, N., & Siegel, M. (2021). Dissociating harmonic and non-harmonic phase-
1151 amplitude coupling in the human brain. *NeuroImage*, 227.
1152 <https://doi.org/10.1016/j.neuroimage.2020.117648>
- 1153 Gips, B., Bahramisharif, A., Lowet, E., Roberts, M. J., de Weerd, P., Jensen, O., & van der
1154 Eerden, J. (2017). Discovering recurring patterns in electrophysiological recordings.
1155 *Journal of Neuroscience Methods*, 275, 66–79.
1156 <https://doi.org/10.1016/j.jneumeth.2016.11.001>
- 1157 Gleeson, P., Davison, A. P., Silver, R. A., & Ascoli, G. A. (2017). A Commitment to Open
1158 Source in Neuroscience. *Neuron*, 96(5), 964–965.
1159 <https://doi.org/10.1016/j.neuron.2017.10.013>
- 1160 Gramfort, A. (2013). MEG and EEG data analysis with MNE-Python. *Frontiers in Neuroscience*,
1161 7, 267. <https://doi.org/10.3389/fnins.2013.00267>
- 1162 Grandy, T. H., Werkle-Bergner, M., Chicherio, C., Schmiedek, F., Lövdén, M., & Lindenberger,
1163 U. (2013). Peak individual alpha frequency qualifies as a stable neurophysiological trait
1164 marker in healthy younger and older adults: Alpha stability. *Psychophysiology*, 50(6),
1165 570–582. <https://doi.org/10.1111/psyp.12043>
- 1166 Grice, J. W., Medellin, E., Jones, I., Horvath, S., McDaniel, H., O'lansen, C., & Baker, M. (2020).
1167 Persons as Effect Sizes. *Advances in Methods and Practices in Psychological Science*,

- 3(4), 443–455. <https://doi.org/10.1177/2515245920922982>
- Groppe, D. M., Bickel, S., Keller, C. J., Jain, S. K., Hwang, S. T., Harden, C., & Mehta, A. D. (2013). Dominant frequencies of resting human brain activity as measured by the electrocorticogram. *NeuroImage*, 79, 223–233. <https://doi.org/10.1016/j.neuroimage.2013.04.044>
- Gross, J. (2014). Analytical methods and experimental approaches for electrophysiological studies of brain oscillations. *Journal of Neuroscience Methods*, 228, 57–66. <https://doi.org/10.1016/j.jneumeth.2014.03.007>
- Gross, J., Baillet, S., Barnes, G. R., Henson, R. N., Hillebrand, A., Jensen, O., Jerbi, K., Litvak, V., Maess, B., Oostenveld, R., Parkkonen, L., Taylor, J. R., van Wassenhove, V., Wibral, M., & Schoffelen, J.-M. (2013). Good practice for conducting and reporting MEG research. *NeuroImage*, 65, 349–363. <https://doi.org/10.1016/j.neuroimage.2012.10.001>
- Haegens, S., Cousijn, H., Wallis, G., Harrison, P. J., & Nobre, A. C. (2014). Inter- and intra-individual variability in alpha peak frequency. *NeuroImage*, 92, 46–55. <https://doi.org/10.1016/j.neuroimage.2014.01.049>
- Hari, R., & Puce, A. (2017). *MEG-EEG Primer*. Oxford University Press.
- Haufe, S., Dähne, S., & Nikulin, V. V. (2014). Dimensionality reduction for the analysis of brain oscillations. *NeuroImage*, 101, 583–597. <https://doi.org/10.1016/j.neuroimage.2014.06.073>
- Haufe, S., Meinecke, F., Görgen, K., Dähne, S., Haynes, J.-D., Blankertz, B., & Bießmann, F. (2014). On the interpretation of weight vectors of linear models in multivariate neuroimaging. *NeuroImage*, 87, 96–110. <https://doi.org/10.1016/j.neuroimage.2013.10.067>
- Haufe, S., Nikulin, V. V., Müller, K.-R., & Nolte, G. (2013). A critical assessment of connectivity measures for EEG data: A simulation study. *NeuroImage*, 64, 120–133. <https://doi.org/10.1016/j.neuroimage.2012.09.036>
- Hauk, O., Stenroos, M., & Treder, M. (2019). EEG/MEG Source Estimation and Spatial Filtering: The Linear Toolkit. In S. Supek & C. J. Aine (Eds.), *Magnetoencephalography* (pp. 1–37). Springer International Publishing. https://doi.org/10.1007/978-3-319-62657-4_85-1
- He, B. J. (2014). Scale-free brain activity: Past, present, and future. *Trends in Cognitive Sciences*, 18(9), 480–487. <https://doi.org/10.1016/j.tics.2014.04.003>
- He, W., Donoghue, T., Sowman, P. F., Seymour, R. A., Brock, J., Crain, S., Voytek, B., & Hillebrand, A. (2019). *Co-Increasing Neuronal Noise and Beta Power in the Developing Brain* (p. 49) [Preprint]. bioRxiv. <https://doi.org/10.1101/839258>
- Hindriks, R., Micheli, C., Mantini, D., & Deco, G. (2017). *Human Resting-State Electrophysiological Networks In The Alpha Frequency Band: Evidence From Magnetoencephalographic Source Imaging* [Preprint]. bioRxiv. <https://doi.org/10.1101/142091>
- Holdgraf, C., Appelhoff, S., Bickel, S., Bouchard, K., D'Ambrosio, S., David, O., Devinsky, O., Dichter, B., Flinker, A., Foster, B. L., Gorgolewski, K. J., Groen, I., Groppe, D., Gunduz, A., Hamilton, L., Honey, C. J., Jas, M., Knight, R., Lachaux, J.-P., ... Hermes, D. (2019). IIEG-BIDS, extending the Brain Imaging Data Structure specification to human intracranial electrophysiology. *Scientific Data*, 6(1), 102. <https://doi.org/10.1038/s41597-019-0105-7>

- Huang, Y., Parra, L. C., & Haufe, S. (2016). The New York Head—A precise standardized volume conductor model for EEG source localization and tES targeting. *NeuroImage*, 140, 150–162. <https://doi.org/10.1016/j.neuroimage.2015.12.019>
- Hyvärinen, A., & Oja, E. (2000). Independent component analysis: Algorithms and applications. *Neural Networks*, 13(4–5), 411–430. [https://doi.org/10.1016/S0893-6080\(00\)00026-5](https://doi.org/10.1016/S0893-6080(00)00026-5)
- Jas, M., & Dupré, T. (2017). Learning the Morphology of Brain Signals Using Alpha-Stable Convolutional Sparse Coding. *Advances in Neural Information Processing Systems*, 1099–1108.
- Jas, M., Engemann, D. A., Bekhti, Y., Raimondo, F., & Gramfort, A. (2017). Autoreject: Automated artifact rejection for MEG and EEG data. *NeuroImage*, 159, 417–429. <https://doi.org/10.1016/j.neuroimage.2017.06.030>
- Jas, M., Larson, E., Engemann, D. A., Leppäkangas, J., Taulu, S., Hämäläinen, M., & Gramfort, A. (2018). A Reproducible MEG/EEG Group Study With the MNE Software: Recommendations, Quality Assessments, and Good Practices. *Frontiers in Neuroscience*, 12, 530. <https://doi.org/10.3389/fnins.2018.00530>
- Jones, S. R. (2016). When brain rhythms aren't 'rhythmic': Implication for their mechanisms and meaning. *Current Opinion in Neurobiology*, 40, 72–80. <https://doi.org/10.1016/j.conb.2016.06.010>
- Kathawalla, U.-K., Silverstein, P., & Syed, M. (2020). *Easing Into Open Science: A Tutorial for Graduate Students* [Preprint]. PsyArXiv. <https://doi.org/10.31234/osf.io/vzjdp>
- Keil, A., Debener, S., Gratton, G., Junghöfer, M., Kappenman, E. S., Luck, S. J., Luu, P., Miller, G. A., & Yee, C. M. (2014). Committee report: Publication guidelines and recommendations for studies using electroencephalography and magnetoencephalography: Guidelines for EEG and MEG. *Psychophysiology*, 51(1), 1–21. <https://doi.org/10.1111/psyp.12147>
- Kiebel, S. J., Tallon-Baudry, C., & Friston, K. J. (2005). Parametric analysis of oscillatory activity as measured with EEG/MEG. *Human Brain Mapping*, 26(3), 170–177. <https://doi.org/10.1002/hbm.20153>
- 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)
- Koles, Z. J. (1991). The quantitative extraction and topographic mapping of the abnormal components in the clinical EEG. *Electroencephalography and Clinical Neurophysiology*, 79(6), 440–447. [https://doi.org/10.1016/0013-4694\(91\)90163-X](https://doi.org/10.1016/0013-4694(91)90163-X)
- 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>
- Kramer, M. A., Tort, A. B. L., & Kopell, N. J. (2008). Sharp edge artifacts and spurious coupling in EEG frequency comodulation measures. *Journal of Neuroscience Methods*, 170(2), 352–357. <https://doi.org/10.1016/j.jneumeth.2008.01.020>
- Krol, L. R., Pawlitzki, J., Lotte, F., Gramann, K., & Zander, T. O. (2018). SEREEGA: Simulating event-related EEG activity. *Journal of Neuroscience Methods*, 309, 13–24. <https://doi.org/10.1016/j.jneumeth.2018.08.001>
- Lachaux, J.-P., Rodriguez, E., Martinerie, J., Varela, F. J., & others. (1999). Measuring phase

- 1256 synchrony in brain signals. *Human Brain Mapping*, 8(4), 194–208.
 1257 [https://doi.org/10.1002/\(SICI\)1097-0193\(1999\)8:4<194::AID-HBM4>3.0.CO;2-C](https://doi.org/10.1002/(SICI)1097-0193(1999)8:4<194::AID-HBM4>3.0.CO;2-C)
- 1258 Lakatos, P., Karmos, G., Mehta, A. D., Ulbert, I., & Schroeder, C. E. (2008). Entrainment of
 1259 Neuronal Oscillations as a Mechanism of Attentional Selection. *Science*, 320(5872),
 1260 110–113. <https://doi.org/10.1126/science.1154735>
- 1261 Lansbergen, M. M., Arns, M., van Dongen-Boomsma, M., Spronk, D., & Buitelaar, J. K. (2011).
 1262 The increase in theta/beta ratio on resting-state EEG in boys with attention-
 1263 deficit/hyperactivity disorder is mediated by slow alpha peak frequency. *Progress in*
 1264 *Neuro-Psychopharmacology and Biological Psychiatry*, 35(1), 47–52.
 1265 <https://doi.org/10.1016/j.pnpbp.2010.08.004>
- 1266 Lendner, J. D., Helfrich, R. F., Mander, B. A., Romundstad, L., Lin, J. J., Walker, M. P., Larsson,
 1267 P. G., & Knight, R. T. (2020). An electrophysiological marker of arousal level in humans.
 1268 *ELife*, 9, e55092. <https://doi.org/10.7554/eLife.55092>
- 1269 Lepage, K. Q., Kramer, M. A., & Eden, U. T. (2013). Some Sampling Properties of Common
 1270 Phase Estimators. *Neural Computation*, 25(4), 901–921.
 1271 https://doi.org/10.1162/NECO_a_00422
- 1272 Lindsley, D. B. (1939). A Longitudinal Study of the Occipital Alpha Rhythm in Normal Children:
 1273 Frequency and Amplitude Standards. *The Pedagogical Seminary and Journal of Genetic*
 1274 *Psychology*, 55(1), 197–213. <https://doi.org/10.1080/08856559.1939.10533190>
- 1275 Lopes da Silva, F. (2013). EEG and MEG: Relevance to Neuroscience. *Neuron*, 80(5), 1112–
 1276 1128. <https://doi.org/10.1016/j.neuron.2013.10.017>
- 1277 Lotte, F. & Cuntai Guan. (2011). Regularizing Common Spatial Patterns to Improve BCI
 1278 Designs: Unified Theory and New Algorithms. *IEEE Transactions on Biomedical*
 1279 *Engineering*, 58(2), 355–362. <https://doi.org/10.1109/TBME.2010.2082539>
- 1280 Lozano-Soldevilla, D., ter Huurne, N., & Oostenveld, R. (2016). Neuronal Oscillations with Non-
 1281 sinusoidal Morphology Produce Spurious Phase-to-Amplitude Coupling and
 1282 Directionality. *Frontiers in Computational Neuroscience*, 10.
 1283 <https://doi.org/10.3389/fncom.2016.00087>
- 1284 Lundqvist, M., Rose, J., Herman, P., Brincat, S. L., Buschman, T. J., & Miller, E. K. (2016).
 1285 Gamma and Beta Bursts Underlie Working Memory. *Neuron*, 90(1), 152–164.
 1286 <https://doi.org/10.1016/j.neuron.2016.02.028>
- 1287 Mahjoory, K., Nikulin, V. V., Botrel, L., Linkenkaer-Hansen, K., Fato, M. M., & Haufe, S. (2017).
 1288 Consistency of EEG source localization and connectivity estimates. *NeuroImage*, 152,
 1289 590–601. <https://doi.org/10.1016/j.neuroimage.2017.02.076>
- 1290 Mahjoory, K., Schoffelen, J.-M., Keitel, A., & Gross, J. (2020). The frequency gradient of human
 1291 resting-state brain oscillations follows cortical hierarchies. *ELife*, 9, e53715.
 1292 <https://doi.org/10.7554/eLife.53715>
- 1293 Maris, E. (2012). Statistical testing in electrophysiological studies: Statistical testing in
 1294 electrophysiological studies. *Psychophysiology*, 49(4), 549–565.
 1295 <https://doi.org/10.1111/j.1469-8986.2011.01320.x>
- 1296 Maris, E., & Oostenveld, R. (2007). Nonparametric statistical testing of EEG- and MEG-data.
 1297 *Journal of Neuroscience Methods*, 164(1), 177–190.
 1298 <https://doi.org/10.1016/j.jneumeth.2007.03.024>
- 1299 Mazaheri, A., Slagter, H. A., Thut, G., & Foxe, J. J. (2018). Orchestration of brain oscillations:

- Principles and functions. *European Journal of Neuroscience*, 48(7), 2385–2388.
<https://doi.org/10.1111/ejn.14189>
- Miller, E. K., Lundqvist, M., & Bastos, A. M. (2018). Working Memory 2.0. *Neuron*, 100(2), 463–475. <https://doi.org/10.1016/j.neuron.2018.09.023>
- Miller, K. J., Honey, C. J., Hermes, D., Rao, R. P., den Nijs, M., & Ojemann, J. G. (2014). Broadband changes in the cortical surface potential track activation of functionally diverse neuronal populations. *NeuroImage*, 85, 711–720.
<https://doi.org/10.1016/j.neuroimage.2013.08.070>
- Miller, K. J., Sorensen, L. B., Ojemann, J. G., & den Nijs, M. (2009). Power-Law Scaling in the Brain Surface Electric Potential. *PLOS Computational Biology*, 5(12), e1000609.
<https://doi.org/10.1371/journal.pcbi.1000609>
- Mortezapouraghdam, Z., Corona-Strauss, F. I., Takahashi, K., & Strauss, D. J. (2018). Reducing the Effect of Spurious Phase Variations in Neural Oscillatory Signals. *Frontiers in Computational Neuroscience*, 12, 82. <https://doi.org/10.3389/fncom.2018.00082>
- Muthukumaraswamy, S. D., & Singh, K. D. (2011). A cautionary note on the interpretation of phase-locking estimates with concurrent changes in power. *Clinical Neurophysiology*, 122(11), 2324–2325. <https://doi.org/10.1016/j.clinph.2011.04.003>
- Newson, J. J., & Thiagarajan, T. C. (2019). EEG Frequency Bands in Psychiatric Disorders: A Review of Resting State Studies. *Frontiers in Human Neuroscience*, 12.
<https://doi.org/10.3389/fnhum.2018.00521>
- Neymotin, S. A., Barczak, A., O'Connell, M. N., McGinnis, T., Markowitz, N., Espinal, E., Griffith, E., Anwar, H., Dura-Bernal, S., Lytton, W. W., Jones, S. R., Bickel, S., & Lakatos, P. (2020). *Taxonomy of neural oscillation events in primate auditory cortex* [Preprint]. bioRxiv. <https://doi.org/10.1101/2020.04.16.045021>
- Neymotin, S. A., Daniels, D. S., Caldwell, B., McDougal, R. A., Carnevale, N. T., Jas, M., Moore, C. I., Hines, M. L., Härmäläinen, M., & Jones, S. R. (2020). Human Neocortical Neurosolver (HNN), a new software tool for interpreting the cellular and network origin of human MEG/EEG data. *ELife*, 9, e51214. <https://doi.org/10.7554/eLife.51214>
- Nikulin, V. V., Nolte, G., & Curio, G. (2011). A novel method for reliable and fast extraction of neuronal EEG/MEG oscillations on the basis of spatio-spectral decomposition. *NeuroImage*, 55(4), 1528–1535. <https://doi.org/10.1016/j.neuroimage.2011.01.057>
- Niso, G., Gorgolewski, K. J., Bock, E., Brooks, T. L., Flandin, G., Gramfort, A., Henson, R. N., Jas, M., Litvak, V., T. Moreau, J., Oostenveld, R., Schoffelen, J.-M., Tadel, F., Wexler, J., & Baillet, S. (2018). MEG-BIDS, the brain imaging data structure extended to magnetoencephalography. *Scientific Data*, 5(1), 180110.
<https://doi.org/10.1038/sdata.2018.110>
- Nunez, P. L., & Srinivasan, R. (2006). *Electric fields of the brain: The neurophysics of EEG* (2nd ed). Oxford University Press.
- O'Neill, G. C., Tewarie, P., Vidaurre, D., Liuzzi, L., Woolrich, M. W., & Brookes, M. J. (2018). Dynamics of large-scale electrophysiological networks: A technical review. *NeuroImage*, 180, 559–576. <https://doi.org/10.1016/j.neuroimage.2017.10.003>
- Oostenveld, R., Fries, P., Maris, E., & Schoffelen, J.-M. (2011). FieldTrip: Open Source Software for Advanced Analysis of MEG, EEG, and Invasive Electrophysiological Data. *Computational Intelligence and Neuroscience*, 2011, 1–9.

- 1344 <https://doi.org/10.1155/2011/156869>
- 1345 Palva, J. M., Wang, S. H., Palva, S., Zhigalov, A., Monto, S., Brookes, M. J., Schoffelen, J.-M.,
1346 & Jerbi, K. (2018). Ghost interactions in MEG/EEG source space: A note of caution on
1347 inter-areal coupling measures. *NeuroImage*, 173, 632–643.
1348 <https://doi.org/10.1016/j.neuroimage.2018.02.032>
- 1349 Palva, S., & Palva, J. M. (2012). Discovering oscillatory interaction networks with M/EEG:
1350 Challenges and breakthroughs. *Trends in Cognitive Sciences*, 16(4), 219–230.
1351 <https://doi.org/10.1016/j.tics.2012.02.004>
- 1352 Parra, L. C., Spence, C. D., Gerson, A. D., & Sajda, P. (2005). Recipes for the linear analysis of
1353 EEG. *NeuroImage*, 28(2), 326–341. <https://doi.org/10.1016/j.neuroimage.2005.05.032>
- 1354 Pascual-Marqui, R. D., Valdes-Sosa, P. A., & Alvarez-Amador, A. (1988). A Parametric Model
1355 for Multichannel EEG Spectra. *International Journal of Neuroscience*, 40(1–2), 89–99.
1356 <https://doi.org/10.3109/00207458808985730>
- 1357 Pernet, C. R., Appelhoff, S., Gorgolewski, K. J., Flandin, G., Phillips, C., Delorme, A., &
1358 Oostenveld, R. (2019). EEG-BIDS, an extension to the brain imaging data structure for
1359 electroencephalography. *Scientific Data*, 6(1), 103. [https://doi.org/10.1038/s41597-019-](https://doi.org/10.1038/s41597-019-0104-8)
1360 [0104-8](https://doi.org/10.1038/s41597-019-0104-8)
- 1361 Pernet, C. R., Garrido, M. I., Gramfort, A., Maurits, N., Michel, C. M., Pang, E., Salmelin, R.,
1362 Schoffelen, J. M., Valdes-Sosa, P. A., & Puce, A. (2020). Issues and recommendations
1363 from the OHBM COBIDAS MEEG committee for reproducible EEG and MEG research.
1364 *Nature Neuroscience*, 23(12), 1473–1483. <https://doi.org/10.1038/s41593-020-00709-0>
- 1365 Pesaran, B., Vinck, M., Einevoll, G. T., Sirota, A., Fries, P., Siegel, M., Truccolo, W., Schroeder,
1366 C. E., & Srinivasan, R. (2018). Investigating large-scale brain dynamics using field
1367 potential recordings: Analysis and interpretation. *Nature Neuroscience*.
1368 <https://doi.org/10.1038/s41593-018-0171-8>
- 1369 Pfurtscheller, G., & Lopes da Silva, F. H. (1999). Event-related EEG/MEG synchronization and
1370 desynchronization: Basic principles. *Clinical Neurophysiology*, 110(11), 1842–1857.
1371 [https://doi.org/10.1016/S1388-2457\(99\)00141-8](https://doi.org/10.1016/S1388-2457(99)00141-8)
- 1372 Piastra, M. C., Nüßing, A., Vorwerk, J., Clerc, M., Engwer, C., & Wolters, C. H. (2020). A
1373 comprehensive study on electroencephalography and magnetoencephalography
1374 sensitivity to cortical and subcortical sources. *Human Brain Mapping*, hbm.25272.
1375 <https://doi.org/10.1002/hbm.25272>
- 1376 Podvalny, E., Noy, N., Harel, M., Bickel, S., Chechik, G., Schroeder, C. E., Mehta, A. D.,
1377 Tsodyks, M., & Malach, R. (2015). A unifying principle underlying the extracellular field
1378 potential spectral responses in the human cortex. *Journal of Neurophysiology*, 114(1),
1379 505–519. <https://doi.org/10.1152/jn.00943.2014>
- 1380 Popov, T., Oostenveld, R., & Schoffelen, J. M. (2018). FieldTrip Made Easy: An Analysis
1381 Protocol for Group Analysis of the Auditory Steady State Brain Response in Time,
1382 Frequency, and Space. *Frontiers in Neuroscience*, 12, 711.
1383 <https://doi.org/10.3389/fnins.2018.00711>
- 1384 Quinn, A. J., van Ede, F., Brookes, M. J., Heideman, S. G., Nowak, M., Seedat, Z. A., Vidaurre,
1385 D., Zich, C., Nobre, A. C., & Woolrich, M. W. (2019). Unpacking Transient Event
1386 Dynamics in Electrophysiological Power Spectra. *Brain Topography*, 32, 1020–1034.
1387 <https://doi.org/10.1007/s10548-019-00745-5>

- Rey, H. G., Ahmadi, M., & Quiñero, R. (2015). Single trial analysis of field potentials in perception, learning and memory. *Current Opinion in Neurobiology*, 31, 148–155. <https://doi.org/10.1016/j.conb.2014.10.009>
- Robertson, M. M., Furlong, S., Voytek, B., Donoghue, T., Boettger, C. A., & Sheridan, M. A. (2019). EEG Power Spectral Slope differs by ADHD status and stimulant medication exposure in early childhood. *Journal of Neurophysiology*, 122(6), 2427–2437. <https://doi.org/10.1152/jn.00388.2019>
- Sameni, R., & Seraj, E. (2017). A robust statistical framework for instantaneous electroencephalogram phase and frequency estimation and analysis. *Physiological Measurement*, 38(12), 2141–2163. <https://doi.org/10.1088/1361-6579/aa93a1>
- Sanz Leon, P., Knock, S. A., Woodman, M. M., Domide, L., Mersmann, J., McIntosh, A. R., & Jirsa, V. (2013). The Virtual Brain: A simulator of primate brain network dynamics. *Frontiers in Neuroinformatics*, 7. <https://doi.org/10.3389/fninf.2013.00010>
- Schaworonkow, N., Blythe, D. A. J., Kegeles, J., Curio, G., & Nikulin, V. V. (2015). Power-law dynamics in neuronal and behavioral data introduce spurious correlations: Power-Law Dynamics in Neuronal and Behavioral Data. *Human Brain Mapping*, 36(8), 2901–2914. <https://doi.org/10.1002/hbm.22816>
- Schaworonkow, N., & Nikulin, V. V. (2019). Spatial neuronal synchronization and the waveform of oscillations: Implications for EEG and MEG. *PLOS Computational Biology*, 15(5), e1007055. <https://doi.org/10.1371/journal.pcbi.1007055>
- Schoffelen, J.-M., & Gross, J. (2009). Source connectivity analysis with MEG and EEG. *Human Brain Mapping*, 30(6), 1857–1865. <https://doi.org/10.1002/hbm.20745>
- Sherman, M. A., Lee, S., Law, R., Haegens, S., Thorn, C. A., Hamalainen, M. S., Moore, C. I., & Jones, S. R. (2016). Neural mechanisms of transient neocortical beta rhythms: Converging evidence from humans, computational modeling, monkeys, and mice. *Proceedings of the National Academy of Sciences*, 113(33), E4885–E4894. <https://doi.org/10.1073/pnas.1604135113>
- Shin, H., Law, R., Tsutsui, S., Moore, C. I., & Jones, S. R. (2017). The rate of transient beta frequency events predicts behavior across tasks and species. *ELife*, 6, e29086. <https://doi.org/10.7554/eLife.29086>
- Stokes, M., & Spaak, E. (2016). The Importance of Single-Trial Analyses in Cognitive Neuroscience. *Trends in Cognitive Sciences*, 20(7), 483–486. <https://doi.org/10.1016/j.tics.2016.05.008>
- Timmer, J., & König, M. (1995). On Generating Power Law Noise. *Astronomy and Astrophysics*, 300, 707–710.
- Urigüen, J. A., & Garcia-Zapirain, B. (2015). EEG artifact removal—State-of-the-art and guidelines. *Journal of Neural Engineering*, 12(3), 031001. <https://doi.org/10.1088/1741-2560/12/3/031001>
- van Diepen, R. M., & Mazaheri, A. (2018). The Caveats of observing Inter-Trial Phase-Coherence in Cognitive Neuroscience. *Scientific Reports*, 8(1). <https://doi.org/10.1038/s41598-018-20423-z>
- van Ede, F., Quinn, A. J., Woolrich, M. W., & Nobre, A. C. (2018). Neural Oscillations: Sustained Rhythms or Transient Burst-Events? *Trends in Neurosciences*, 41(7), 415–417. <https://doi.org/10.1016/j.tins.2018.04.004>

- 1432 van Vugt, M. K., Sederberg, P. B., & Kahana, M. J. (2007). Comparison of spectral analysis
1433 methods for characterizing brain oscillations. *Journal of Neuroscience Methods*, 162(1–
1434 2), 49–63. <https://doi.org/10.1016/j.jneumeth.2006.12.004>
- 1435 VanRullen, R. (2016). Perceptual Cycles. *Trends in Cognitive Sciences*, 20(10), 723–735.
1436 <https://doi.org/10.1016/j.tics.2016.07.006>
- 1437 Varela, F. J., Lachaux, J.-P., Rodriguez, E., & Martinerie, J. (2001). The brainweb: Phase
1438 synchronization and large-scale integration. *Nature Reviews Neuroscience*, 2(4), 229–
1439 239. <https://doi.org/10.1038/35067550>
- 1440 Vaz, A. P., Yaffe, R. B., Wittig, J. H., Inati, S. K., & Zaghloul, K. A. (2017). Dual origins of
1441 measured phase-amplitude coupling reveal distinct neural mechanisms underlying
1442 episodic memory in the human cortex. *NeuroImage*, 148, 148–159.
1443 <https://doi.org/10.1016/j.neuroimage.2017.01.001>
- 1444 Vidaurre, D., Quinn, A. J., Baker, A. P., Dupret, D., Tejero-Cantero, A., & Woolrich, M. W.
1445 (2016). Spectrally resolved fast transient brain states in electrophysiological data.
1446 *NeuroImage*, 126, 81–95. <https://doi.org/10.1016/j.neuroimage.2015.11.047>
- 1447 Voytek, B. (2016). The Virtuous Cycle of a Data Ecosystem. *PLOS Computational Biology*,
1448 12(8), e1005037. <https://doi.org/10.1371/journal.pcbi.1005037>
- 1449 Voytek, B., & Knight, R. T. (2015). Dynamic Network Communication as a Unifying Neural Basis
1450 for Cognition, Development, Aging, and Disease. *Biological Psychiatry*, 77(12), 1089–
1451 1097. <https://doi.org/10.1016/j.biopsych.2015.04.016>
- 1452 Voytek, B., Kramer, M. A., Case, J., Lepage, K. Q., Tempesta, Z. R., Knight, R. T., & Gazzaley,
1453 A. (2015). Age-Related Changes in 1/f Neural Electrophysiological Noise. *Journal of*
1454 *Neuroscience*, 35(38), 13257–13265. <https://doi.org/10.1523/JNEUROSCI.2332-14.2015>
- 1455 Wacker, M., & Witte, H. (2013). Time-frequency Techniques in Biomedical Signal Analysis: A
1456 Tutorial Review of Similarities and Differences. *Methods of Information in Medicine*,
1457 52(04), 279–296. <https://doi.org/10.3414/ME12-01-0083>
- 1458 Wang, X.-J. (2010). Neurophysiological and Computational Principles of Cortical Rhythms in
1459 Cognition. *Physiological Reviews*, 90(3), 1195–1268.
1460 <https://doi.org/10.1152/physrev.00035.2008>
- 1461 Waschke, L., Donoghue, T., Fiedler, L., Smith, S., Voytek, B., & Obleser, J. (2021). *Modality-*
1462 *specific tracking of attention and sensory statistics in the human electrophysiological*
1463 *spectral exponent* [Preprint]. bioRxiv. <https://doi.org/10.1101/2021.01.13.426522>
- 1464 Watrous, A. J., & Buchanan, R. J. (2020). The Oscillatory ReConstruction Algorithm adaptively
1465 identifies frequency bands to improve spectral decomposition in human and rodent
1466 neural recordings. *Journal of Neurophysiology*, 124(6), 1914–1922.
1467 <https://doi.org/10.1152/jn.00292.2020>
- 1468 Watrous, A. J., Miller, J., Qasim, S. E., Fried, I., & Jacobs, J. (2018). Phase-tuned neuronal
1469 firing encodes human contextual representations for navigational goals. *ELife*, 7,
1470 e32554. <https://doi.org/10.7554/eLife.32554>
- 1471 Wen, H., & Liu, Z. (2016). Separating Fractal and Oscillatory Components in the Power
1472 Spectrum of Neurophysiological Signal. *Brain Topography*, 29(1), 13–26.
1473 <https://doi.org/10.1007/s10548-015-0448-0>
- 1474 Wessel, J. R. (2020). β -Bursts Reveal the Trial-to-Trial Dynamics of Movement Initiation and
1475 Cancellation. *The Journal of Neuroscience*, 40(2), 411–423.

- 1476 <https://doi.org/10.1523/JNEUROSCI.1887-19.2019>
- 1477 Whitten, T. A., Hughes, A. M., Dickson, C. T., & Caplan, J. B. (2011). A better oscillation
1478 detection method robustly extracts EEG rhythms across brain state changes: The
1479 human alpha rhythm as a test case. *NeuroImage*, 54(2), 860–874.
1480 <https://doi.org/10.1016/j.neuroimage.2010.08.064>
- 1481 Widmann, A., Schröger, E., & Maess, B. (2015). Digital filter design for electrophysiological data
1482 – a practical approach. *Journal of Neuroscience Methods*, 250, 34–46.
1483 <https://doi.org/10.1016/j.jneumeth.2014.08.002>
- 1484 Wilson, G., Bryan, J., Cranston, K., Kitzes, J., Nederbragt, L., & Teal, T. K. (2017). Good
1485 enough practices in scientific computing. *PLOS Computational Biology*, 13(6),
1486 e1005510. <https://doi.org/10.1371/journal.pcbi.1005510>
- 1487 Wutz, A., Melcher, D., & Samaha, J. (2018). Frequency modulation of neural oscillations
1488 according to visual task demands. *Proceedings of the National Academy of Sciences*,
1489 115(6), 1346–1351. <https://doi.org/10.1073/pnas.1713318115>
- 1490 Zich, C., Quinn, A. J., Mardell, L. C., Ward, N. S., & Bestmann, S. (2020). Dissecting Transient
1491 Burst Events. *Trends in Cognitive Sciences*, 24(10), 784–788.
1492 <https://doi.org/10.1016/j.tics.2020.07.004>
- 1493