Title: **Neural harmonics of syntactic structure**

Authors: Alessandro Tavano[1*], Stefan Blohm[2], Christine A. Knoop[2], R

Muralikrishnan[1], Mathias Scharinger[2,3], Valentin Wagner[2], Dominik Thiele[1], Oded

Ghitza[4], Nai Ding[5], Winfried Menninghaus[2], David Poeppel[1,6]

Affiliations:

1. Department of Neuroscience, Max Plank Institute for Empirical Aesthetics,

60322 - Frankfurt am Main, Germany

2. Department of Language and Literature, Max Plank Institute for Empirical

Aesthetics, 60322 - Frankfurt am Main, Germany

3. University of Marburg, 35037 – Marburg, Germany

4. Department of Biomedical Engineering, Boston University, 02215 - Boston,

USA

5. College of Biomedical Engineering and Instrument Sciences, Zhejiang

University, 310027 - Hangzhou, Zhejiang Province, China

6. Department of Psychology, New York University, 10003 - New York City, New

York, USA

* Corresponding author: alessandro.tavano@ae.mpg.de, david.poeppel@ae.mpg.de

Abstract

Can neural rhythms reflect purely internal syntactic processes in multi-word

constructions? To test this controversial conjecture - relevant to language in

particular and cognition more broadly - we recorded electroencephalographic and

behavioural data as participants listened to isochronously presented sentences of

varying in syntactic complexity. Each trial comprised ten concatenated sentences

and was either fully grammatical (regular) or rendered ungrammatical via randomly

distributed word order violations. We found that attending the regular repetition of

abstract syntactic categories (phrases and sentences) generates neural rhythms

whose harmonics are mathematically independent of word rate. This permits to

clearly separate endogenous syntactic rhythms from exogenous speech rhythms.

We demonstrate that endogenous but not exogenous rhythms predict participants'

grammaticality judgements, and allow for the neural decoding of regular vs. irregular

trials. Neural harmonic series constitute a new form of behaviourally relevant

evidence for syntactic competence.

Introduction

Speech and language are related but distinct dimensions of verbal communication [1–4]. In a nutshell, speech is linear, while language is hierarchically organized. Temporal regularities linearly guide speech chunking in ways that are relevant for language production and comprehension [3, 5–13]): for example, entrainment to syllabic rhythm is relevant for word formation processes [3,6,14]. However, the core of language processing lies in the internal and tacit knowledge of syntactic rules, which determine how words combine into meaningful utterances [15]. The challenge is then to extract syntactic hierarchy from neural responses to speech acoustics. Valuable information has been inferred about the human brain's sensitivity to syntactic rules by measuring event-related potentials responses (ERPs) to their violation [16-17]. However, ERPs reflect a mixture of abstract functional and physical properties of each individual word [18-19], making it difficult to capture a signature of syntactic competence spanning over multiple words. To go the extra mile, a link can be established between the duration of multi-word syntactic units, whose extraction presupposes competence, and slow brain rhythms. Ding and colleagues [20] made such connection by asking native speakers of English and Mandarin Chinese to listen to trials of ten spoken sentences composed of monosyllabic words presented at a fixed rate of 4 Hz. Forcing word-to-word isochrony into sentence perception allows to 'frequency tag' [21] the duration of abstract syntactic units – phrases and sentences – which regularly repeat albeit with ever differing words. Ding and colleagues [20] tested one syntactic structure: Two-word noun phrases (NP = a phrase headed by a noun, such as "My shoe") followed by two-word verb phrases (VP = a phrase headed by a verb, such as "is wet", see Figure 1a). A Fast Fourier Transform (FFT) analysis of chains of syntactically identical sentences composed of

different words highlighted significant peaks of spectral energy not only at 4 Hz, the

stimulation or word frequency, but also at 2 Hz and 1 Hz, which the authors

interpreted as tracking the period of phrases (2-word units = 2 Hz) and sentences (4-

word units = 1 Hz). If this were the case, an isomorphic mapping between neural

rhythms and levels of syntactic structure would have been established. Of note,

linguistic rhythms emerged only if participants listened to their own language [22].

Could the hierarchy of rhythms reflect the online incremental combination of words

into phrases, and phrases into sentences [23]?

On reflection, the fact that the human brain readily distinguishes between NPs and

VPs [24–29] weakens the evidence for brain rhythms-syntax isomorphism. In Ding et

al.'s [20] experimental setting, each of the phrasal nodes repeats once every second

(1 Hz), and not twice a second, making it impossible for phrasal nodes to

independently generate a 2 Hz rhythm. Instead, since phrasal and sentence units

repeat with the same period of 1unit/sec, they must jointly contribute to the 1 Hz

rhythm. We term this the *Categorical* rhythm, as it reflects the repetition of abstract

syntactic categories, phrases and sentences. It follows that the 2 Hz rhythm would

simply be its first harmonic process (see Figures 1a,b,c), effectively barring any

direct, one-to-one mapping from linguistic to brain hierarchy.

An independent phrasal rhythm could still emerge if and only if there exited a

superordinate phrase category, which we term *sPhrase*, that would cycle through all

NPs and VPs, blind to their lexical head type (see Figures 1d,e). In the example of

Ding et al. [20], this would generate an independent 2 Hz rhythm, and bring the

hypothesis of isomorphism back on the table. The idea that phrase structure may be

independent of lexical categories has exerted a large influence in mainstream

syntactic theory [30], and although rebutted [31], it has never been tested from the

view-point of brain rhythms.

We set out to compare the categorical harmonics and sPhrase models by

systematically varying the size of phrasal units in six different sentence structures in

the German language. We recorded encephalographic (EEG) and behavioural data

while 31 participants discriminated regular from irregular trials composed of ten

isochronized sentences. Regular trials (50%) contained only grammatical sentences,

while for irregular trials two randomly chosen sentences were rendered

ungrammatical by violating word order (random word distribution).

We derived inter-trial phase coherence estimates for each sentence type from the

FFT analysis. Results do not lend support to the existence of a superordinate

sPhrase category, and question attempts to directly map brain hierarchy to syntactic

hierarchy within a frequency tagging approach. Rather, attending the regular

repetition of abstract linguistic categories – NPs and VPs together with the sentence

node – generates a single categorical rhythm with its own harmonic structure. Neural

harmonics of categorical rhythms are mathematically independent from word rate,

making it possible to conclusively dissociate entrainment to word rate from top-down

rhythms reflecting the repetition of multi-word syntactic categories. Neural responses

to categorical harmonics significantly predict participants grammaticality judgements,

and – in contrast to word rate – successfully classify grammatically regular from

irregular trials.

Figure 1. Stimulus structure and neural response models: a. A syntactic description

of a sample sentence structure which, when presented in chains of ten sentences

with the same structure but realized using different lexical items, would permit

distinguishing harmonic peaks from sPhrase peaks in the frequency domain. Three

levels of linguistic investigation are highlighted: word (stimulus rate), phrase and

sentence. b. For five-word sentences running at 4 Hz, the categorical model

assumes that NPs and VPs repeat every 0.8 seconds, on a pair with sentences. c.

Consequently, an FFT analysis of neural data should highlight – beside a word rate

peak – a 0.8 Hz peak of energy, representing the categorical rhythm (Cat), and its

harmonics (first harmonic – H1 – 1.6 Hz, second harmonic – H2 – 2.4 Hz). d. The

sPhrase model would suggest instead the existence of phrasal tracking blind to their

lexical head, and therefore yielding, in the example at stake, a 2 Hz rhythm (for the

two-word NP) and 1.33 Hz rhythm (for the three-word VP). e. The neural responses

of sentences and phrases would be independent.

Results

Grammatical judgments. Trials (ten concatenated sentences) were organized into

three contexts: 1) fast four-word sentences, with monosyllabic words running at 4

Hz; 2) fast five-word sentences, with monosyllabic words running at 4 Hz; 3) slow

five-word sentences, built from disyllabic words running at 2 Hz. Within each context,

two sentence structures were created by varying the relative duration of noun and

verb phrases, in order to test their import on neural responses (see Figure 2).

Fast Four words

Fast Five words

Slow Five words

Figure 2. Sentence contexts: Sentence structures 1 and 2 have four monosyllabic

words running at a 4 Hz, with structure 1 composed of a two-word NP and a two-

word VP - replicating the stimuli of Ding et al. [20] -, and structure 2 of one-word NP

followed by a three-word VP. Structures 3 and 4 have five monosyllabic words

running at 4 Hz: Structure 3 sentences are composed of a three-word NP followed

by a two-word VP, while structure 4 sentences flips that pattern. Structures 5 and 6

replicate structures 3 and 4 with disyllabic words running at 2 Hz. Syntactic trees of

exemplary sentences and corresponding waveforms are provided for illustrative

purposes. Black and grey colours identify even and odd sentence structures within

each context (1-2, 3-4, 5-6, respectively). Notice that although current theories see

determiners as heads of the determiner phrases (DP), with NPs as complements, for

our purposes we follow a more traditional labelling convention. See Supporting

Information for a complete list of sentences (N = 300).

Using a signal detection approach [32], we estimated task sensitivity and response

bias for each sentence structure. The discriminability index (d') measures task

sensitivity: For all sentence structures, participants successfully distinguished

irregular from regular trials, suggesting that the grammatical nature of the task was

well understood: all $ts_{(30)} \geq 7.94$, all $ps \leq 7.205 \times 10^{-09}$, FDR-corrected (threshold p =

0.008. See Figure 3, left panel). There was a significant difference among sentence

structures: $F_{(5,185)} = 4.16$, p = 0.001, $\eta^2 = 0.04$. When FDR-corrected (threshold p =

0.006), type 1 differed from type 4, $t_{(30)} = -3.34$, p = 0.001, suggesting that sensitivity

for structure 1 (mean = 1.83, SD = 1.24) was lower than for structure 4 (mean =

2.67, SD = 1.75). Response bias, measured by the decision criterion, showed that

participants tended to deem irregular trials as regular for sentence types 1, 4, 5 and

6: all $ts_{(30)} \geq 3.65$, all $ps \leq 9.749*10^{-04}$, FDR-corrected (threshold p = 0.021. See

Figure 3, right panel). This suggests a conservative bias in grammaticality judgment.

From this, we can infer that that the impoverished acoustic quality of stimuli following

the pitch-flattening procedure did not confound participants' comprehension. No

significant difference in criterion estimates among sentence structures was found,

suggesting that the stimulus generation procedure was bias-free: $F_{(5,185)} = 1.81$, p =

0.115.

Figure 3, Behavioural accuracy. Upper panel: Participants discriminated between

regular and irregular trials for all six sentence structures (middle bar, color-coded). A

significant difference between structures 1 and 4 suggests that discriminating

irregular from regular trials in structure 1, the original condition of Ding et al. [20],

was more difficult than in structure 4. Middle panel: sentence structure label. Lower

panel: A positive bias for structures 1, 4, 5 and 6 suggests that participants adopted

a conservative criterion when operating grammatical judgments: They were more

likely to respond that a given trial was regular, rather than irregular.

Controlling for speech and acoustic cues. Phonemic transitions between successive

words could implicitly cue phrasal segmentation. To verify whether this was the case,

we calculated phonemic transition probabilities at critical boundaries within and

between phrases in all experimental sentences, and found no significant differences

(See Supporting Information, Figure S1a,b). Word frequency differed across

sentence structures (Figure S1c), as expected due to the asymmetry in the

distribution of monosyllabic and disyllabic words in natural languages (for the

German language, see [33]).

Acoustic waveforms were subject to a Fast Fourier Transform analysis on a trial-by-

trial basis. We extracted normalized power estimates using the complex modulus,

and then calculated signal-to-noise ratio estimates (SNR). Results suggest the

presence of a significant word rate effect for all sentence structures: all $ts_{(30)} \geq 55.04$,

all $ps \leq 1.080*10^{-31}$, FDR-corrected (threshold $p = 0.001$, Figure S2a,b,c). No other

significant rhythmic components were found, suggesting that acoustic input could not

drive higher-order, syntactic rhythms (see Supporting Information).

Categorical harmonics

For neural data, we calculated inter-trial phase coherence estimates as the ratio of

the FFT output and the amplitude of the signal, normalized by the number of trials for

each sentence type and participant (See Materials and Methods). Neural frequency

tagging effects were investigated using all trials, regardless of grammaticality. Figure

4, left column, illustrates phase coherence peaks for each sentence structure.

Statistical analyses were run on standardized phase coherence signal-to-noise ration

(SNR) scores. T-tests to zero showed robust entrainment to word rate for all

sentence structures: all $ts_{(30)} \geq 9.21$, all ps $\leq 1*10^{-08}$, FDR-corrected (threshold p =

0.008), and no significant difference between sentence structures within each

context: all $ts_{(30)} \leq 0.27$, all ps ≥ 0.788. Similarly, a significant harmonic series, here

defined by the emergence of a categorical rhythm and its first two harmonics, H1 and

H2, was found for all sentence structures: Category, all $ts_{(30)} \geq 3.61$, all ps ≤ 0.001;

H1, all $ts_{(30)} \geq 3.28$, all ps ≤ 0.002; H2, all $ts_{(30)} \geq 3.09$, all ps ≤ 0.004 (all FDR-

corrected, threshold p = 0.008; See Figure 4, middle column). Hence, differences in

the internal phrasal organization of sentences did not influence the

presence/absence of a harmonic series. Crucially, the duration of abstract categories

in sentence structures 3, 4, 5 and 6 ensures that first and second harmonic

processes cannot be mathematically derived as subharmonics of stimulus rate [34]:

for example, for sentence types 3 and 4 there exists no integer factor which

multiplied by the first harmonic at 1.6 Hz would yield the 4 Hz word rate. The same

holds for sentence types 5 and 6. This simple fact grants functional independence of

each harmonic series relative to exogenous word rhythm, and proves the existence

of purely endogenous, top-down rhythms.

Syntactic variability within each sentence context modulated the strength of

harmonic peaks. For the fast four-word context, a main effect of peak strength was

found: $F_{(2,185)} = 4.83$, p = 0.01, $\eta^2 = 0.04$. H2 differed from both categorical rhythm

and H1: all $ts_{(30)} \geq 2.75$, all ps ≤ 0.01. The responses for categorical rhythm (mean =

0.47 standardized SNR, SD = 0.35) and H1 (mean = 0.44, SD = 0.29) were larger

than for H2 (mean = 0.27, SD = 0.25). In the fast five-word context, a significant peak

by sentence structure interaction ($F_{(2,185)}$ = 5.06, p = 0.009, η^2 = 0.03) highlighted a

difference in H1 responses between sentence structures 3 and 4: $t_{(30)}$ = -4.08, p =

$3.618*10^{-05}$. H1 in structure 3 (mean = 0.22, SD = 0.33) was smaller than in structure

4 (mean = 0.63, SD = 0.45). In the slow five-word context, a main effect of sentence

structure was found: $F_{(1,185)}$ = 14.16, p = 0.007, η^2 = 0.05. Neural responses were

larger in structure 5 (mean = 0.48, SD = 0.07) than in structure 6 (mean = 0.26, SD =

0.04).

Figure 4, Neural rhythms. Left column shows phase coherence results for all trials,

for each sentence context (sentence structure pairs: 1-2, 3-4, 5-6). Categorical

rhythms and their harmonics, along with the positional distribution of possible

sPhrase rhythms, are color-coded and indicated by vertical hyphenated bars. No

significant peaks were found for any of the hypothesized sPhrase rhythms.

Conversely, evidence for a categorical harmonics model was found for all sentence

structures. The central column confirms the difference between categorical and

sPhrase frequencies in all sentence contexts, using standardised SNR estimates. All

probability values, here and in the text, are deemed significant or not within an FDR

approach. Significance is marked f as follows: *** $\leq$ 0.0001, ** $\leq$ 0.001, ns = non-

significant. Horizontal bars on right column figures represent 95% posterior odds

credible intervals.

Model comparison

Sentence structure 1 is based on the original stimuli of Ding et al. [20].

Consequently, sPhrase and H1 coincide, making sentence structure 1 uninformative

in adjudicating between categorical harmonics and sPhrase models. In sentence

structure 2, the sPhrase model would expect a 1.33 Hz neural rhythm stemming from

3-word VPs (the 4 Hz rhythm stemming from the one-word NP is confounded with

the stimulation frequency; Figure 4a, left column). Structures 3 to 6 allow for a full

test of the internal generation of neural rhythms, as each phrase should create its

own sPhrase peak (Figures 4b and 4c, left column; See also Figure 1).

T-tests to zero showed that no sPhrase peak reached significance: all $ts_{(30)} \leq 0.71$,

all ps $\geq$ 0.062. We then averaged SNR values across sentence structures (Figure 4,

middle column), and run a series of one-way rmANOVAs, one for each sentence

context, with peak frequency as a fixed factor. In all cases, categorical harmonic

rhythms differed from sPhrase rhythms: all $Fs_{(3,123)} \geq 6.76$, all ps $\leq$ 0.001, all $\eta^2 \geq$

0.13. Post-hoc comparisons showed that categorical harmonic structures

outperformed sPhrase rhythms: all $ts_{(30)} \geq 2.73$, all ps $\leq$ 0.01, FDR-corrected

(threshold $\geq$ 0.016).

We evaluated the amount of evidence provided by phase coherence data using a

Bayesian rmANOVA. The Bayes factor indicates that the data – across sentence

contexts – were 156.786 times more likely under the model that includes peak

frequency as a fixed factor, compared to the null model, which assumes no

difference between peaks. Post-hoc comparisons highlight the difference between

sPhrase and categorical harmonic peaks in the distribution of model-averaged

posterior odds (normalized to the mean of all data). The sPhrase peak vs. H2 in the

fast, four-word context revealed posterior odds of 1.782 against the null hypothesis,

which indicates moderate evidence in favour of the alternative hypothesis ($BF_{10} =$

4.301). For all remaining comparisons in the fast four-word context, as well as in the

remaining sentence contexts, the evidence in favour of the alternative hypothesis

was very strong: all $BF_{10} \geq 47.309$ (Figure 4, right column).

Grammaticality in the frequency domain

We have determined that the brain encodes the duration of regularly repeated

abstract syntactic nodes – NP, VP, sentence – within a harmonic structure of their

period. However, the emergence of significant peaks of neural activity does not imply

that they encode perceived grammaticality. To test this, we re-calculated SNR

estimates of inter-trial phase coherence separately for hits (correctly identified

irregular trials) and correct rejections (correctly identified regular trials). Omnibus

rmANOVAs with factors frequency peak, sentence structure and grammaticality,

separately for each sentence context, highlighted a significant grammaticality effect:

all $Fs_{(1,371)} \geq 9.50$, $p \leq 0.01$, $\eta^2 = 0.02$. In all cases, phase coherence for grammatical

trials was larger than for trials containing ungrammatical sentences: fast four words,

grammatical mean = 0.23, SD = 0.16, ungrammatical mean = 0.11, SD = 0.16; fast

five words, grammatical mean = 0.35, SD = 0.23, ungrammatical mean = 0.21, SD =

0.16; slow five words, grammatical mean = 0.26, SD = 0.14, ungrammatical mean =

0.24, SD = 0.13 (see Figure 5a). For the fast five words context, a significant

difference between types was also found, mirroring model comparison results: $F_{(1,371)}$

= 14.53, p = 0.0006, η^2 = 0.04. No other significant main effect or interaction was

found: all $Fs_{(1,371)} \leq$ 2.60, p $\geq$ 0.08.

Using a Bayesian rmANOVA with factors sentence context and grammaticality

(Figure 5b, left panel), the Bayes factor indicates that the data were at least

$1.185*10^{07}$ times more likely under the model that includes grammaticality and

sentence context as main factors: updated probability = 0.89, prior null model

probability = 0.20. The evidence for the interaction component was very small, as

adding it changed prior odds only marginally - BF_M = 0.433 -, as compared to simply

adding main effects: BF_M = 32.352. Post-hoc comparisons of grammatical vs.

ungrammatical trials revealed a BF_{10} of 40847.319 against the null hypothesis, which

indicates decisive evidence in favour of the alternative hypothesis. As for the context

factor, decisive evidence was obtained for the difference between fast five- and four-

word sentence contexts (BF_{10} = 394.811, uncorrected for multiple comparisons), and

strong evidence for the difference between fast and slow five-word sentences (BF_{10}

= 17.039, uncorrected). We conclude that the strength of categorical harmonic

structures, resulting from attending to the repetition of abstract syntactic categories,

is modulated by trial grammaticality.

Figure 5, Neural rhythms of grammaticality: a. An omnibus rmANOVA analysis on

averaged endogenous rhythms shows a significant effect of grammaticality in each

sentence context. b. A Bayesian rmANOVA highlights two main effects: sentence

context and grammaticality. The grammaticality effect provides decisive evidence for

endogenously-driven phase coherence. The sentence context effect provides

decisive evidence suggesting that the fast five-word context generates the largest

phase coherence scores. Significance is marked as follows: *** $\leq$ 0.0001, ** $\leq$

0.001, ns = non-significant. Horizontal bars on right column figures represent 95%

posterior odds credible intervals.

Harmonics encode grammaticality judgements

When participants decide whether a trial contained ungrammatical sentences or not

(irregular trial), their judgment must be driven by implicit syntactic competence of the

German language. However, violations of expected word order also represent a form

of sensory surprise, which per se can capture attention [35] and independently

predict behavioural accuracy. To partition these two contributions, we created two

simple neural predictors: an endogenous one, by calculating for each participant the

mean standardized phase coherence score across categorical harmonic peaks (Cat,

H1 and H2), for all sentence structures, separately for correct rejections and hits;

and an exogenous one, by averaging the strength to which each participant

entrained to external word rate, again across sentence structures and separately for

regular and irregular trials. Each neural predictor was then regressed onto task

sensitivity scores, averaged across all sentence structures. A functionally relevant

dissociation emerged between endogenous and exogenous neural rhythms

extracted from regular vs. irregular trials. For fully grammatical, regular trials, the

endogenous index significantly predicted participants' ability to decide whether a trial

was regular or irregular: $F_{(1,29)} = 15.93$, $p = 4.096*10^{-04}$, $adjR^2 = 0.33$. However, the

exogenous index did not: $F_{(1,29)} = 0.70$, $p = 0.407$ (see Figure 6a). This suggests that

only endogenous categorical harmonics reflect access to internal models of syntactic

knowledge. For irregular trials, however, both indexes significantly predicted

participants' ability to decide whether a trial was regular or irregular: endogenous,

$F_{(1,29)} = 23.15$, $p = 4.277*10^{-05}$, $adjR^2 = 0.42$; exogenous, $F_{(1,29)} = 13.68$, $p = 0.0011$,

$adjR^2 = 0.28$ (see Figure 6b). To decide whether the results for exogenous rhythms

in irregular trials reflect sensory surprise or grammatical violation detection, we

resorted to a neural decoding approach. We calculated standardized phase

coherence scores separately for each relevant frequency peak – categorical, H1, H2

and word rate –, and used a linear discriminant analysis (LDA) with electrodes as a

feature dimension to classify hits and correct rejection trials from neural rhythms.

The performance of the classifier – scored by computing the nonparametric measure

of the area under the curve (AUC) of the receiver-operating characteristic – was

significant for all rhythms belonging to categorical harmonic series: Categorical

rhythm = .58, CI: 0.47 – 0.53; H1 = .64, CI: 0.46 – 0.52; H2 = .65, CI: 0.46-0.53.

Crucially, however, it was at chance for entrainment to word rate: .50, CI: 0.47-0.53

(see Figure 6c). Hence, we conclude that only top-down rhythms, constituting a

harmonic series of syntactic node repetition period, encode information about a trial

grammaticality.

Figure 6, Brain/behaviour fit: a. Accuracy fit of endogenous and exogenous rhythms

extracted from regular trials. Only endogenous rhythms predict behavioural

accuracy. Thick lines represent best fit. Thin, dotted lines indicate each regression's

95% confidence interval. b. When extracted from irregular trials, both endogenous

and exogenous rhythms significantly predict grammaticality judgements, suggesting

the contribution of both syntactic competence and sensory surprise to neural activity.

c. However, AUC scores from an LDA analysis show that it is possible to correctly

classify regular vs. irregular trials using endogenous rhythms, but not entrainment to word rate. Shaded color-coded areas report 95% confidence intervals, obtained via Montecarlo resampling. Significance: *** ≤ 0.001, * ≤ 0.05, ns = non-significant.

Discussion

We aimed at extracting a neural signature of purely internal syntactic information from multi-word speech units. Since speech acoustic and linguistic information are intertwined in verbal communication, to zoom in on syntactic competence we reduced the perceptual significance of speech cues to comprehension by forcing words to onset at regular intervals. This experimental strategy may appear artificial, because intuitively we associate the naturalness of language to the naturalness of speech. However, once the difference between the hierarchical nature of syntax vs. the linear nature of speech is highlighted, it becomes evident that isochronizing speech leaves the core of language competence, and hence its naturalness, intact. That intuition founded Ding et al.'s [20] milestone attempt to go beyond single-word neural responses (e.g., ERPs), and measure neural rhythms to multi-word language units. We provide evidence that frequency tagging goes deep in that it captures internal syntactic processes relevant for grammaticality judgements. However, this result does not rest on an isomorphic mapping between the levels of syntactic analysis (word, phrase, sentence) and the hierarchy of neural rhythms (word rhythm, phrase rhythm, sentence rhythm). Frequency tagging does not allow to separately read out sentences and phrases, because their abstract syntactic nodes are repeated with the very same period (for a congruent lexical approach, see [36]). The repetition of linguistic categories – both phrases and sentences – with the same period generates a categorical neural rhythm, which gives rise to a harmonic series

that encodes grammaticality judgements. Thus, harmonic series represent an

indirect mapping between syntactic competence and neural rhythms.

Importantly, our experimental manipulations document that the harmonic series

generated by category-driven syntactic rhythms are independent of entrainment to

word rate. For example, in conditions 3 and 4, the first and second harmonics at 1.6

and 2.4 Hz cannot be mathematically derived as sub-harmonic processes of the 4

Hz word rate. This strongly supports a principled distinction between endogenous

rhythms, which follow abstract node repetitions and encode grammaticality, and

exogenous rhythms, which entrain to word rate and likely encode sensory surprise,

when extracted from irregular trials (See Figure 6c).

Notice that categorical rhythms and their harmonics emerge regardless of internal

sentence structure, although the latter modulates the strength of the harmonics,

possibly highlighting differences in the ease of syntactic processing.

The strength of harmonic series predicts the degree to which native German

speakers are sensitive to word order violations, i.e., the grammaticality manipulation

we selected for our experiment. The online comprehension of German relies

substantially on word order within and across phrases [37]. However, within the

Germanic language family, English, sensibly, assigns word order more weight for

comprehension than German does [38]. We therefore predict that the sensitivity to

word order violations, as reflected by the strength of neural harmonics, should be

larger for languages which strongly depend on word order for comprehension, such

as English, and smaller for morphologically richer languages, such as Italian, in

which comprehension is locally bound by rules of declension. This perspective opens

new, exciting avenues of cross-linguistic testing of different types of grammaticality

constraints in multilanguage acquisition settings, as well as for language recovery in

clinical settings.

By correctly characterizing the correspondence between linguistic functions and

brain data, we contribute to linguistic theory by proving no evidence for the existence

of a superordinate phrasal category – *sPhrase* – cycling through all phrases

regardless of their lexical head can be posited. A proposal along these lines was first

put forward by Chomsky [30], following the observation of similarities in the internal

structure of NPs and VPs (see also [39-40]). In this respect, a frequency tagging

approach to abstract multi-word categories has the potential to contribute directly to

linguistic theories. However, there are also limitations to our experimental setup and

analysis. First, FFT-base analyses of repeated sentence structures convey

information on the *result* of top-down comprehension processes – that is, whether

syntactic representations have been accessed from long-term memory – , but cannot

provide any evidence for the incremental bottom-up processes combining words into

phrases, and phrases into sentences (for a different take, see [23]). More generally,

endogenous rhythms can in principle simply reflect the outcome of a decoding

process that originates when participants encounter the first few trials, and is just

top-down replicated for the rest of the trials. Second, endogenous neural rhythms

generated via frequency tagging are uninformative relative to the temporal dynamics

of internal representations in continuous speech perception. Third, the distinction

between linear speech perception and hierarchical language comprehension sits well

with our stimuli, as we use of monosyllabic and simple disyllabic units, but must not

be overstated, as in many languages morphological processes contributing to word

formation also have a hierarchical nature.

The delta band (0.5-4 Hz) and infra-slow (< 0.5 Hz) rhythms we measured lie within

natural spectral niches for linguistic operations because their extended time windows

match the ranges of spontaneous language segment duration [41- 43]. However,

delta rhythms also robustly entrain to salient events in continuous speech [44-45], as

well as to meaningless word lists [22]. Frequency tagging offers a way to partition the

contribution of language knowledge from that of speech, and is easily applicable to

the study of top-down knowledge effects in other domains, such as music

perception. An interesting point for future research pertains to the type of top-down

knowledge that frequency tagging can tap into, from capturing complex syntactic

structures (combining sentences within sentences) to detecting degrees of rule

violations as a function of neural response strength.

In conclusion, we have shown that internally constructed neural rhythms and their

harmonic series reflect the top-down application of syntactic rules determining the

temporal boundaries of phrases and sentences, independently of entrainment to

word rate.

Materials and methods

Participants. Thirty-two young adults participated in the Experiment (Age range: 19-

30, 8 males). Sample size was determined using values derived from the inspection

of Figure 1c in Ding et al. [20]: N = 16, mean response across peaks ± 12 dBs (SD ±

8 dBs), effects size = 1.5, power = 0.99. We hypothesized a reduction of 50% in

effect size due to EEG volume conduction, leading to a required sample size of 30

with alpha = 0.05. The EEG data from one participant were not faithfully recorded,

and were thus discarded. All analyses were run on the remaining participants (N =

31): they were all German native speakers, right-handed (Oldfield Test), and self-

reported normal hearing, normal-to-corrected vision, no medical history of treatments

affecting the central nervous system, and no psychiatric disturbances. They were

compensated with 10 euros per hour (~3 hours, including instruction, EEG montage

and cleaning). All experimental procedures were approved by the Ethics Committee

of the Max Planck Society, and were undertaken with a written informed consent

signed by each participant.

Stimuli. To reduce implicit word- and phrase-level prosodic cues, word stimuli were

individually generated using the MacinTalk Text-to-Speech Synthesizer (OSX El

Capitan, voice Anna, f0 set at 220 Hz), and then pitch-flattened to 220 Hz across all

time points (fundamental frequency estimated using an autocorrelation approach,

Praat, v. 6.0.16, www.praat.org). If an original stimulus was longer than the relative

SOA, its duration was compressed to fit the stimulation window while preserving

pitch.

Individual trials were composed of ten sentences, identical from the viewpoint of

main phrasal units, but differing in the types of lexical heads and internal composition

of phrases. Sentences were randomly selected from six predefined lists of 50

sentences, one list per sentence structure (see Procedure, and Supporting

Information). Within a trial, no sentence was ever repeated. There were two types of

trial: regular trials, which contained only grammatically correct sentences; and

irregular trials, which contained also two ungrammatical sentences, obtained by

randomly shuffling word positions across two successive sentences. There were fifty

trials per sentence structure, half were regular and half irregular, thereby yielding

unbiased estimates of accuracy in discriminating regular from irregular trials, a task

akin to classic grammaticality judgments.

Procedure. Participants sat comfortably in an IAC 40a sound attenuating and

electrically shielded recording booth (IAC Acoustics, iacacoustics.com),

approximately 1m from an LCD computer screen. They were instructed to fixate a

cross at the centre of the screen and listen attentively to each trial; When a trial

ended, they were asked to determine whether it was regular or irregular. They were

asked to be as accurate as possible, with no time limit. Participants used their right

hand to press on two pre-defined buttons (arrow up regular, arrow down irregular) on

a keyboard. Stimuli were delivered diotically at 75 dBs SPL via loudspeakers

positioned at circa 1.2 m from participants, 25 cm to the left and right of the LCD

screen. On presentation, stimuli were further attenuated using a fixed -20 dB SPL

step, resulting in a comfortable and perceptually controlled environment. Brief rest

periods between blocks were self-determined by each participant.

Trial presentation was blocked by sentence structure. Stimuli within a trial were

presented at a constant Stimulus Onset Asynchrony (isochronous SOA): 250 ms for

monosyllabic words, 500 ms for disyllabic words. There was no gap between

sentences within a trial: Participants listened to a continuous flow of words equally

spaced in time, further reducing prosodic cues at phrasal and sentence level.

The main experimental manipulation pertained to the size - in number of words - of

Noun Phrases (NP) and Verb Phrases (VP) composing each sentence structure. An

NP is a phrase with a noun as head (e.g., "The girls"), while a VP has a verb as head

(e.g., "play rugby"). NPs usually perform the grammatical functions of verb subject or

verb object. In our experiments, all grammatical sentences contained an NP

functioning as verb subject, and a VP, which either included a second NP functioning

as a verb object or a different phrasal component. The Appendix lists all token

sentences. See Figure 2 for an exemplary analysis of a sentence used in each

condition. Stimulus sequences were created using custom scripts written in Matlab

(R2015b, 64 bit, mathworks.com). Sequence delivery was controlled by

Psychophysics Toolbox Version 3 (PTB-3, psychtoolbox.org) for Matlab, running on

a Windows 7 computer (ASIO sound card for optimal stimulus latency control).

Data Recording and Analysis. Behavioural accuracy data were subject to a signal

detection theory approach, obtaining measures of task sensitivity (d' = zscore(Hits)

minus zscore(False Alarms)) and response bias (criterion = -

0.5*(zscore(Hits)+zscore(False Alarms))) for each participant and condition [32].

We used an actiCAP 64-channel, active electrode set (10-10 system, Brain Vision

Recorder, Brain Products, brainproducts.com) to record electroencephalographic

(EEG) activity at a sampling rate of 1KHz, with a 0.1 Hz online filter (12 dB/octave

roll-off). Additionally, electrocardiographic (ECG) and electrooculographic (EOG)

signals were recorded using a standard bipolar montage. All impedances were kept

below 10 kOhm. Data were recorded with an on-line reference to FCz channel, and

offline re-referenced to the average activity of all scalp channels, and downsampled

to 250 Hz. Using the EEGLAB toolbox for Matlab [46] (sccn.ucsd.edu), continuous

EEG data were first visually inspected to remove large non-stereotypical artifacts

(e.g., sudden head movements, chewing), digitally filtered at 35 Hz low-pass (Kaiser

window, beta = 5.65326, filter order 93, transition bandwidth 10 Hz), high-pass

filtered at 1 Hz (filter order 455, transition bandwidth 2 Hz) to ensure data

stationarity, submitted to an automatic artifact rejection based on spectrum

thresholding (threshold = 10 standard deviations, 1-35 Hz), and then decomposed

into Independent spatial components using the Infomax algorithm, which allows for

optimal source separation [47]. The resulting Independent Components (ICs) were

tested using the SASICA toolbox for EEGLAB [48]: ICs reflecting blinks/vertical eye

movements, lateral eye movements and heart-beat, were detected by means of a

correlation threshold (0.7) with bipolar Vertical, Horizontal EOG, and ECG channels,

respectively, and found to be present in all participants (range: 1-3 ICs for vertical

eye movements, 1-2 ICs for horizontal, 1 IC for heart beat). To reduce inter-trial

signal variability, ICs reflecting muscle artifacts were identified via autocorrelation

(threshold = 0, lag = 20 s), while ICs reflecting focal topography was used for bad

electrodes (= 7 standard deviations threshold relative to the mean across

electrodes). ICA results were copied back to the original EEG datasets (0.1 Hz high-

pass), and ICs marked as artefactual were rejected before trial epoching. Each ten-

sentence trial varied in duration between 10 seconds (4 words per sentence, running

at 4 Hz) and 25 seconds (5 words per sentence, running at 2 Hz). Individual trial

power estimates were extracted for each electrode from Hann-windowed and

standardized data using the complex modulus of the Fast Fourier Transform,

correcting for the Hann window loss of power (sqrt(1.5)). FFT trials were then

transformed into phase coherence estimates by dividing by the amplitude of the

signal, summating the angles, taking the absolute value and normalizing by the

number of trials for each sentence type and participant. Signal-to-noise ratio

estimates (SNR) were calculated for each frequency point by normalizing its value by

the average of two samples before and two samples after it.

To verify the presence of significant neural peak and the effect of syntactic

manipulations, a series of T-tests, repeated measures analysis of variance

(rmANOVAs) and Bayesian rmANOVAs [49], were run, verifying the evidence for the

categorical harmonics model. False-discovery-rate (FDR) correction with Q value =

0.05 was applied in all cases of multiple comparisons [50]. A machine learning

approach to trial classification was run on the standardized (z-scored) phase

coherence scores for each participant and each harmonic series peak, with the

addition of the word rate peak. Three independent, numerically balanced subsets

("folds") were created for each peak from the scores of all participants ("samples"),

and an iterative process was repeated independently for each fold, while the other

two were used a training set to train an LDA (Linear Discriminant Analysis) classifier

[51]. Electrodes counted as features in this analysis. The resulting classifiers were

tested by predicting whether a trial label was irregular or regular. Accuracy was

quantified using the area under the curve of the receiver-operating characteristic

(ROC AUC)

This analysis was repeated three times, with each trial subset serving as testing set

once. A Montecarlo resampling approach (1000 repetitions) was used to obtain 95%

confidence intervals, by shuffling the order of the trial labels and replicating the LDA

classification.

Data Availability

Anonymised pre-processed datasets for this experiment, as well as the relevant

analysis scripts, are in the process of being made available in OSF

(https://osf.io/rqfws/, currently private).

Acknowledgements

The authors would like to thank Cornelius Abel, Jana Gessert, Freya Materne,

Claudia Lehr, Alexander Lindau, Georg-Friederich Paasch for help with stimulus

creation and data collection, and Dobromir Dotov for critical feedback on data

analysis.

References

1. Hickok G, Poeppel D. Dorsal and ventral streams: a framework for understanding

aspects of the functional anatomy of language. Cognition 2004; 92: 67–99.

2. Hickok G, Poeppel D. The cortical organization of speech processing. Nat Rev

Neurosci. 2007;8: 393–402.

3. Giraud AL, Poeppel D. Cortical oscillations and speech processing: emerging

computational principles and operations. Nat Neurosci. 2012;15(4) :511–517.

4. Tavano A, Scharinger M. Prediction in speech and language processing. Cortex

2015; 68: 1–7.

5. Ding N, Patel AD, Chene L, Butler H, Luo C, Poeppel D. Temporal modulations in

speech and music. Neurosci Biobehav Rev. 2017;81: 181–187.

6. Ghitza O, Greenberg S. On the possible role of brain rhythms in speech

perception: intelligibility of time-compressed speech with periodic and aperiodic

insertions of silence. Phonetica 2009;66: 11326.

7. Ghitza O. Acoustic-driven delta rhythms as prosodic markers. Lang Cogn

Neurosci. 2017;32: 54561.

8. Assaneo F, Poeppel D. The coupling between auditory and motor cortices is rate-

restricted: Evidence for an intrinsic speech-motor rhythm. Sci Adv. 2018;4(2):

eaao3842.

9. Steinhauer K, Alter K, Friederici AD. Brain potentials indicate immediate use of

prosodic cues in natural speech processing. Nat Neurosci. 1999;2: 191—196.

10. Hwang H, Steinhauer K. Phrase length matters: The interplay between implicit

prosody and syntax in Korean garden path sentences. J Cogn Neurosci.

2017;23(11): 295– 300.

11. Tavano A, Pesarin A, Murino V, Cristani M. Automatic conversational scene

analysis. low-level acoustic feature classification of speech activity in children with

Asperger syndrome and typically developing peers. PLoS One 2014;9(1): e85819.

12. Scharinger M, Steinberg J, Tavano A. Integrating speech in time depends on

temporal expectancies and attention. Cortex 2017;93: 28–40.

13. Keitel A, Gross J, Kayser C. Perceptually relevant speech tracking in auditory

and motor cortex reflects distinct linguistic features. PLoS Biology 2018;16(3):

e2004473.

14. Rimmele, JM, Sun, Y, Michalareas, G, Ghitza, O, Poeppel D. Dynamics of

functional networks for syllable and word-level processing. Biorxiv 2019, doi:

https://doi.org/10.1101/584375.

15. Chomsky N. Language architecture and its import for evolution. Neurosci

Biobehav Rev. 2017;81: 295-300.

16. Friederici AD, Hahne A, Mecklinger A. Temporal structure of syntactic parsing:

early and late event-related brain potential effects. J Exp Psychol Learn Mem Cogn.

1996;22(5): 1219-48.

17. Bornkessel-Schlesewsky I, Schlesewsky M. Toward a Neurobiologically

Plausible Model of Language-Related, Negative Event-Related Potentials. Front

Psychol. 2019;10: 298.

18. Brouwer H, Crocker MW, Venhuizen NJ, Hoeks JCJ. A Neurocomputational

Model of the N400 and the P600 in Language Processing. Cogn Sci. 2017;41(6):

1318-1352.

19. Regel S, Meyer L, Gunter TC. Distinguishing neurocognitive processes reflected

by P600 effects: evidence from ERPs and neural oscillations. PLoS One. 2014

20;9(5): e96840.

20. Ding N, Melloni L, Zhang H, Tian X, Poeppel D. Cortical tracking of hierarchical

linguistic structures in connected speech. Nat Neurosci. 2016;19: 15864.

21. Rossion B. Understanding face perception by means of human

electrophysiology. Trends Cogn Sci. 2014 Jun;18(6):310-8.

22. Ding N, Pan X, Luo C, Su N, Zhang W, Zhang J. Attention is required for

knowledge-based sequential grouping: Insights from the integration of syllables into

words. J Neurosci. 2017;38(5): 1178-1188.

23. Martin AE, Doumas LA. A mechanism for the cortical computation of hierarchical

linguistic structure. PLoS Biology 2017;15(3): e2000663.

24. Pallier C, Devauchelle AD, Dehaene S. Cortical representation of the constituent

structure of sentences. PNAS USA 2011;108: 2522–2527.

25. Pylkknen, Liina B, Bemis DK, Elorrieta EB. Building phrases in language

production: an Meg study of simple composition. Cognition 2014;133(2): 371–384.

26. Fruchter J, Linzen T, Westerlund M, Marantz A. Lexical preactivation in basic

linguistic phrases. J Cogn Neurosci. 2015;27(10): 1912–1935.

27. Pylkknen L. The neural basis of combinatory syntax and semantics. Science

2019;366(6461): 62–66.

28. Zaccarella E, Mayer L, Makuuchi M, Friederici AD. Building by syntax: The

neural basis of minimal linguistic structures. Cereb Cortex 2017;27(1): 411–421.

29. Nelson MJ, El Karoui I, Giber K, Yang X, Cohen L, Koopman H et al.

Neurophysiological dynamics of phrase-structure building during sentence

processing. PNAS USA 2017;114: E3669–E3678.

30. Chomsky, N. Remarks on Nominalizations. In: Jacobs R, Rosenbaum P, editors.

Readings in the English Transformational Grammar. Waltham, MA: Ginn and Co.;

1970. pp. 184-221.

31. Giorgi A, Longobardi, G. The syntax of noun phrases. Cambridge, UK:

Cambridge University Press; 1991.

32. Macmillan NA, Creelman CD. Detection theory: A user's guide. 2nd ed. New

York: Lawrence Erlbaum Associates Publishers; 2005.

33. Kupietz, M. BCKH, Witt A. The German reference corpus Dereko: A primordial

sample for linguistic research. In: Calzolari N, Choukri K, Maegaard B, Mariani J,

Odijk J, Piperidis S, et al., editors. Proceedings of the 7th conference on international

language resources and evaluation (Irec 2010). Valletta, Malta: European language

resources association (Elra); 2010. pp. 1848-1854.

34. Zhou H, Melloni L, Poeppel D, Ding N. Interpretations of frequency domain

analyses of neural entrainment: Periodicity, fundamental frequency, and harmonics.

Front Hum Neurosci. 2016;10: 274.

35. Schröger E, Marzecová A, SanMiguel I. Attention and prediction in human

audition: a lesson from cognitive psychophysiology. Eur J Neurosci. 2015;41(5):641-

664.

36. Frank SL, Yang J. Lexical representation explains cortical entrainment during

speech comprehension. PLoS ONE 2018;13(5): e0197304.

37. Weyerts H, Penke M, Münte TF, Heinze HJ, Clahsen H. Word order in sentence

processing: An experimental study of verb placement in German. J Psycholing Res.

2002;31(3): 21168.

38. Koster J. Word order in sentence processing: An experimental study of verb

placement in German. In: Abraham W, editor. Groninger arbeiten zur

germanistischen linguistik. Groningen, The Netherlands: University of Groningen;

1999. pp. 1-42.

39. Jackendoff R. $\overline{\text{X}}$ syntax: A study of phrase structure. Cambridge, MA: The MIT

Press; 1977.

40. Chomsky N. Bare phrase structure. Cambridge, MA: The MIT Press; 1994.

41. Bieber D. Variation across speech and writing. Cambridge: Cambridge University

Press; 1988.

42. Greenberg S, Carvey H, Hitchcock L, Chang S. Temporal properties of

spontaneous speech - a syllable-centric perspective. J Phonetics 2003;31(3): 465–

485.

43. Park H, Ince RAA, Schyns PG, Thut G, Gross J. Frontal top-down signals

increase coupling of auditory low-frequency oscillations to continuous speech in

human listeners. Curr Biol. 2015;25: 1649–1653.

44. Keitel A, Robin AA Ince, Joachim Gross CK. Auditory cortical delta-entrainment

interacts with oscillatory power in multiple fronto-parietal networks. Neuroimage

2017;147: 32–42.

45. Delorme A, Makeig S. EEGlab: an open source toolbox for analysis of single-trial

eeg dynamics. J Neurosci Methods 2004;134: 9–21.

46. Bell A, Sejnowski T. An information-maximization approach to blind separation

and blind deconvolution. Neural Computation 1995;7: 1129–1159.

47. Chaumon M. BD, Busch N. A practical guide to the selection of independent

components of the electroencephalogram for artifact correction. J Neurosci Methods

2015;250: 47–63.

48. Genovese CR, Lazar NA, Nichols T. Thresholding of statistical maps in functional

neuroimaging using the false discovery rate. Neuroimage 2002;15: 870–878.

49. JASP Team (2020). JASP (Version 0.14) [Computer software].

50. Baayen H PR, Gulikers L. The Celex lexical database [cd-rom]; 1995.

Philadelphia, PA: Linguistic data consortium, University of Pennsylvania.

51. Treder MS. MVPA-Light: A Classification and Regression Toolbox for Multi-

Dimensional Data. Front. Neurosci. 2020;14:289. doi: 10.3389/fnins.2020.00289