Integrating When and What Information in the Left Parietal Lobe Allows Language Rule Generalization

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Abstract

A crucial aspect when learning a language is discovering the rules that govern how words are

combined in order to convey meanings. Since rules are characterized by sequential co-occurrences

between elements (e.g. '***These*** cupcake*s **are un**believ**able'**), tracking the statistical relationships

between these elements is fundamental. However, statistical learning alone cannot fully account for

the ability to create abstract rule representations that can be generalized, a paramount requirement

of linguistic rules. Here, we provide evidence that, after the statistical relations between words have

been extracted, the engagement of *goal-directed attention* is key to enable rule generalization.

Incidental learning performance during a rule-learning task on an artificial language revealed a

progressive shift from statistical learning to goal-directed attention. In addition, and consistent with

the recruitment of attention, fMRI analyses of late learning stages showed left parietal activity

within a broad bilateral dorsal fronto-parietal network. Critically, rTMS on participants' peak of

activation within the left parietal cortex impaired their ability to generalize learned rules to a

structurally analogous new language. No stimulation or rTMS on a non-relevant brain region did

not have the same interfering effect on generalization. Performance on an additional attentional task

showed that rTMS on the same parietal site hindered participants' ability to integrate *what* (stimulus

identity) and *when* (stimulus timing) information about an expected target. The present findings

suggest that learning rules from speech is a two-stage process: following statistical learning, goal-

directed attention –involving left parietal regions– integrates *what* and *when* stimulus information to

facilitate rapid rule generalization.

Introduction

Our increasing understanding of the interplay between domain-specific and domain-general cognitive processes has gradually broadened our views on language learning. Apparently simple feats such as the learning of new words are no longer thought to result from the sole workings of a system specialized for language but are known to involve general-purpose mechanisms of statistical learning[1], memory consolidation[2], attention[3] or reward[4]. Besides words, linguistic proficiency requires the learning of and, particularly, the ability to *generalize rules*, which involves the development of abstract representations of grammatical categories and an understanding of their inter-relations. While this has been the topic of much research, a fundamental question remains unresolved: what are the brain mechanisms that support language rule learning and generalization?

Given its core relevance, the mechanisms supporting rule learning have been subject to intense debate. By their very nature, language rules are characterized by sequential co-occurrences, often between non-adjacent elements (e.g. **These** cupcake*s* **are un**believ*able*). Accordingly, tracking statistical relations is known to be a key computation in this context[5] as it is in many other domains[6,7]. Yet, statistical learning alone cannot fully account for the generalization of non-adjacent dependencies[8–10], and an additional role for attention has been posited as necessary[11–17]. The deployment of attention has been shown to be crucial in determining what information will be learned[16] and to bias subsequent learning[18]. Indeed, it is often the case in natural languages that dependencies occur at salient positions such as edges, with edges acting as perceptual anchor points that could additionally help to learn the positional information of the related elements[19]. In line with this, attention has been also proposed to underpin the ability to transfer the rules of a first language to a structurally similar second language[20,21], and diverting attention interferes with rule generalization[22]. However, and despite the increasing support for the implication of attention in rule learning and generalization, there is currently no direct evidence for how attention interacts with statistical learning during the extraction of language rules, and how this interaction facilitates

generalization. Moreover, the brain networks enacting this interaction for language rule learning and generalization remain largely unspecified.

In the current article, we propose that attention is essential in the process leading from the extraction of statistical regularities to the generalization of rules. In particular, we capitalized on prior research on attention to test the hypothesis that different types of attention interact with statistical learning to support language rule learning and generalization.

Research in the domain of attention posits the distinction between two networks for attention orienting[23,24]: a ventral fronto-parietal network, running along perisylvian areas from the inferior parietal lobule (IPL) to the inferior frontal gyrus (IFG), involved in the automatic detection of behaviorally relevant stimuli[23,24] (i.e., *stimulus-driven attention*); and a more dorsal and bilateral fronto-parietal attention network, including the dorsolateral prefrontal cortex (dlPFC) and the intra-parietal sulcus (IPS) extending to the superior parietal lobe (SPL)[23], involved in orienting attention towards stimulus features based on internal goals (i.e., *goal-directed attention*).

An important feature of the ventral attention network is its sensitivity to input statistics, such as the probability of a cue predicting an upcoming target[25]. Critically, while predicting a target at a specific spatial location correlates with right-lateralized activity, predicting *when* a target will appear elicits a left-lateralized response[26,27]. Given that the relationship between rule elements in speech necessarily involves a temporal dimension, we hypothesized that the tracking of statistical relationships for rule learning will predominantly engage left-lateralized ventral network activity. Supporting this hypothesis, a variety of studies on language rule learning[28–30], as well as on statistical learning for word segmentation from speech[31], report the involvement of a left-lateralized fronto-parietal network in the early stages of exposure to a new language, highlighting an overlooked overlap between the left ventral attention network for temporal orienting and the classic language network in its fronto-parietal component. Importantly, the ventral network is also thought to inform[24] the dorsal network for the generation and updating of internal models of the

environment[25] that will ultimately guide goal-directed attention. Moreover, the right/left

specialization in spatial vs. temporal domains observed in the ventral network is also observed in

the dorsal network[27] for goal-directed attention.

In line with these observations, we hypothesize that rule learning is a two-stage process involving

an initial statistical learning stage, recruiting the (left) ventral attention network, followed by the

engagement of goal-directed attention as learning proceeds, associated with the dorsal fronto-

parietal attention network. Therefore, we expected goal-directed attention to play a more prominent

role at later stages of learning, and to be critical for rule generalization to new languages by

providing the relevant temporal structure. Our specific predictions were that: i) the ventral attention

network will be primarily activated in response to rule elements in the learning of their statistical

contingency; ii) informed by this, the dorsal attention system will, in turn, direct attention towards

these relevant elements of the speech stream, both in terms of their identity (i.e., *what* information;

e.g. "un" predicting "able" in **un**forgett**able**) and the specific position (i.e. moment in time) in

which these elements are expected to occur (i.e., *when* information; e.g. word initial/final positions);

and iii) the interaction between ventral and dorsal attention networks will be particularly critical in

the *generalization* of learned rules to new instances in a language with analogous dependencies

between elements. More specifically, we predicted that goal-directed attention to the time-positions

of rule elements will facilitate the fast learning of new dependencies by the binding of new *what*

(stimulus identity) and *when* (stimulus temporal position) information[32].

Results

In order to test our hypotheses, we followed the protocol illustrated in Figure 1. Participants were

exposed to an artificial language in two different sessions: Session 1 tested rule learning, and

Session 2 tested rule generalization. In all cases, the artificial language/s used comprised three-word

phrases with no semantic content and an embedded rule consisting of a dependency between the

initial and final elements of a phrase. Importantly, the results obtained with this kind of stimulus

material are similar to those observed with real language rules[33,34] (e.g., *is* sing*ing, is* play*ing*).

Figure 1. Schematic overview of the protocol. Sessions 1 and 2 were conducted a week apart. Session 1 assessed

learning using a single language. In Session 2, generalization was assessed with two new languages that followed an

analogous structure to the language learned in Session 1. Languages were counterbalanced across the protocol. rTMS

intervention order (lPL/POz) was counterbalanced between participants in the Intervention group and performed on the

same day. The control group followed the exact same protocol but had no fMRI or rTMS intervention. lPL, left parietal

lobe; POz, midline posterior location according to the 10-20 system electrode location. L1, L2, L3 = Languages 1-3.

Session 1 consisted of two parts: in Part 1, all participants were exposed to an artificial language

(L1) in an incidental learning task. In Part 2, participants were re-exposed to the same (L1)

language and task, but divided in two groups depending on whether they were fMRI scanned

(Intervention group, N = 22) or performed the task outside of the scanner (Control group, N = 32).

The Control group was included to assess task repetition effects. Session 2 was conducted one week

after the initial session. This session was designed to test participants' ability to generalize the rule

learned on Session 1 to new but structurally analogous artificial languages (L2 and L3; i.e., L2/L3

followed the same structure as L1 but instantiated in totally new words). In Session 2, participants

in the Intervention group (top row, Fig. 1) received repetitive transcranial stimulation (rTMS) on the

peak of activation within the *left parietal lobe* (lPL), and on a control site (POz), successively.

Participants learned the new languages under the effects of the stimulation of each site. The order of

stimulation was counterbalanced across participants. The hotspot in lPL was individually identified

from the fMRI scans of the second part of Session 1. We predicted this to reflect goal-directed

attention engaged at a later stage of learning, following Part 1. Targeting the lPL with rTMS in

Session 2 was thus expected to hinder participants' goal-directed attention and hence the

generalization of the rule learned in Session 1. The Control group (bottom row, Fig. 1) was tested

with the same language tasks and with the same delays and order as the Intervention group, but

without receiving rTMS. Performance from the Control group additionally informed us about the

expected effects of language generalization without rTMS.

In each session, we evaluated rule learning of non-adjacent dependencies in artificial languages[16].

We used two types of blocks. In rule blocks, participants were exposed to three-word phrases

containing AXC-type dependencies with the first word A predicting the last word C irrespective of

the intermediate word X (e.g. 'jupo [variable word] runi'). In no-rule blocks, three-word phrases

with no dependencies (XXX-type and XXC-type strings) were used. Each three-word phrase was

considered a trial. Rule and no-rule trial blocks were interleaved (see Methods). Participants were

not informed about the presence of rules; their task was to detect the presence or absence of a given

target word (e.g. *runi*) which always took the third position in the phrase, in both rule and no-rule

blocks.

The word-monitoring task measures incidental learning of the dependencies[16] such that, as

learning progresses, participants are expected to start anticipating the presence/absence of the target

word (C) upon hearing the first word (A) of the phrase in the rule blocks. Learning progression,

evidenced by a reaction time *gain* over trials, can be approximated as a *learning slope* via

regression analysis (Fig. 2). A negative learning slope can thus be interpreted as an indication of

statistical learning in the initial stages (Fig. 2; see also *Methods*). As learning accrues, we expected

participants' reaction times to gradually reach a plateau, indicating that they can now use the initial

element of each dependency to consistently anticipate the dependent element. Such plateau should

be expressed in later learning stages as stable RTs over the course of the trials (relatively flat

learning slope) for the rule blocks, that is a slope similar to no-rule blocks albeit with faster mean

reaction times (RTs) in rule blocks compared to no-rule blocks. This RT advantage in rule-blocks

(henceforth, *rule effect),* is calculated as the difference in the mean reaction time between rule and

no-rule blocks; Fig. 2).

Figure 2. Hypothesized reaction time slopes for rule and no-rule blocks over repetitions of the incidental rule-

learning task. For each artificial language learned, participants were exposed to blocks with rules, where the initial

word determined the identity of the last word of the phrase, and no-rule blocks where the final word could not be

predicted based on the first one. A. Part 1: Reflecting statistical learning, rule blocks are expected to exhibit a greater

gain in reaction times across trials than no-rule blocks, as a consequence of the ability to predict the upcoming occurrence

or absence of the target word. The difference between rule and no-rule slopes (Learning Slope) is thus a measure of

statistical learning indicating progressive rule learning in the early stages. **B.** Part 2: If participants can benefit from

previous learnings to orient attention to the initial element to consistently anticipate the final one, their reaction times for

rule blocks should plateau in later learning stages and show a sustained difference compared to no-rule blocks throughout

(Rule Effect, i.e., the mean difference in reaction time between rule and no-rule trials). **C.** A plateau should also be

observed for participants that generalize their attentional focus on initial and final elements to a new language with the

same type of dependencies (i.e. rule).

This pattern of behavior is to be expected if participants can apply the knowledge acquired during

the initial stage to focus their attention in a goal-directed manner on the initial element of the phrase

in order to consistently predict the appearance of the target word at the end of the phrase. Finally, a

stable rule effect (i.e., a sustained difference in reaction time between rule and no-rule phrases)

should also be expected in the learning of a new but structurally analogous language, given by the

generalization of the learned attentional focus to the new material[20,21].

During the first session, participants (N = 54, 39 women, mean age= 22.61 years, *sd* = 5.75)

performed the incidental rule-learning task on an artificial language (e.g. L1). A short break was

provided midway through the session (Session 1, see Fig. 1). Out of the main cohort, a subgroup of

participants (Intervention group; N = 22, 13 women, mean age = 23.63 years, *sd* = 4.67) was

randomly selected to undergo the rTMS intervention in Session 2. In order to obtain the appropriate

coordinates for rTMS stimulation, participants in the Intervention group performed the second part

of the rule-learning task in the MRI scanner (i.e. Rule-learning 1, Part 2). To assess task repetition

effects, the remaining participants (Control group; N = 32, 26 women, mean age = 21.9 years, *sd* =

6.37) were behaviorally tested with the exact same protocol as the Intervention group but without

fMRI in Session 1 or rTMS in Session 2.

Figure 3 illustrates changes in RT across trials as the task was repeated across the two parts of

Session 1. A linear mixed model approach was used to obtain the *learning slopes* for the two rule

learning parts of Session 1, and to compare rule vs. no-rule performance within each part of the

session. In addition, *rule effects* were calculated by contrasting mean RTs in rule and no-rule blocks

within each part and were assessed via paired *t*-tests (see *Methods*; four subjects were excluded

from Part 1 due to missing data).

Figure 3. Incidental rule-learning task results for Session 1. Slopes for rule and no-rule blocks (N = 54) over task
repetitions derived from the mixed model analysis conform to the expected pattern (Fig. 2), with a significant Learning
Slope in Part 1 and a significant Rule Effect with a flat (i.e., non-significant) Learning Slope in Part 2. Actual data shown
averaged into 6 trial bins (for visualisation purpuses only; the analysis did not bin the data) with the SEM, over the slopes
for rule and no-rule derived from the mixed model analysis.

For the entire sample, a steeper *learning slope* (i.e. more negative) for rule blocks compared to no-
rule blocks was observed during Part 1 ($ß_{diff}$ = -0.73, t = -3.86 p < 0.0002; $ß_{diff}$ is the estimate of the
difference in learning slopes between the rule and no-rule conditions, see *Methods*). The significant
difference indicates that learning of the statistical relations between words occurred at this stage. As
expected (Fig. 2), non-significant (flatter) slopes for rule blocks were then obtained the second time
the task was performed (Part 2: $ß_{diff}$ = -0.36, t = -1.81, p > 0.07). Crucially, the slope in the rule
condition in Part 1 was significantly more pronounced than that of Part 2 ($ß_{diff}$ = 0.69, t = 3.32, p <
0.001). These changes in the slope were accompanied by a significant *rule effect* in both Part 1
($t(49)$ = 3.63, p < 0.001, d_{Cohen} = 0.513) and Part 2 ($t(53)$ = 4.00, p < 0.001, d_{Cohen} = 0.545). The
rule effect was equivalent in Parts 1 and 2 (non-significant difference between the two, p = 0.649,
d_{Cohen} = 0.065). The combined findings of a non-significant learning slope (significantly flatter than
that for Part 1) and a significant rule effect in Part 2 supports the view of rule-learning as a two
stage-process (Fig. 2). An analysis of the Intervention and Control groups separately indicated a

similar pattern of results in the two groups, except that in the Intervention group some learning was

still in progress in Part 2 (Fig. S1 and Fig. S2).

Functional localization of parietal areas for stimulation

With the purpose of selecting the coordinates for the rTMS intervention, we identified parietal

regions engaged in the late stages of learning of Session 1 at the individual level. In particular, we

localized the peak voxel with the highest activity from single-subject contrast maps for the rule vs.

no-rule contrast within the left parietal lobe. The decision to stimulate the left parietal lobe was

taken given the critical role this region has goal-directed attention in the temporal domain [27]. As

mentioned in the introduction, we expected this process to be relevant for rule learning in later

stages (Part 2, Session 1; see Fig. 2) as well as during generalization (i.e. Rule Learning Part 2 and

beyond). In the cases where the rule vs. no-rule contrast showed no significant activity difference

(uncorrected $p < 0.05$), we used the contrast rule condition against the implicit baseline (N=5). In

order to illustrate the variability in individual activation during rule learning, masks for each

participant's activation pattern were calculated using a $p < 0.005$ uncorrected threshold as a cut-off.

These individual masks were then added to create a group overlap map. Figure S3 depicts the

resulting overlap, showing voxels that were activated in at least 10 participants at the individual

level during the rule-learning task. Two subjects were excluded from this analysis due to technical

failure during fMRI scanning.

Exploration of the brain networks engaged during late learning stages (Part 2, Session 1)

Given the potential mixture of slow and fast learners in the Intervention group (Fig. S2), we

performed an exploratory individual differences analysis of their fMRI activity. This analysis was

aimed at investigating the networks related to statistical learning and goal-directed attention by

correlating participants' corresponding behavioral measures (i.e. learning slope and rule effects)

with the appropriate fMRI contrasts.

To identify the neural activity related to statistical learning, we derived individual *learning slopes*

for rule blocks ran in the fMRI (Part 2, Session 1). These individual learning slopes were then

correlated with the BOLD signal change in the same blocks (see *Methods*). Covarying with

individual differences in the slope for rule blocks, this analysis revealed a ventral fronto-parietal

network as related to statistical learning (see Fig. 4A). This ventral network included two large

clusters of perisylvian areas in the left hemisphere including the left inferior frontal gyrus (IFG) and

the left inferior parietal lobule (IPL), and a smaller cluster centered on the right IFG (Table S1).

Figure 4. A. BOLD signal over a ventral fronto-parietal network in rule blocks significantly co-varies with their

corresponding measure of statistical learning (Learning Slope); the more activity, the greater (i.e. more negative) the slope

(see also Figure S4). **B.** BOLD signal activity over a dorsal fronto-parietal network (rule blocks – no rule blocks) co-

varying with the measure of goal-directed attention (Part 2 Rule Effect – Part 1 Rule Effect); the more activity, the larger

the effect. Only significant results ($p < 0.05$ FWE-corrected at the cluster level, with an additional $p < 0.001$ at the voxel

level and 50 voxels of cluster extent) are shown for both analysis. IFG = Inferior Frontal Gyrus; IPL = Inferior Parietal

Lobule; SMG = Supra-Marginal Gyrus; INS = Insula; MFG = Middle Frontal Gyrus; PostCG = Post-Central Gyrus; SPL

= Superior Parietal Lobe; MidCing = Mid-Cingulum; Precu = Precuneus. Neurological convention is used with MNI

coordinates shown at the bottom right of each slice.

Since we hypothesized that goal-directed attention would increase over exposure to the same

artificial language, to identify neural activity related to this process, we estimated individual

measures corresponding to the change in the *rule effect* (rule vs. no-rule RTs) from Part 1 to Part 2

(i.e. Part 2 Rule Effect – Part 1 Rule Effect). We then correlated this *rule effect* increment with the

rule *vs.* no-rule fMRI contrast. Individual differences in the increment of the *rule effect* from Part 1

to Part 2 covaried with activity in a more dorsal and bilateral fronto-parietal network, consistent

with the engagement of goal-directed attention. This dorsal network included two large clusters

centered on bilateral and predominantly superior parietal regions extending to the precuneus, as

well as left middle and superior frontal gyri. In addition, there was also an involvement of the left

IPL and bilateral basal ganglia (Table S2).

Note that, due to the limited number of subjects, these results can only be interpreted tentatively.

However, they do hint in the direction of the hypothetized gradual transition from more ventral

fronto-parietal regions for statistical learning to more dorsal fronto-parietal areas for goal-directed

attention in later stages of learning.

Is the left parietal lobe causally related to rule generalization and goal-directed attention?

The same participants (Intervention and Control groups) came to a second session one week later

(Session 2: Fig. 1) to test whether the left parietal lobe areas activated in the late stages of learning

(functionally localized in Session 1) were causally related to rule *generalization*, and (ii) whether

this area was indeed involved in goal-directed attention for the binding of *what* and *when*

information. Session 2 thus assessed participants' rule learning performance on two new languages

(L2 and L3) that followed the same rule structure as the language (L1) used in Session 1 (i.e. where

the initial word determined the identity of the final word). Participants in the Intervention group

performed the rule learning task after rTMS on the lPL (individually determined by the functional

localization analysis; see *Methods*) and a control site (POz; order of stimulation site and language

counterbalanced). Participants in the Control group were also tested in Session 2 with the same new

languages (L2 and L3), just like the Intervention group, but without rTMS (Control group Session

2, see Fig. 1). Their performance was used to measure the effects of task repetition and to have a

measure of the baseline levels of generalization without rTMS.

We expected participants' goal-directed attention to be engaged in the transfer of the rule

knowledge acquired in Session 1 to structurally similar languages in the rule-learning tasks of

Session 2. A mixed model analysis of their RTs indicated, as predicted, that rule-performance in the

Control group in Session 2 followed the same pattern observed in Part 2 of Session 1 despite

performing the task on a new language. That is, participants showed faster RTs in rule blocks

compared to no-rule blocks (rule effect: $t(31) = 2.434$, $p < 0.021$, $d_{Cohen} = 0.430$) but no significant

slope difference between the blocks ($t(31) = -1.6$, $p > 0.11$; Fig. 5). Moreover, the slope for the

Control group during Session 2 was not significantly different from their slope for Part 2 in Session

1 ($t(31) = -0.84$, $p > 0.4$).

The same pattern was observed for the Intervention group after rTMS to the control site POz (rule

effect: $t(19) = 2.85$, $p < 0.01$, $d_{Cohen} = 0.638$; rule *vs.* no-rule slope difference: $t(19) = 0.182$, $p >$

0.85). Indeed, both the rule effect and the rule slope under rTMS POz were comparable to those of

the Control group (respectively, $t(50) = 2.85$, $p = 0.271$, $d_{Cohen} = 0.317$ and t(51) = -0.79, p > 0.4).

This indicates that stimulation to POz did not influence task performance (i.e., the Control group

and the Intervention group during rTMS POz have comparable slopes and rule effects) and thus

confirms rTMS POz as a suitable control site for participants' performance under rTMS lPL. This

pattern of results supports the idea that participants' knowledge of the rule structure acquired early

on in learning (Session 1) was used to perform the task in this later phase. Thus, the behavioral

plateau achieved in the later stages of Session 1 (Part 2) was maintained when a new language

following the same type of rule was being learned and could be used in generalization.

Finally, we assessed the effects of rTMS over lPL on rule generalization performance. As can be

observed in Figure 5, the pattern of performance in the Intervention group after the application of

rTMS on the lPL stood in stark contrast to the pattern seen in both the Control group and in the

Intervention group after rTMS to the control site POz (Session 2). First, the slope difference for rule

blocks compared to no-rule blocks was significant after rTMS stimulation to the lPL (β_{diff} = -0.95, t

= -3.75, $p < 0.0002$; Fig. 5), in line with the pattern observed in the early stages of rule learning

(i.e., Session 1, Part 1). The slope for rule blocks in this phase was significantly more negative than

both the slope after rTMS stimulation to the control site POz (β_{diff} = -0.67, t = -2.66, p = 0.008) and

that of the Control group Session 2 (β_{diff} = -0.57, t = -2.49, p = 0.013), suggesting that after rTMS

over the lPL subjects performed the task capitalizing on statistical learning. Note that the effects of

rTMS on the lPL were not driven by the performance in the no-rule blocks: there were no

significant differences between the slopes for the no-rule blocks in rTMS lPL and rTMS POz (β_{diff} =

0.29 , t = 1.17 , p = 0.24), rTMS POz and Control group Session 2 (β_{diff} = 0.21, t = 0.94, p = 0.35),

or rTMS lPL and Controls group Session 2 (β_{diff} = -0.08, t = -0.33, p = 0.74). To rule out any order

effects (lPL stimulated first/second), *order* was added as a factor to the model for both rTMS lPL

and rTMS POz phases (*Methods*), yielding non-significant effects in both cases ($p > 0.1$). Finally,

we note that the overall faster RTs for rTMS conditions compared to RTs for the Control group

may be due to non-specific rTMS effects[35] and/or the lower latency equipment employed for

testing in the rTMS facilities and do not affect the main outcome.

Figure 5. Incidental rule-learning task results for Session 2. Both rTMS POz intervention (left panel) and Control

group Session 2 (right panel) show the expected pattern of rule learning with a significant Rule Effect and a non-

significant Learning Slope, indicating that attentional focus generalized to the learning of the new languages. In contrast, a

significant Learning Slope under rTMS lPL effects (center panel) suggests a return to the progressive rule-learning of

early learning stages (i.e., statistical learning). Actual data shown averaged into 6 trial bins (for visual purpuses only; the

analysis did not bin the data) with the SEM, over the slopes for rule and no-rule derived from the mixed model analysis.

The effects of rTMS on IPL on goal-oriented attention

To discern the precise role of the lPL, participants in the Intervention group performed an additional

attention task specifically designed to assess the ability to use content (*what:* e.g. "un" predicting

"able" in **un**forgett**able**) and temporal cues (*when:* e.g. word initial/final positions). The attention

task was administered under the effects of rTMS to lPL and POz, just like the rule learning task

(Fig. 1), with the order of tasks counterbalanced between participants (Fig. 6). Participants were

asked to judge whether the pitch of a target syllable at the end of a sequence of alternating syllables

was higher or lower than the preceding sequence. In order to build a task as similar as possible to

the language paradigm used for the rule learning experiment, we used syllables as the auditory

stimuli on which to manipulate attention. More critically, in the language task, the temporal interval

between words was fully predictable. Thus, for the Attention task, temporal orienting (*when*

information) was manipulated by presenting the sequences of syllables in either rhythmic (with

isochronous inter-stimulus intervals) or non-rhythmic form (variable time interval between

syllables) while keeping the overall trial length constant. Therefore, in both tasks the temporal

relation between elements follows the same implicit temporal expectation. We chose rhythm

because the empirical and theoretical foundations of isochronous-anisochronous event timing are

well-established as a manipulation of temporal attention[36–38]. In addition, there is solid evidence

with respect to the effect of rhythm in speech and rule-dependency processing and learning in

language and an extensive theoretical background relating temporal attention to speech processing

[39–43]. On the other hand, in the language task, the identity of the first element informed the

identity of the last element in rule blocks while this was uninformative in the no-rule blocks. Thus,

in the attention task, the first syllable of the sequence was informative or uninformative of the

identity of the final target syllable. Attention to the content (*what* information) was manipulated by

asking the participants to focus their attention to the identity of the first syllable since this would

help them to perform the task. The attention manipulations used and the orthogonality of the

participant's goal (i.e. pitch discrimination) are comparable to those previously used in temporal

attention research[37,44,45]. Participants were familiarized with the attention task before receiving

rTMS.

Figure 6. Illustration of the experimental design of the Attention Task. Participants had to judge the pitch of a target

syllable presented after a sequence of alternating syllables. The pitch of the target syllable could be either higher or lower

than that of the preceding sequence of syllables. Sequences of syllables were presented either rhythmically to engage

temporal orienting (attention to *when*) or non-rhythmically (50%), with an otherwise constant trial length. At the same

time, the initial syllable of each sequence could be informative or non-informative of the identity of the target syllable

(50%), manipulating identity-based attention (i.e. attention to *what*).

In order to test whether *what* and *when* cues where used differently in the two intervention

conditions we performed a 2x2x2 repeated measures ANOVA on reaction times to the target

syllable (correct responses only), with Intervention (rTMS_lPL, rTMS_POz), Rhythm (rhythmic,

non-rhythmic) and Identity (informative, non-informative) as within-subject factors. This analysis

revealed a main effect of Intervention ($F(1,18) = 13.36$, p = 0.002, Partial $\eta^2 = 0.426$), with overall

reaction times slower after rTMS to the lPL compared to rTMS POz. Regarding the use of *when*

cues, pitch discrimination in rhythmic sequences was faster than in non-rhythmic sequences (main

effect of Rhythm: $F(1,18) = 24.62$, $p < 0.001$). Concerning the effect of *what* cues, there was also a

facilitation in reaction times for informative compared to non-informative sequences (main effect of

Identity: $F(1,18) = 9.01$, $p = 0.008$). The main effects of Rhythm and Identity thus indicated that the

attentional manipulations were effective and participants benefited from the use of both *what* and

when information. Crucially for our hypothesis, we observed a significant triple interaction between

Intervention, Identity and Rhythm ($F(1,18) = 7.54$, $p = 0.013$, Partial $\eta^2 = 0.295$), not affected by

the order of rTMS intervention ($F < 1$), suggesting differences in how *when* and *what* cues were

used under each rTMS intervention (Fig. 7). We then proceeded to unpack the interaction by

analyzing how information about *what* was affected by the presence or absence of predictive

information about *when* the target is expected in each rTMS intervention. In sequences in which

when information was unpredictable (non-rhythmic sequences), Identity had a comparable

facilitatory effect in both rTMS sites (Identity: $F(1,19) = 9.58$, $p = 0.006$, Partial $\eta^2 = 0.335$;

Intervention x Identity: $F(1,19) = 1.12$, p > 0.3, Partial $\eta^2 = 0.056$). However, in sequences in which

when information was predictive (rhythmic sequences), Identity had a facilitatory effect (Identity:

$F(1,19) = 4.64$, p = 0.044, Partial $\eta^2 = 0.196$) that differed in the two rTMS stimulated sites

(Intervention x Identity: $F(1,19) = 13.18$, $p < 0.002$, Partial $\eta^2 = 0.410$). While participants could

benefit from knowing *what* the identity of the target was while stimulated at the control site (rTMS

POz, $t(19) = 5.51$, $p < 0.0001$, $d_{Cohen} = 0.596$), this facilitation disappeared under rTMS lPL

stimulation ($t(19) = -0.22$, $p > 0.8$, $d_{Cohen} = 0.027$). These results indicated that while able to benefit

from the *what* information in isolation, participants were impaired in their ability to benefit from it

when both *what* and *when* cues were presented in *conjunction*.

Figure 7. Attention task results. Black and grey shaded bars represent mean RTs with their SEM for informative and
non-informative trials, respectively, for the phases with rTMS on POz and lPL, separated into rhythmic and non-rhytmic
conditions (***p < 0.0001, main effect of Identity; * p < 0.006, pairwise comparison).

The rTMS lPL intervention therefore appeared to impact the interaction between the two forms of

attention, that is, on the joint use of *both when* and *what* cues to optimize performance in the pitch

discrimination task. In sum, while rhythm and information about future content seem to act jointly

to decrease RTs for discrimination, this benefit was lost under rTMS at lPL, where either content or

rhythm but *not both concurrently* could be used to improve task performance.

Discussion

In this study, we provide converging behavioral and neuroimaging data in favor of the hypothesis

that successful rule learning is a two-stage process. In the early exposure to an artificial language,

participants showed a gradual decrease in response times for rule compared to no-rule phrases

consistent with the progressive learning of the statistical relationship between rule elements (Fig. 3,

left panel). In the latter part of exposure, a sustained advantage in response times for rule phrases

over no-rule phrases (Fig. 3, right panel) is supported by a dorsal fronto-parietal network (Fig. 4,

and Fig. S2). This advantage in response times was maintained for new languages with the same

kind of rule. This is consistent with the hypothesis that goal-directed attention in later learning

stages facilitates the abstraction of the underlying rule, allowing participants to generalize their

acquired knowledge regarding the position (i.e. initial-final) of the elements of the rule to new

material (Fig. 5; Control group; and Intervention group, left panel). Moreover, rTMS to a key node

of the dorsal network (the lPL) interfered with rule generalization (Fig. 5, Intervention group, right

panel), pointing to a further role for goal-directed attention in this stage. Finally, by means of an

additional attention task, we were also able to specify the mechanism underlying generalization as

carried out by the dorsal network. In particular, rTMS to the lPL hindered participants' ability to

integrate stimulus temporal and identity information, that is, information about *what* element is

expected, and *when* (Fig. 7), suggesting that rule generalization involves the integration of such

features.

Here, by contrasting reaction times over trials for rule and no-rule phrases in an orthogonal word-

monitoring task[16], we show that rule learning is a two-stage process involving: i) the progressive

learning of statistical contingencies, followed by ii) the engagement of goal-directed attention for

rule extraction and generalization. The use of this *online* measure of learning thus enabled us to tap

into processes that parallel those arising in natural language learning, where rule knowledge accrues

from successive encounters with a particular grammatical rule. Our results -replicated across

different sessions and cohorts- further speak for the reliability of this measure of *online* learning,

and add substantially to the knowledge gained by the use of test measures acquired after language

exposure (i.e. *offline*), which are blind to the learning dynamics[46,47].

We hypothesize that, while still tied to specific content (e.g. the particular words forming a

dependency), the ability to direct attention to particular moments in time at later learning stages

leads to the emergence of abstract knowledge regarding classes of words (i.e. categories of words

that can occur at specific positions in a phrase), which facilitates generalization (*cf.*[48]). Note that

category knowledge implies the notion of time-bound category *slots* and *placeholders*, relating to

how specific content is apt to take a particular slot in time. What is further required for

generalization is the transfer of the learned temporal structure (i.e., the attentional focus) to the new

material, as it occurs in the acquisition of a second language[20]. That is, generalization simply

involves the reverse process of filling in the category slots with the specific word-forms of the new

language.

The two-stage account of rule learning evidenced at the behavioral level is also supported by the

observed shift in the cortical networks involved. Approximated as a learning slope, statistical

learning in the early stages correlated with brain activity in a left-lateralized *ventral* fronto-parietal

network (Fig. 4A). In contrast, later-arising sustained rule effects (mean reaction time differences

between rule and no-rule phrases) engaged a more *dorsal* and bilateral fronto-parietal network (Fig.

4B) related to goal-directed attention[23]. Although these fMRI analyses should be taken with

caution due to the limited sample size, activity in the reported areas is consistent with the existing

literature on both language learning (24,26,27,34–38) and attention[23].

Regarding language learning, several studies have previously related regions of the ventral network,

such as the inferior frontal gyrus and parietal lobule, to statistical word[51–53] and rule

learning[13,28,30,31,49,50,54,55]. These studies have significantly added to a wealth of literature

implicating the IFG in syntax processing more generally (e.g.[29,56–58]) by extending its role to

the *learning* of, at least, simple syntactic rules. Opitz and Friederici (2004), for example, reported

an increase in the activity of ventral premotor cortex following a rule violation during the learning

of an artificial grammar[28]. More anterior areas in the IFG have also been related to successful

non-adjacent dependency learning more specifically[59], along with the caudate nucleus of the

basal ganglia[31]. Goranskaya et al. (2016), additionally, found that non-adjacent dependency

learners activated a broader fronto-parietal network, including not only left frontal but also bilateral

parietal regions[30]. By bridging rule learning and attention, our study now sheds light onto the

precise role of these regions. Indeed, the same fronto-parietal network has been classically

associated with the detection of behaviorally relevant stimuli (i.e. stimulus-driven attention[24]).

More recently, however, this ventral attention network has also been shown to respond to statistical

regularities in the environment (cue-target contingencies[25]. This may be particularly interesting

for the interpretation of our results, as the engagement of the ventral attention network in our work

could reflect the detection of rule elements (predictive and predicted), which become relevant by

virtue of their statistical relationship. This link between the learning of statistical dependencies and

stimulus driven attention is in line with previous studies relating language rule learning to an event-

related potential (the P2) associated with this form of attention[13,54,55]. Finally, a left-

lateralization of the ventral network has been shown for stimulus-driven temporal orienting[27].

This is consistent with our reported left-lateralized activity in this network in response to rules

instantiated in *speech*, which necessarily imposes a temporal dimension.

In contrast to the ventral network, the increment in *rule effect* arising in later learning stages was

related to activity in a more dorsal and bilateral fronto-parietal network. This dorsal network has

been traditionally associated with the goal-directed orienting of attention, in both the spatial[23] and

temporal[27] domains. We further confirmed the causal implication of this network in rule

generalization by the interference of its left parietal node, identified for each individual during later

learning. Indeed, rTMS to this region hindered participants' ability to use acquired rule knowledge

for the fast learning of new dependencies. It is important to mention though that the causal role of

this region does not imply that other brain areas within this network, including the contralateral

parietal lobe, do not play a role. Further research is necessary to answer this question. It is

interesting as well that while the same participants stimulated on a non-relevant cortical area (POz)

showed the expected generalization behavior, in line with the Control group, participants under the

effects of rTMS lPL were seen to return to progressive rule-learning, suggesting that left parietal

stimulation did not hinder statistical learning. In other words, rTMS on the lPL did not impede the

computation and use of statistical and temporal order information for the gradual learning of the

rules. This further speaks for the relative independence of the two mechanisms, congruent with their

putative support by the two reported cortical networks.

Finally, we devised an attention task to clarify the mechanisms underlying the lPL function. In

particular, we measured participants' ability to use temporal and stimulus identity cues (i.e.

information about *when* and *what*, respectively) under the same rTMS effects. Our results show

that, although participants could still benefit from either *what* or *when* -type cues, rTMS on the left

parietal cortex interfered with the use of both kinds of cues *in conjunction*. This suggests that the

binding of stimulus temporal and identity information is required for successful generalization, and

is a characteristic of goal-directed attention in latter learning stages. We were careful to design the

attention task to match the language task as closely as possible, and the results obtained support our

hypothesis. The lPL was proven to be relevant for the integration of what and when information, a

mechanism that we claim is key in the rule generalization process. The exact same region was

critical for rule generalization. Despite this, we acknowledge that the fact that the same lPL region

has both an effect on generalization and in the integration of *what* and *when* information does not

necessarily imply that the two functions are linked. It could simply be due to the lPL supporting

both functions without them being related. Further investigations will shed light onto this issue.

However, it is worth noting that our proposal coincides with the role of the parietal lobe in structure

learning very recently proposed in the visual domain. This function is argued to derive from this

area encoding the "relative position of objects in space" and "the relations among entities in abstract

conceptual space" [60].

On a more general note, our results are consistent with the dual-process hypothesis of rule learning

from speech that we recently proposed in the developmental domain[17]. Specifically, we

hypothesized that the maturation of the dorsal fronto-parietal network facilitates the exploitation of

knowledge acquired through statistical learning, via attentional mechanisms. Hence, the gradual

acquisition of statistical contingencies via incidental learning will allow children after their second

year of life to form abstract, long-lasting rule-based knowledge. The results of the present study

suggest a similar dual-process mechanism in adult learners when facing a new language.

In conclusion, our data supports the view of two distinct stages in rule-learning, predominantly

characterized by the engagement of stimulus-driven and goal-directed attention, respectively.

Importantly, we have established a causal link between left parietal function as engaged in later

learning stages and the abstraction and subsequent generalization of language rules. Interfering with

this region caused both an impairment in goal-directed attention and in the ability to generalize

rules in language. In considering the precise implication and role of attentional mechanisms,

therefore, this view of rule-learning integrates seemingly irreconcilable experimental findings and

theoretical proposals[8,9,11,22,61].

Materials and Methods

Participants

The study included one group of participants (N = 54, 39 women, mean = 22.61, *sd* = 5.75) who

completed the whole protocol (Session 1 and Session 2) under different conditions. A subgroup of

22 right-handed participants (Intervention group: 13 women, mean age = 23.63 years, *sd* = 4.67),

underwent an fMRI session (part 2, Session 1) and rTMS (Session 2). The remaining 32 right-

handed participants formed the Control group (26 women, mean age = 21.9 years, *sd* = 6.37) who

was exposed to the artificial languages the same number of times as the Intervention group but

without fMRI acquisition or rTMS. The Control group was crucial to characterize the course of

events under repeated exposure to the same language task and to have a baseline measure of

generalization effects under no intervention. Since there was no comparable study using TMS and a

similar paradigm, the sample size was determined based on previous studies applying TMS to the

parietal cortex. The sample size in these studies is between 12 subjects (e.g.[62,63]) and 18 subjects

(e.g.[64]) with some studies reaching 24 participants (e.g.[62,65]). The initial sample size was set to

22 with a final sample size of 20 that was thus expected to be largely sufficient considering that we

were the in the range of the maximum sample size used the TMS studies and the main comparisons

were within subjects for the rTMS intervention.

All participants were native Spanish speakers and had no history of neurological or auditory

problems. Participants in the Intervention group were screened for compatibility with fMRI and

rTMS procedures[66]. The study was approved by the ethics committee of the Universitat de

Barcelona, the Universitat Pompeu Fabra (CIEC Parc de Mar) and the European Reseach Council

ethics scientific office, and was conducted in accordance with the Declaration of Helsinki.

Participants were remunerated and completed a written informed consent.

In each part and session of the experiment, several participants within each group were excluded for

technical reasons or after being identified as outliers within their group. We considered participants

as outliers when their learning slopes were 2 SD above or below the mean for their group and part.

In Session 1, two participants in Part 1 (final sample N=30) and one participant in Part 2 (final

sample N=31) of the Control group were identified as outliers. For the Intervention group in Session

1, five participants were excluded from Part 1 (one outlier and four for technical reasons; final

sample N=17) and two were identified as outliers during Part 2 (fMRI; N=19). In Session 2, none of

the participants in the control group were identified as outliers (final sample N=32). Two

participants from the Intervention group were excluded (final sample N=20) because they were

stimulated > 2 mm from the appropriate coordinate within the lPL during rTMS (see Repetitive

TMS protocol section below) due to an Experimenter error.

 Experimental design

The overall experiment (main text Fig. 1) comprised two sessions that took place approximately a

week apart from one another. In Session 1, participants performed an artificial language incidental

rule-learning task twice (Part 1, behavioral only for both the Intervention and Control groups; Part

2, behavioral for the Control group and during fMRI scanning for the Intervention one), followed

by an offline recognition test (see *Supporting Information* for further details on the offline

recognition test). The artificial language (L1) and its non-adjacent dependencies were the same for

both Part 1 and 2 of Session 1. For the Intervention group, the fMRI data was used to identify

relevant activity within the lPL during late stages of rule learning in order to select appropriate

coordinates for rTMS stimulation. For the Intervention group only, during Session 2, 1Hz rTMS (15

minutes off-line) was used to stimulate the individually determined maximum peak of activation

within the left parietal lobe (rTMS lPL phase) or a task-irrelevant brain area (POz electrode

location; rTMS POz phase). The order between the rTMS lPL phase and the rTMS POz phase was

counter-balanced across participants. After rTMS application, participants performed two tasks: a

language task similar to that in Session 1; and an attention task, which tested for the effects of

rTMS (lPL/POz) on the goal-directed use of content and temporal cues (i.e., *what* and *when*

information). Importantly, the total duration of the attention plus rule learning tasks was kept within

30 minutes so as to ensure post rTMS performance under the estimated rTMS effects (see section

on Repetitive TMS protocol below). During Session 2, the Control group completed the same

attention and language tasks as the Intervention group but without rTMS stimulation.

Rule Learning Task

Three different artificial languages counting twenty-eight bi-syllabic (consonant-vowel-consonant-

vowel) nonsense words each were created. One language (L1) was used in Session 1, the remaining

two languages (L2 and L3) in Session 2 (main text, Fig. 1). The order of languages was

counterbalanced between sessions and participants (see Table S5 for the full list of stimuli used).

Twenty-eight words were created for each language stream and synthesized using Mbrola speech

synthesizer software[67] by concatenating diphones from the Spanish male database

(http://tcts.fpms.ac.be/synthesis/) at 16 KHz. Words (385 ms), were combined using Adobe

Audition® software to form three-word phrases with 100 ms gaps between word. Phrase stimuli

were presented via Presentation® software (Neurobehavioral Systems) through appropriate

headphones and at a volume level adjusted for the participant.

A total of 96 rule and 96 no-rule phrases (trials) were used in this task. Rule phrases conformed to

an AXC structure whereby the initial word (A) always predicted the final word (C), while the

middle word (X) was variable. Two different A_C dependencies (A1_C1 and A2_C2) were created

out of 4 words from the total word pool. The remaining 24 served as middle (X) elements for each

of the two A_C dependencies. The transitional probability was always 1 between A and C elements,

0.04 between A and X, and 0.5 between X and C. Half of the no-rule trials consisted in the

combination of 3 of the 24 X elements and so took the form XXX, with the only constraints that

each X had an equal probability to appear in each position but could never appear twice in the same

phrase. The other half of the no-rule trials consisted of the combination of two XX elements

(following the same constraints as for XXX) followed by the participant's target word (C1 or C2).

The probability of target occurrence in both the rule and the no-rule blocks was therefore 50%. Note

that in the set of no-rule trials, the C element (the participant's target) occurred also in the last

position but, in contrast to the rule block, this could not be predicted on the basis of previous

elements.

Participants were presented with the randomized 96 rule and 96 no-rule phrases in four alternated

rule and no-rule blocks, with the order of blocks counterbalanced between participants. In the fMRI

version of the task, data was acquired in two runs, including a block of rule and no-rule each

(counterbalanced). A short break was given between runs in the fMRI. A single offline recognition

test was issued after the fourth block (see Supporting Information). In order to obtain a measure of

incidental rule-learning, participants performed a cover word-monitoring task. Specifically, they

were instructed to detect, as fast and accurately as possible via a button press, the presence or

absence of a given target word, which was always one of the C elements (C1 or C2,

counterbalanced). A given target word remained constant for each participant throughout the

experiment and was displayed in the middle of the screen at all times for reference during the

blocks. Participants were not informed about the presence of rules. Inter-trial interval was jittered,

using pseudo-random values between 1000 and 3000 ms for optimal fMRI acquisition, and fixed at

500 ms in the remaining phases. A maximum of 1000 ms after the end of a given phrase was

allowed for participants to respond before the next trial started. Reaction times (RTs) were

calculated from the onset the last word in the phrase until button press. Performance in interleaved

rule and no-rule blocks was jointly analyzed by concatenating blocks of a same kind. Only correct

response trials with RTs within mean $\pm$ 2sd were included for the analysis (mean over groups and

conditions of $5.57 \pm 3.55\%$ of total trials removed; note that the rejection rate for the comparisons

of interest was similar between groups: $4.79 \pm 1.36\%$ for the rTMS IPL phase and $4.81 \pm 1.61\%$ for

the rTMS POz phase in the Intervention group and $5.73 \pm 3.44\%$ for the rTMS phase in the Control

group).

We reasoned that if incidental rule-learning occurs over exposure in the rule block, participants'

gradual ability to predict the appearance or non-appearance of a target word C_j on the basis of the

identity of the initial word A_j should be reflected in a RT gain (i.e., faster RTs) over trials within the

blocks (see section Linear Mixed Models analysis). We also expected an overall RT advantage over

target words in the no-rule blocks (*rule effect*), where prediction is possible (no prediction can be

made during no-rule blocks). Participants' *rule effect* for the different parts/sessions was calculated

as the mean RT difference between no-rule and rule trials.

Linear Mixed Models analysis

In order to assess online rule learning in the Rule Learning Task within each experimental session,

we used a linear mixed model approach to fit learning slopes that reflect reaction time gains over

trials for the rule and no-rule conditions. The use of mixed models to compare the slope between

conditions allows the use of RT data for all trials and subjects, which results in a more sensitive

measure of the online learning process than a single mean value per subject[46,68]. Analyses were

performed using the *lme4*[69] and lmerTest[70] packages as implemented in the R statistical

language (R Core Team, 2012). Our basic model included reaction time (*rt*) and *trial* as continuous

variables, no-rule/rule (*NR_R*) and target/no target (*TNT*) as 2-level factors, and *subject* as a factor

with as many levels as participants.

$rt \sim NR\_R + trial + NR\_R*trial + TNT + (1|subject)$ (1)

$rt = ß0 \ (intercept) + ß1(NR\_R) + ß2(trial) + ß3(TNT) + ß4(NR\_R*trial)$ (2)

$rt \sim TNT + (1+NR\_R*trial|suje)$ (3)

As detailed in (1), which shows the specified model, *NR_R*, *trial* and their interaction were

introduced as fixed effects terms. *TNT* was also included here albeit as a predictor of no interest. To

allow for a different intercept per participant and so account for inter-subject variability in basal

response speed, *subject* was introduced as a random effect. The algebraic expression of the fixed

effects part of the model is given in (2). Note that, in this this model, ß4 (*NR_R*trial*) represents an

estimate of the difference in learning slopes between the rule and no-rule conditions and can thus be

used as a de-trended learning slope estimate for the Rule condition. For the sake of clarity, we have

referred to this estimate as ß$_{diff}$. A statistically significant negative ß$_{diff}$ value therefore indicates that

online rule learning effectively took place in that experimental phase over and beyond any reaction

time gain that may be attributed to within-phase practice effects. Individual slopes were estimated

for correlations via a random slopes model specifying the interaction term in the random effects part

(3).

Attention task

In addition to the Rule Learning Task, during Session 2, participants in the Intervention group also

performed an attention task with the aim of better characterizing the impact of rTMS lPL and rTMS

POz. Note that all tasks were completed within 30 minutes in order to ensure post rTMS

performance under the estimated rTMS effects for both tasks. Half of the subjects performed the

language task first and then the Attention Task while the other half took the reverse order under

each of the rTMS interventions (see main text Fig 1).

In the Attention Task (main text Fig. 6), participants were asked to make a pitch discrimination

judgment (higher or lower) on an isolated syllable that followed a sequence of alternating syllables.

A pool of 200 ms-long syllables at three different pitch heights (249, 440 and 554 Hz) was created

from four original tokens ('ba', 'co', 'pi' and 'te') previously recorded by a female native Spanish

speaker. All syllables were normalized for homogeneous output volume using ADOBE AUDITION

(Adobe Systems Software Ireland Ltd.). Variants at 440 Hz were used to build sequences of six

alternating stimuli consisting of either the syllables '/ba' and '/pi' or the syllables '/co' and '/te'. A

higher (554 Hz) or lower pitch (440 Hz) syllable was then attached 800 ms following the end of

each sequence and represented the target syllable upon which the pitch discrimination was to be

judged. Sequences could be either informative, the first syllable matching the target syllable in

identity, or non-informative, the identity of the target syllable could not be anticipated (50%

probability) from the first syllable. The time interval between the syllables of a given sequence was

also manipulated to be either constant (fixed at 400 ms; rhythmic condition) or pseudo-random

(ranging between 100-700 ms; mean 400 ms; non-rhythmic condition). In both cases the duration of

a trial was the same. The overall design, therefore, conformed to a standard 2x2 orthogonal design

with factors Identity (informative/non-informative) and Rhythm (rhythmic/non-rhythmic).

Participants were instructed to judge whether the target syllable was either higher or lower in pitch

compared to the preceding syllables in the sequence, responding as fast as possible while avoiding

errors. Participants were also informed about the Identity factor and they were asked to focus on the

identity of the first syllable to anticipate target appearance. No reference to sequence rhythm was

made. Each trial began with a fixation cross of variable duration (500-1000 ms) followed by the

syllable sequence and target syllable. A maximum of 2000 ms was allowed for response after target

presentation. An inter-trial interval of 1900 ms followed responses, after which the next trial started.

Participants ran a set of 16 practice trials (with only rhythmic sequences) in which speed and

accuracy feedback was given after each trial. This was included in order to ensure proper

understanding of the task. A familiarization session was then completed, consisting of three blocks

of 32 sequences each. The Attention Task was completed twice, after rTMS lPL and rTMS POz.

Each run consisted of two blocks of 32 sequences each. Experimental conditions were manipulated

within blocks, pseudo-randomly. Only correct responses were included for the analysis. Outlier

trials per condition on the basis of a mean ± 2.5 *sd* criterion were also removed from further

analysis (rTMS POz: 6.7% of trials; rTMS lPL: 6.3% of trials). A repeated measures ANOVA was

used to assess the effects of the factors [within subject factors: Intervention (rTMS lPL, rTMS

POz), Rhythm (rhythmic, non-rhythmic), Identity (informative/non-informative); and between

subject factor: Intervention Order]. An ANOVA for each intervention type was performed, with

interactions therein explored via paired-samples *t*-tests.

fMRI acquisition and apparatus

Functional T2*-weighted images were collected using a General Electric MRI 3T scanner and a

gradient echo-planar imaging sequence to measure blood oxygenation level dependent (BOLD)

contrast over the whole brain [repetition time (TR) 2000 ms, echo time (TE) 30 ms; 35 slices

acquired in ascending interleaved order; slice-thickness: 3 mm with a 0.3 mm gap, 64×64 matrix,

in plane resolution = 3×3 mm; flip angle, 90 º]. Two main fMRI runs with 325 images were

acquired for the rule-learning task. Structural images were collected using the MPRAGE-sequence

with the following parameters: TR = 7.3 ms, TE/-TI = 2.6, flip angle = 90°, FOV = $256 \times 256 \times 160$

mm, spatial resolution = $1mm^3$/voxel. Auditory stimuli were presented using an amplifier

(Sensimetrics[TM], Malden, MA, USA) attached to the earphones, and visual stimuli were presented

using MRI-compatible goggles.

fMRI preprocessing and regression analysis

Data were preprocessed using Statistical Parameter Mapping software (SPM8, Wellcome Trust

Centre for Neuroimaging, University College, London, UK, www.fil.ion.ucl.ac.uk/spm/). The two

functional runs were first realigned and a mean image of all the EPIs was created. This mean image

was then co-registered to the T1 image, which was then segmented into grey and white matter by

means of the Unified Segmentation algorithm[71]. The resulting normalization parameters were

applied to normalize the whole functional set to the MNI space. Finally, functional EPI volumes

were re-sampled into $2\times2\times2$ mm voxels and spatially smoothed with an 8 mm FWHM kernel.

An event-related design matrix was specified using the canonical hemodynamic response function.

Trial onsets were modeled at the moment of the presentation of the first word in a sentence. Two

main conditions were modeled: rule and no-rule. A temporal 1st order modulator and a parametric

modulator (we included, for each trial, its reaction time) were added to each condition. Data were

high-pass filtered (to a maximum of 1/90 Hz) and serial autocorrelations were estimated using an

autoregressive (AR(1)) model. Remaining motion effects were minimized by also including the

estimated movement parameters in the model. First-level contrasts were specified for all

participants for the main conditions against the implicit baseline. In addition a rule vs. no-rule

contrast was also calculated at the individual level. First level contrast images were fed to two

second level one sample models with one covariate each to calculate correlations between brain

activity and behavior: rule vs implicit baseline contrasts were correlated with individual learning

slopes and rule vs no-rule contrasts were correlated with the increment in the rule effect in Part 2

over Part 1.

Group results are reported at a FWE $p < 0.05$ corrected threshold at the cluster level with 50 voxels

of cluster extent, with an additional uncorrected $p < 0.005$ threshold at the voxel level. Anatomical

and cytoarchitectonical areas were identified using the Automated Anatomical Labelling Atlas[72]

included in the xjView toolbox (http://www.alivelearn.net/xjview8/).

Repetitive TMS protocol

Transcranial magnetic stimulation was delivered through a figure eight coil attached to a standard

MagStim Rapid 2 stimulator (Maximum stimulation output 2.2T). The rTMS was delivered off-line,

15 minutes before the Rule Learning and the Attention tasks, at a frequency of 1Hz and an intensity

of the 60% of the maximum stimulation output. This protocol is known to decrease the excitability

in the cortical regions beneath the coil position[73] for an average duration of 30 minutes[74] after

the end of the stimulation period.

Participants performed two rTMS conditions on the same day (Session 2) corresponding to: i) the

target stimulation site (obtained from the peak activation of the BOLD signal on the lPL; rTMS lPL

phase); and ii) the control stimulation site (POz electrode location according to the 10/20 EEG

international system; rTMS POz phase). rTMS was applied within an average of 3 mm from that

peak.

The Control group did not receive rTMS but followed the same behavioral protocol. Since no rTMS

was applied, data from L2 and L3 sessions were merged for this group.

When applied onto the lPL, the handle of the coil was angled at 45° away from the midline. In

contrast, the coil was placed in a vertical position (with the coil handle pointing backwards) for

stimulation at POz. Stimulation sites were identified on subject's scalp using the SofTaxic navigator

system (EMS, Italy) according to the T1 image and a marker of the individual target location, using

the coordinates of the functional localizer (as described above). Mean stimulation MNI coordinates

located at the lPL were x = -47.85 ± 6.91; y = -39.6 ± 8.62; z = 48.8 ± 5.52 (see Table S3 for

individual coordinates). SofTaxic navigator system (EMS Italy) was also used for coil maintenance

during the rTMS application (allowed error < 2 mm in 3D space). Due to an Experimenter error,

two subjects were stimulated beyond 3 mm from the peak of activation and were thus excluded

from rTMS analyses.

Author Contributions

R. d. D.-B., J. L. A and A. C. designed the research. J. O.**,** P. R., M. R., J. L. A., A. C., S. S.-F., and

R. d. D.-B. wrote the manuscript. M. R., J. L. A., A. M. acquired the data. J.O.**,** P. R. analyzed the

data.

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Supplementary Information Captions

Figure S1. Control group incidental rule-learning task results for Session 1. Slopes for rule and no-rule blocks over task repetitions derived from the mixed model analysis. The Control Group showed the expected transition from a significant learning slope in Part 1 (ßdiff = -0.8, t = -3.1, $p < 0.002$) to a non-significant learning slope (ßdiff = 0.06, t = 0.252, $p > 0.8$) with a significant rule effect in Part 2 ($t(30)$ = 4.49, $p < 0.001$) and Part 1 ($t(29)$ = 3.6, $p < 0.002$). Actual data shown averaged into 6 trial bins (for visual purpuses only; the anlaysis did not bin the data) with the SEM, over the slopes for rule and no-rule derived from the mixed model analysis.

Figure S2. Intervention group incidental rule-learning task results for Session 1. Slopes for rule and no-rule blocks over task repetitions derived from the mixed model analysis. the Intervention Group –perhaps owing to the smaller sample and/or scanner effects– appeared to comprise slower learners who still showed a significant learning slope during fMRI phase (Part 2: ßdiff = -0.8, t = -2.63, $p < 0.01$; Part 1: ßdiff = -0.56, t = -2.13, $p < 0.034$), as well as the expected rule effects (Part 1: $t(16)$ = 2.19, $p < 0.044$; Part 2: $t(19)$ = 3.08, $p < 0.007$). Actual data shown averaged into 6 trial bins (for visual purpuses only; the anlaysis did not bin the data) with the SEM, over the slopes for rule and no-rule derived from the mixed model analysis.

Figure S3. Individual fMRI enhanced activity during later stages of rule learning and rTMS stimulation sites. In red-yellow, overlap of individual masks for each participant's activation pattern. Only voxels in which at least 10 participants showed individual fMRI enhanced activity during Rule learning are shown. In blue, the sites for rTMS stimulation for all participants is shown (for clarity purposes, 4 mm spheres were created around the stimulation centers). Neurological convention is used with MNI coordinates shown at the bottom right of each slice.

Fig. S4. Overlap (yellow) between the ventral network correlating with Statistical Learning (green) and the main rule effect (Rule vs. Baseline; in green). In the lPL, there is an overlap between the contrast showing regions in which the BOLD signal significantly co-varies with the measure of statistical learning (Learning Slope) and the contrast showing the brain regions in which activity increases during Rule blocks: the more activity in the lPL during rule blocks, the greater (i.e. more negative) the slope and the faster the statistical learning occurs. Only significant results ($p < 0.05$ FWE-corrected at the cluster level, with an additional $p < 0.001$ at the voxel level and 50 voxels of cluster extent) are shown over a canonical template with MNI coordinates on the bottom right of each slice. L, Left Hemisphere.

Table S1. Whole brain fMRI activity related to individual differences in rule slopes. Group-level fMRI local maxima for areas correlating with negative (i.e. RTs are still decreasing) rule slopes during the rule blocks in the fMRI phase (see also red-yellow regions in **Fig. 4A**, main text). Results are reported at a FWE $p < 0.05$ corrected threshold at the cluster level with 50 voxels of cluster extent, with an additional uncorrected $p < 0.005$ threshold at the voxel level. MNI coordinates were used. BA, Brodmann Area.

Table S2. Whole brain fMRI activity related to individual differences in the rule effect increment for Part 2. Group-level fMRI local maxima for areas correlating with rule effect increments in Part 2 for the rule vs. no-rule contrast during the fMRI phase (see also red-yellow regions in **Fig. 4B**). Results are reported at a FWE $p < 0.05$ corrected threshold at the cluster level with 50 voxels of cluster extent, with an additional uncorrected $p < 0.005$ threshold at the voxel level. MNI coordinates were used. BA, Brodmann Area.

Table S3. Coordinates for parietal stimulation sites

Table S4. Results for the d' calculated with the order or the dependency as false alarms

Table S5. Stimuli set for the rule-learning task