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Main Manuscript for

Action planning modulates perceptual confidence through action monitoring processes

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modulates perceptual confidence, *International Conference of Cognitive Neuroscience*

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insights from EEG signals, *Association for the Scientific Study of Consciousness 2022*,

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Abstract

Dominant models of metacognition suggest that sensory information quality determines

perceptual confidence, but recent accounts propose that motor signals also affect

confidence judgments. In this study, we investigated the impact of motor planning of

perceptual responses on decision confidence, testing two hypotheses. The "fluency

hypothesis" suggests that ease of motor response selection and preparation enhances

confidence. In contrast, the "monitoring hypothesis" posits that increased action

monitoring during response selection boosts confidence, potentially counteracting

response fluency. In three pre-registered experiments, participants reported the orientation

of a stimulus and indicated their confidence in their response. A cue induced action

planning that was either congruent or incongruent with the response side used to report the stimulus orientation. Across experiments, we consistently observed higher confidence when participants prepared spatially incongruent actions compared to congruent ones, regardless of response accuracy. In the third experiment, electroencephalography (EEG) revealed an increased fronto-central P2 amplitude for incongruent actions, suggesting that incongruent action planning heightened early attentional resources needed to resolve response conflict. Incongruent action plans also modulated post-response ERPs at centro-parietal channels (e.g., Pz), typically linked to confidence and error monitoring. These findings align with the "monitoring hypothesis" suggesting that the degree of action monitoring during response selection modulates retrospective decision confidence.

Keywords: confidence, decision-making, action monitoring, action fluency, sense of control, EEG

Public Significance Statement

While virtually every decision we make leads to an action, the role of motor processes in decision making has been largely neglected. Our results show that retrospective confidence in a perceptual discrimination task is boosted when the motor execution is spatially incongruent with motor preparation, independently of the correctness of the response. Electroencephalography recordings indicate that this effect could be explained by a larger involvement of early attentional resources related to action monitoring, which has an impact on confidence computations. Taken together, these results suggest that action

75 planning information might trigger monitoring mechanisms susceptible to alter

76 retrospective confidence in our decisions, implying that motor processes are not only the

77 output, but also an input of the decision mechanisms.

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Introduction

In a world where feedback is sparse and uncertain, adaptive behavior crucially relies

on our ability to estimate how confident we are in the accuracy of our perceptual

representations and decisions (Mamassian, 2016). Research showed that the sense of

confidence plays a prominent role in learning (Guggenmos et al., 2016), information

seeking (Desender et al. 2018), change of mind (Rollwage et al. 2020; Sanchez et al. 2024),

and social interactions (Bahrami et al. 2010). Delineating the factors influencing the

formation of confidence is therefore crucial for the understanding of adaptive behavior

(Rahnev et al., 2021).

Perceptual confidence is typically measured and dissociated from accuracy by asking

participants to provide two types of responses during a perceptual decision task. First, they

make a perceptual decision about a stimulus, and then they rate their confidence in how

accurate that decision was, using a scale (cf. Mamassian, 2016). Classical models of

metacognition argue that, just like accuracy, perceptual confidence is determined by the

quality of sensory information -i.e., confidence increases with stronger perceptual evidence

(Kiani & Shadlen, 2009). In other words, confidence and accuracy are two different

descriptions of the same internal variable. However, recent studies suggest that the

characteristics of the action used to bring about a perceptual decision can also shape

perceptual confidence (Fleming et al., 2015; Gajdos et al., 2019; Pereira et al., 2020;

Siedlecka, Koculak, & Paulewicz, 2021; Siedlecka, Paulewicz, & Koculak, 2020; Turner

et al., 2021; Wokke, Achoui, & Cleeremans, 2020). Hence, besides sensory evidence,

motor signals can modulate confidence judgments. However, it remains unclear what

specific aspect of motor processing (i.e., motor preparation or motor execution processes)

is responsible for this modulation, and how this modulation takes place.

Two alternative hypotheses, each providing contradictory predictions, can be proposed

regarding how motor information is used for confidence estimations. The "fluency

hypothesis" suggests that the ease and smoothness in the selection and execution of a motor

response, may serve as a cue for confidence judgments (Fleming et al., 2015; see also

Wenke et al., 2010; Oppenheimer, 2008; Brouillet et al., 2022; 2023; for notions related to

fluency and action fluency). Specifically, fluent actions may lead to higher confidence

judgments compared to non-fluent actions. This hypothesis is supported by a study

conducted by Fleming et al. (2015), where transcranial magnetic stimulation (TMS) was

used to manipulate motor representations in the premotor cortex. The authors found that

stimulating premotor areas linked to the chosen perceptual response increased confidence.

They hypothesized that this stimulation enhanced action fluency, which, in turn, boosted

confidence judgments. The notion of fluency aligns also with the observation that, other

things being equal, faster responses are associated with higher confidence reports (Kiani et

al., 2014). In sum, the idea underlying this hypothesis is that confidence judgments rely in

part on heuristics employing proxies of response accuracy (Ackerman, 2019; Maniscalco

et al., 2016; Turner et al., 2021; Van Marcke et al., 2022), and action fluency would be one

of these proxies. Alternatively, according to the "monitoring hypothesis", the modulation

of action signals on confidence occurs through the involvement of action monitoring

processes (cf. Anzulewicz et al., 2019; Gajdos et al., 2019; Sanchez et al., 2024). Contrary

to the fluency hypothesis, this perspective suggests that it is not the fluency of an action

that leads to higher confidence estimates, but rather the degree of control exerted and the

level of monitoring required for implementing a perceptual decision. The higher the control

required for implementing a response, the higher the confidence in the perceptual decision.

This hypothesis is supported by neuroimaging research highlighting the critical role of the

Prefrontal Cortex (PFC) in both perceptual confidence and in the monitoring of self-

generated decisions (Fleming et al., 2012), as well as by more recent studies. For instance,

Turner et al. (2021) showed that effortful perceptual responses are associated with higher

confidence than less effortful motor decisions (but see Hagura et al. 2023). Similarly,

Sanchez et al. (2024) showed that enhanced visuo-motor control towards perceptual

decisions increase confidence judgments. Finally, Gajdos et al. (2019) observed that when

undesired motor responses are correctly inhibited perceptual confidence increases. Overall,

these studies suggest that when action monitoring processes are more strongly involved -

either because actions are more effortful or unwanted responses need to be controlled and

inhibited (Anzulewicz et al., 2019; Gajdos et al., 2019; Morel et al., 2017; Sanchez et al.,

2024; Turner et al., 2021)- confidence increases, even if action fluency is hindered.

To confront these two hypotheses and test their predictions, we conducted three pre-

registered experiments which aimed specifically at investigating how motor preparatory

signals of perceptual responses are used in confidence estimations. Specifically,

participants reported the orientation of a Gabor (vertical or horizontal) with their index

fingers and judged their confidence in their decision. Using a novel motor priming

paradigm, we prompted participants to prepare an action in advance that could either be

compatible or incompatible with the action ultimately chosen to report the Gabor

orientation.

The "fluency hypothesis" would predict that perceptual confidence increases when the

perceptual response is compatible with the primed action. Indeed, priming a compatible

aspect of the perceptual response should facilitate the planning and selection of that

response, thereby increasing fluency (e.g., Chambon & Haggard 2012). Thus, higher

confidence is expected when participants prepare a congruent action (e.g., a left response

is primed and the Gabor is reported with a left response) compared to incongruent actions

(e.g., a left response is primed but the Gabor is reported with a right response). In contrast,

the "monitoring hypothesis" would predict the opposite: perceptual confidence is higher

when the primed action is incompatible with the subsequent perceptual response. In

incompatible trials, increased monitoring is required to control and inhibit unwanted motor

plans. Accordingly, higher confidence may be expected when participants prepare an

incongruent action compared to congruent actions.

In Experiment 3, we used electroencephalography (EEG) to investigate the specific

brain processes modulating confidence, focusing in particular on the frontocentral P2/N2

complex. The P2 component is linked to attentional resource allocation for feature-based

stimulus evaluation and response selection (Darriba et al., 2018; Xie et al., 2020), while

the N2 is associated with motor inhibition and it is typically observed in go/no-go tasks

(Folstein et al., 2008). We predicted that priming incompatible responses would require

higher monitoring and control, leading to increased P2 and N2 components.

In a nutshell, we found that confidence was higher when participants prepared incongruent rather than congruent actions, supporting the monitoring hypothesis over the fluency hypothesis: the more individuals monitor and control their motor performance, the higher their perceptual confidence. This finding is important as it suggests that action processes are not merely auxiliary to decision-making; instead, they shape high-level cognitive constructs, such as retrospective confidence.

Materials and Methods

Materials.

Participants. For Experiment 1, data collection was stopped after gathering the data from 16 participants that did not violate our data inclusion criteria. The inclusion criteria required participants to achieve perceptual performance levels suitable for the study of confidence (i.e., an accuracy rate between 60% and 90%), and to use each level of confidence at least once, indicating a proper understanding of the task (for a complete description of the inclusion criteria, see Supplementary Information -SI-, section *Experimental Procedure*). Crucially, we verified that our results remained consistent with both the inclusion and exclusion of outlier participants (SI *Table S6, S7 & S8*). This sample size was estimated by performing a statistical power analysis (using G*Power; Faul et al., 2007), based on a pilot study that investigated the effect of our motor priming paradigm on

response times (see OSF pre-registration: MG1). Ten participants were not included in the sample since they did not fulfil our inclusion criteria. The remaining 16 participants (11 participants reported their gender as female, 5 as male, mean age = 24.9, SD = 4.18) were analyzed. Based on the results of the first experiment (see OSF pre-registration: MG2) and on the decision to increase statistical power, the data collection of Experiment 2 was stopped after gathering the data from 24 participants. In total 32 adults were recruited for this experiment, 8 of them were not included in the sample since they did not fulfil our inclusion criteria. The remaining 24 participants (14 participants reported their gender as female, 10 as male, average age = 25.8, SD = 3.68) were analyzed. For Experiment 3, we replicated the procedure of Experiment 2 (see OSF pre-registration: MG3) and no participant was excluded, hence 24 participants (12 participants reported their gender as female, 12 as male, average age = 25.4, SD = 4.36) were recruited for this experiment and included in the analyses. Participants were paid 10€/hour in gift card (and 15€/hour for Experiment 3 since it involved EEG). All participants had normal or corrected-to-normal vision, were right-handed, and were naïve regarding the hypothesis under investigation. They all gave written and informed consent before participating in the experiment. This study was conducted in agreement with the requirements of the Declaration of Helsinki and approved by the ethics committee of Université Paris Cité (Number: 00012024-51).

Equipment. Participants performed the experiment in a dark room. They were comfortably seated in front of an LCD monitor (in Experiments 1 and 2 we used a 20-inch Asus PG248Q with 1920 by 1080 screen resolution and a refresh rate of 60 Hz, in Experiment 3 we used Display ++ with a resolution of 1920 x 1080 pixels, a size of 32 inches, and refresh rate of

60 Hz). The experiment was programmed with Python (Python Software Foundation.

Python Language Reference, version 3.9. Available at http://www.python.org) and the

stimuli were generated using the Psychopy software (Peirce et al., 2019). A button response

box (Millikey Response Box MH-5) and foot pedals (Accuratus X3P Footswitch) were

used to collect participants' hand and feet responses, respectively. For Experiment 3, two

thumb-press buttons were specifically built for convenience (See SI, *Figure S1*). EEG

signals were recorded with 64 Ag/AgC1 electrodes mounted on an elastic cap and

amplified by an ActiCHamp Plus amplifier (Brain Products GmbH, Munich, Germany).

Electrodes were arranged according to the international 10–20 systems. EEG data were

online referenced to M2. EEG signal was sampled at a digitization rate of 500 Hz.

Methods.

Experimental paradigm. In the three experiments, we manipulated the motor preparation

of perceptual responses, by randomly interleaving the trials of a perceptual task with the

trials of a speeded Reaction Time (RT) task. In each trial (of both the perceptual and the

speeded RT task) a visual cue was displayed (e.g., an arrow in Experiment 2 and 3, see

below). The visual cue indicated the action to perform (e.g., left or right button press) in

the speeded RT task. This action had to be executed as soon as a white flash was presented.

However, in part of the trials, participants rather than viewing a white flash, were presented

with a Gabor at perceptual discrimination threshold. Thresholds were calculated for each

participant in a preliminary calibration phase (see Calibration Phase in the Supplementary

Material). In those trials, they were instructed to ignore the speeded RT task and to perform

instead a perceptual task, which consisted in indicating the orientation of the Gabor.

Crucially, the task to perform at each trial was not known at the time of the cue onset, only

the presentation of the white-flash or the Gabor indicated the participant which task to

perform. We expected that the presentation of the visual cue would push the participants

to prepare in advance a response in order to meet the temporal constraint imposed by the

speeded RT task. The action preparation induced by the cue would then facilitate (or

interfere with) the selection/execution of a compatible (or incompatible) action required to

report the orientation of the Gabor.

Experiment 1

Each trial started with a presentation of a green fixation cross (0.4 by 0.4 degrees of visual

angle, dva) at the center of the screen for a duration of 300 ms (see Figure 1). A visual cue

was then displayed for 200 ms. The cue was the same fixation cross but presented with a

different color (e.g., blue or yellow). In the speeded RT task trials participants were

instructed to report the color of the cue by executing a left or a right response as soon as a

white flash (1.8 dva diameter) was displayed (e.g., left-response if the fixation turned blue

and right-response if the fixation turned yellow). In half of the trials, participants reported

the color of the fixation with their index fingers (the same pair of effectors used in the

perceptual task described below, i.e., same effector block), while in the other half of the

trials, they reported the color by using their feet (different effector block). The white flash

was presented 800 ms after the onset of the cue and displayed for a duration of 50 ms.

Participants had 800 ms to respond to the cue. If no response was given within this delay,

the trial was interrupted and a message 'too late!' was displayed. The action-color mapping

and the order of the blocks (same and different effector) were counterbalanced across

participants.

Experiment 1   Experiment 2 and Experiment 3 (EEG)

Figure 1. Experimental paradigms. Participants are initially presented with a visual cue (i.e., a

colored fixation in Experiment 1, and an arrow in Experiments 2 and 3). Subsequently, they view

either a white flash or a Gabor patch. If a white flash is displayed, participants have to report the

color of the visual cue (Experiment 1) or its pointing direction (Experiment 2 and 3) by pressing a

left or a right button (speeded RT task). If a Gabor is presented, they have to report its orientation

by executing a left or a right button-press (e.g., left for horizontal and right for vertical; perceptual

task). After their perceptual decision, participants report their confidence on a scale from 1 to 4.

For Experiment 2 and 3, participants could also report an error instead of a confidence level.

In one third of the trials, rather than viewing a white flash, participants were presented with

a noise texture (1.8 dva diameter) for 50 ms. A Gabor patch (1.8 dva diameter) was then

displayed 800 ms after the onset of the noise stimulus, for a duration of 50 ms. This delay

of 800 ms was introduced in Experiment 1 to help participants to prepare for reporting the

orientation of the Gabor. In these trials, participants were instructed to forget about the

speeded RT task and were asked to perform a **perceptual task** instead. Importantly,

perceptual and speeded RT trials were randomly interleaved. The perceptual task consisted

in discriminating the orientation (horizontal or vertical) of the Gabor patch (see Figure 1).

The Gabor was embedded in a noisy texture and displayed with a contrast level supporting

72.5% correct discrimination performance calculated separately for each participant in a

preliminary calibration phase (see SI *Experimental procedure*). Perceptual responses were

provided by executing a left or right index finger key-press as quickly and as accurately as

possible (Gabor-response mapping was counterbalanced across participants). Participants

had 800 ms to respond. If no response was given within this delay, the trial was interrupted

with an error message and added to the trial list.

Critically, the action performed to report the orientation of the Gabor could be

compatible or incompatible with the action prepared to report the visual cue. In particular,

we manipulated two sorts of compatibility. Firstly, the action prepared to report the cue

could be spatially congruent or incongruent with the response selected to report the

orientation of the Gabor. For instance, in spatially incongruent trials, the cue induced the

preparation of a left (index finger or foot) response while the participant performed a right

(index finger) response to report the Gabor. Secondly, the action prepared to report the cue

could involve the same (index fingers) or a different pair of effectors (feet) than those used

by the participant to report the Gabor. Hence, the experiment implemented a 2 by 2 factorial

design with *Spatial congruency* (congruent and incongruent) and *Effector compatibility*

(same and different).

After the perceptual response, participants reported their confidence on a scale from

1 to 4 corresponding to the probability that their perceptual decision was correct (1:

guessing, 2: barely confident, 3: fairly confident, 4: certainly correct). The confidence level

numbers were displayed inside four circles randomly displayed around the center of the

screen (Figure 1). There was no time pressure to report confidence ratings. To indicate the

level of confidence at each trial, since participants used four effectors in the experiment,

each of the four effectors were associated with a confidence circle.

300   The experiment lasted about one hour and thirty minutes and was divided into two

main blocks (same and different effector blocks). Each block was composed of 11 mini-

blocks of 36 trials each (24 speeded RT task trials and 12 perceptual task trials) for a total

of 792 trials evenly split into 22 blocks. Before the main experiment, a training phase and

a calibration phase were performed (see SI *Experimental procedure*).

Experiment 2

For Experiment 2, we made a few changes to the design of Experiment 1. In order to make

the speeded RT task easier and more intuitive, the visual cue was a white arrow (1 dva by

1 dva) displayed at the center of the screen pointing either to the left or to the right.

Participants were instructed to indicate its direction with a left or right response

respectively (Figure 1). Furthermore, in the *different effector* block we replaced left and

right foot responses with left and right middle finger button-presses. We changed the

effectors because in Experiment 1 we observed a clear deterioration of performances when

switching from lower- to upper-limb actions. Since the speeded RT task was easier, it

enabled us to bring the onset of the Gabor patch closer to the cue. Hence, we removed the

warning noise texture presented before the occurrence of the Gabor, which was now

displayed 800 ms after the cue (thus, respecting the timing of the presentation of the visual-

flash in the speeded RT task).

We also increased the response time limit of the perceptual task to 1500 ms. Additionally,

in case participants made an involuntary mistake in the perceptual task (e.g., they executed

a right key-press but they wanted to perform a left key-press), they could report it during

the confidence judgment by pressing the space bar (Figure 1).

The experiment lasted ~45 minutes and was divided into two main blocks (same effector

and different effector). Each block was composed of 5 mini-blocks of 40 trials each (20

cue and 20 Gabor trials) for a total of 400 trials evenly split into 10 blocks.

Experiment 3

In Experiment 3, the design was identical to Experiment 2, except for the following

changes. Since effector compatibility did not seem to impact confidence, we removed the

different effector condition, so the speeded RT task and the perceptual task were performed

with the thumbs. Participants indicated their confidence by moving an arrow pointing

towards one of the confidence levels. Participants could move the arrow either clockwise

(right hand) or anti-clockwise (left hand) and validate their confidence level with a double

button-press. To report an error, participants had to validate when the arrow was pointing

toward the box 'E' (Figure 1). Additionally, the confidence display was presented 1.25

seconds after the perceptual response to leave some time without visual stimulation for

EEG considerations. The experiment was divided into 10 mini-blocks of 40 trials each (20

cue and 20 Gabor trials) for a total of 400 trials lasting about 45 minutes.

Data analyses

Behavioral data. Data were analyzed using R (R Core Team, 2020). We used generalized

linear mixed-effect models (inverse-gaussian with identity log function) for Response

Times (RT), generalized linear mixed-effects models (binomial with logit link function)

for response accuracy, and ordinal regression with cumulative link model for confidence.

Linear and generalized linear regressions were performed with the package lme4 (Bates et

al., 2015) and ordinal regressions with the package ordinal (Christensen et al., 2018).

Linear regressions were performed with the restricted maximum likelihood fitting method,

and *p*-values for coefficients were computed with Satterthwaite's method using the

lmerTest package (Kuznetsova et al., 2017). Ordinal regressions were performed with

Laplace approximation, and *p*-values were computed for Wald tests. The models were

adjusted to accommodate convergence or singularity issues. The structure of random

effects was determined based on the parsimony principle (Bates et al., 2015). Notably, for

some analyses, the inclusion of the maximal random structure led to convergence failures.

In those cases, we performed a Principal Component Analysis (PCA) to isolate the random

effects that the least contributed to model fitting and we removed them one-by-one from

the final model, until there were no convergence failures. We used an alpha level of 0.05

for all statistical tests. Factors were coded using sum contrasts, and response times were

centered (with respect to the across-subjects mean). In addition, our results were replicated

with the inclusion of outliers (see SI *Tables S6, S7 & S8*), and they were also replicated

using Bayesian methods (see SI *Bayesian analyses, Tables S3, S4 & S5*). For graphical

representations, the package ggplot2 was used (Wickham, 2009).

In Experiment 2 and 3, participants were able to report errors they made consciously

when indicating the orientation of the Gabor. These trials were omitted from further

analysis. In Experiment 2 they represented 6.0 % of the total number of trials (i.e., 290

errors over 4800 trials in total). Specifically, participants reported 151 errors in the different

effector condition (44 in the incongruent and 107 in the congruent condition) and 139 in

the same effector condition (52 in the incongruent and 87 in the congruent condition). In

Experiment 3 they represented 5.0 % of the trials (i.e., 230 error over 4800 trials in total).

Specifically, participants reported 133 errors for congruent and 74 for incongruent trials.

EEG Data. EEG data were analyzed using MNE-Python 1.3.0 (Gramfort et al., 2013,

2014) and re-referenced to the left and right mastoids (M1 and M2). The signal was band-

pass filtered between 0.05 and 48 Hz by a non-causal infinite impulse response filter. The

raw data was inspected visually to remove time periods containing large artifacts.

Subsequently, we performed an independent component analysis (Hyvärinen, 1999), to

identify and remove components representing blinks or eye movements. We then extracted

epochs going from -1900 ms to 2500 ms time-locked to Gabor onset. A second artifact

rejection was performed. Epochs containing amplitudes greater than 100 µV or less than -

100 µV were marked as potential artifacts and then removed after confirmation through

visual inspection. This led to the removal of 224 out of 4628 trials, i.e., a proportion of

0.048 trials. EEG activity in specific time windows for ERP components was quantified

through mean amplitudes calculation and tested with ANOVA and t-tests as they are robust

to different numbers of trials across conditions (Luck, 2012). If no specific time windows

were identified from the literature for a component, we performed a nonparametric cluster-

based permutation test (Maris and Oostenveld, 2007) to isolate the time windows

exhibiting a difference between congruent and incongruent trials.

Firstly, we performed a temporal cluster-based permutation test comparing the

voltage amplitude observed in congruent and incongruent trials on a time window going

from 0 to 350 ms post-stimulus (we selected a time window in which no perceptual

response could have already occurred) at electrode Fz, Cz and FCz. Stimulus-locked

segments were corrected with a baseline of 200 ms prior to Gabor onset (Folstein & Van

Petten, 2008; Luck et al., 2012). This analysis was conducted to identify the time window

and potential ERP component (e.g., P2 or N2) that differed between congruent and

incongruent trials.

Secondly, we analyzed post-response ERPs. Response-locked epochs were

corrected with a 100 ms baseline, going from -100 to 0 ms prior to the response (Boldt &

Yeung., 2015). We focused and analyzed a time period going from 0-500 ms post-response

at the electrode Pz, given that this time window and electrode were already identified in

previous study of EEG correlates of confidence (Davies et al., 2001; Wang et al., 2020;

Rausch et al., 2020). We then performed a cluster-based permutation tests on the difference

between congruent and incongruent between 0 ms to 500 ms post-response at Pz to identify

potential components exhibiting a modulation of spatial congruency. Two temporal

clusters were identified (see Results). To investigate their relation with confidence

judgments we binned the four levels of confidence into low (confidence ratings equal 1 or

2) and high (confidence ratings equal 3 or 4) trials and performed ANOVAs on the by-

participant average amplitude observed in the two time periods reported above. The

ANOVA included Confidence (high, low) and Congruency (congruent, incongruent) as

factors. Note that we also analyzed the amplitude of post-stimulus frontal theta oscillations.

These analyses did not yield conclusive results and are reported and discussed in the

supplementary materials.

Furthermore, we analyzed the Lateralized Readiness Potential (LRP), traditionally

associated with motor preparation (Coles, 1989). Our motor priming paradigm aimed to

induce the preparation of a specific action prior to the perceptual response. Hence, we

expected shorter response times for congruent compared to incongruent actions, along with

similar changes in the latency of congruent LRPs. LRPs were calculated using the double

subtraction method, where the average effector-specific activity over the left and right

motor cortices (electrodes C3 and C4) are subtracted (Coles, 1989). Response-locked LRP

epochs (going from -800 to 0 ms time-locked to perceptual response onset, Smulders &

Miller, 2012) were corrected with a baseline going from -1000 and -800 ms prior to the

perceptual response (Kraemer & Gluth, 2023). A jackknife-based method with a one-third

threshold was employed to investigate potential differences in LRP onset latencies between

congruent and incongruent responses, as this method has been shown to be a powerful tool

for analyzing LRP onset latencies (Ulrich & Miller, 2001).

Additional analyses on motor preparatory signals and the P300 were conducted as

sanity checks and are reported in the Supplementary Information. Specifically, we

examined the relationship between confidence and the stimulus-locked P300 component,

given that previous research linked the P300 to evidence accumulation, sensory

uncertainty, and confidence judgments (Herding et al., 2019). Our findings indeed revealed

a correlation between confidence and the P300 (see SI, EEG analyses and results, Figure

S6).

 Transparency and openness

The hypotheses, methods, and statistical procedure for the behavioral data were pre-

registered before the experiment. Pre-registration, data and codes for analysis are publicly

available at https://osf.io/sb45z/. However, the analysis of response time was not included

in the pre-registration. In addition, we made some adjustments to the pre-registered

statistical procedure due to convergence issues with mixed models (see Methods). In

particular, for the analysis of confidence, in Experiment 1, we removed the interaction

between congruency and effector from the random effect structure; and in Experiment 3,

we removed the interaction between congruency and response time from the random effect

structure. For the analysis of accuracy, in Experiment 1, we only kept response time in the

random effect structure; in Experiment 2, we only kept response time and effector in the

random effect structure; and in Experiment 3, we only kept response time in the random

effect structure. Regarding the EEG analyses, only some general aspects were specified in

the pre-registration, as the exact signals and methods of investigation were not fully

determined at the time. We indicated in the pre-registration that we would examine brain

oscillations, such as post-stimulus theta activity (4-7 Hz) in fronto-parietal electrodes, as a

marker of cognitive control (Eisma et al. 2021). Although these analyses were performed,

they are not included in the manuscript because they did not yield clear results. This is

mentioned and discussed in the Supplementary Material.

Results

Behavioral Result

The analyses encompassed a total of 4,224 trials for Experiment 1; 4,510 trials for

Experiment 2; and 4,570 trials for Experiment 3. The average task accuracy for Experiment

1 (M = 0.68, SE = 0.03), Experiment 2 (M = 0.73, SE = 0.01), and Experiment 3 (M = 0.73,

SE = 0.02), and the average confidence ratings for Experiment 1 (M = 2.34, SE = 0.12),

Experiment 2 (M = 2.51, SE = 0.13), and Experiment 3 (M = 2.1, SE = 0.7) were

appropriate for investigating confidence judgments, aligning with previous studies

(Fleming & Lau, 2014; Ferrigno et al. 2019).

Figure 2. Average response times (A), and average accuracy (B) in the perceptual task, as a function

of spatial congruency for each experiment. The graph C depicts perceptual confidence as a function

of perceptual accuracy (C). Error bars represent confidence interval regarding the dispersion of

individual averages corrected by the Cousineau & Morey method (Morey, 2008).

Accuracy

We analyzed accuracy using a binomial generalized linear mixed-effects model, with a

logit link function. Statistical significance was assessed with the Wald test. Formally, we

estimated the following models:

Experiment 1: accuracy ~ congruency * effector * effector_order * RT + (1 + RT | subject)

Experiment 2: accuracy ~ congruency * effector * effector_order * RT + (1 + effector + RT |

subject)

Experiment 3: accuracy ~ congruency * RT + (1 + RT | subject)

Detailed results are gathered in Table S1 in the Supplementary Material. No effect of

spatial congruency on accuracy was observed in Experiment 1 (p = 0.91) and Experiment

2 (p = 0.1). Incongruent responses were slightly more accurate (M = 0.74, se = 0.02) than

congruent responses (M = 0.72, se = 0.02) in Experiment 3 (odds-ratio = 0.92, p = 0.02).

Accuracy was higher in the same effector block compared to the different effector block in

Experiment 1 (odds-ratio = 1.1, p < 0.001), with lower accuracy when the visual cue primed

feet responses (M = 0.64, se = 0.02) compared to hand responses (M = 0.71, se = 0.01),

indicating that the visual task was harder when it required feet responses. This effect was

not observed in Experiment 2 (p = 0.69). No interaction between congruency and effector

(Experiment 1: p = 0.81; Experiment 2: p = 0.92). In Experiment 1, we observed a

significant interaction between effector order and response times on accuracy (odds-ratio

= 0.36, p = 0.003, see SI, *Figure S4*), indicating that only for incorrect responses, response

times were faster when participants completed the same effector block first compared to

when they began with different effector block.

Response time

Response times (RT) were calculated with respect to the onset of the Gabor patch. They

were analyzed with a generalized (inverse-gaussian with identity link function) linear

mixed-effects model, and p-values were computed with Satterthwaite's method.

Experiment1: RT ~ accuracy * congruency * effector * effector_order + (1 + accuracy + effector |

subject)

Experiment 2: RT ~ accuracy * congruency * effector * effector_order + (1 + accuracy + effector

+ effector_order | subject)

Experiment 3: RT ~ accuracy * congruency + (1 + accuracy_gabor + congruency | subject).

Detailed results are gathered in Table S2 in the Supplementary Material. No effect of

spatial congruency on RT was observed in Experiment 1 (p = 0.17). This suggests that the

presentation of the noise patch (see Method section) 800 ms before the Gabor allowed

participants to more easily control primed responses without any impact on response times.

However, RTs were faster for congruent (Experiment 2: M = 695 ms, se = 20 ms;

Experiment 3: M = 673 ms, se = 23 ms) than incongruent trials (Experiment 2: M = 713

ms, se = 17 ms; Experiment 3: M = 694 ms, se = 23 ms) in Experiment 2 (β = -11.5, p <

0.001) and Experiment 3 (β = -13.7, p = 0.008). This suggests that our action priming

successfully facilitated (impeded) the selection of spatially congruent (incongruent)

perceptual responses. In Experiment 1, we found an interaction between effector and

effector order (β = -31.38, p < 0.001; see SI, Figure S4 and post-hoc tests), indicating that

participants were overall faster in the second block. Notably, if they started with the same

effector block, their reaction times were faster in the different effector block (i.e., the

second block), and those who started with the different effector block showed faster

responses in the same effector block (i.e., the second block for those participants; see SI,

Figure S4 & Post-hoc tests).

Confidence

We analyzed confidence using a cumulative link mixed-effects model, with a probit link

function. Statistical significance was assessed with the Wald test. Formally, we estimated

the following models:

Experiment 1: confidence ~ accuracy * congruency * effector * effector_order + RT + (1 + accuracy

+ congruency + effector + effector_order + RT | subject)

Experiment 2: confidence ~ accuracy * congruency * effector * effector_order + RT + (1 + accuracy

+ congruency * effector + effector_order + RT | subject)

Experiment 3: confidence ~ accuracy * congruency + RT + (1 + accuracy + congruency + RT |

subject).

Detailed results are gathered in Table 1 below. As expected, confidence was greater for

correct trials than for errors in all three experiments (Experiment 1: odds-ratio = 1.9, p <

0.001; Experiment 2: odds-ratio = 1.4, p < 0.001; Experiment 3: odds-ratio = 1.6, p <

0.001). We also found in Experiments 2 and 3 the classical relation between RT and

confidence, with faster responses leading to higher confidence levels (Experiment 2: odds-

ratio = 0.07, p < 0.001; Experiment 3: odds-ratio = 0.14, p < 0.001). This relation was

however not observed in Experiment 1 (p = 0.17). This might be due to a floor effect,

response time being much faster in Experiment 1 than in the two others (see Figure 2).

There was no main effect of effector compatibility on confidence (Experiment 1: p = 0.78;

Experiment 2: p = 0.67), and no significant interaction between spatial congruency and

effector compatibility (Experiment 1: p = 0.22; Experiment 2: p = 0.26, see Table 1). We

also observed some interactions involving effector compatibility, though these were not

consistent across experiments. These interactions are discussed and reported in full in the

Supplementary Material. Overall, they suggest that switching from upper to lower limb

responses in the speeded RT task of Experiment 1 introduced an attentional cost affecting

both confidence and accuracy (i.e., perceptual and metacognitive accuracy decreased in the

different effector block). Additionally, effector compatibility interacted with block order.

Accordingly, the difficulty of switching from upper to lower limbs varied depending on

whether participants began the task with hand or foot responses. These interactions did not

seem to be consistent as they were not replicated across experiments.

Figure 3 The graphs in A show the across-participant average confidence and individual confidence

values as a function of spatial congruency for each experiment. The graph in B shows the across-

participant average confidence as a function of effector compatibility for Experiments 1 and 2.

Error bars represent confidence intervals indicating the dispersion of individual averages, corrected

using the Cousineau & Morey method (Morey, 2008). Note that confidence averages are presented
only for illustrative purposes. Statistics were not performed on averages, since confidence were
treated as an ordinal variable.

Importantly, priming the side of an action consistently modulated confidence across

the three experiments, over and above the specific effector primed, accuracy and reaction

times. Indeed, we observed a main effect of spatial congruency, with higher confidence in

incongruent than congruent trials in the three experiments (Experiment 1: odds-ratio =

0.94, p = 0.004; Experiment 2: odds-ratio = 0.94, p = 0.017; Experiment 3: odds-ratio =

0.90, p < 0.001). This effect was slightly larger for errors than for correct trials in

Experiment 1 (odds-ratio = 1.04, p = 0.028). However, the interaction between congruency

and accuracy was not significant in Experiment 2 (p = 0.94), nor in Experiment 3 (p =

0.18). Only in Experiment 2 we observed a three-way interaction involving accuracy,

spatial congruency and effector order (odds-ratio = 0.98, p = 0.024, see SI Figure S3).

However, post-hoc tests revealed no significant contrasts (see SI, post-hoc tests), leaving

some statistical ambiguity regarding the interpretation of this interaction.

Hence, the consistent finding across experiments is the modulation of spatial

congruency on confidence, with higher confidence for spatially incongruent actions

compared to spatially congruent ones. Importantly, this effect was observed even after

controlling for accuracy and response times in the models, and even when perceptual

accuracy did not significantly differ between congruent and incongruent trials (see

Experiments 1 and 2). The direction of the congruency effect on confidence supports the

action monitoring hypothesis rather than the action fluency hypothesis.

Table 1. Hierarchical regression tables predicting confidence from accuracy, congruency, and reaction time for Experiment 3 (right), plus effector and effector order for Experiment 1 (left), 2 (middle), and 3 (right). Coefficients were coded as follows: Accuracy: error = -1, correct = 1; Congruency: incongruent = -1, congruent = 1; Effector compatibility: different effector (feet in Experiment 1, middle fingers in Experiment 2) = -1, same effector (index fingers) = 1; Bloc order: same effector first = -1, different effector first = 1. Response time is mean-centered. The models tested were the following:

Experiment 1: confidence ~ accuracy * congruency * effector * effector_order + RT + (1 + accuracy + congruency + effector + effector_order + RT | subject)

Experiment 2: confidence ~ accuracy * congruency * effector * effector_order + RT + (1 + accuracy + congruency * effector + effector_order + RT | subject)

Experiment 3: confidence ~ accuracy * congruency + RT + (1 + accuracy + congruency + RT | subject)

	Confidence (Exp. 1)			Confidence (Exp. 2)			Confidence (Exp. 3)		
Predictors	Odds Ratios	CI	p	Odds Ratios	CI	p	Odds Ratios	CI	p
1|2	0.66	0.50 – 0.86	**0.002**	0.43	0.31 – 0.60	**<0.001**	0.85	0.74 – 0.99	**0.037**
2|3	1.66	1.27 – 2.17	**<0.001**	1.23	0.89 – 1.70	0.209	2.36	2.03 – 2.74	**<0.001**
3|4	4.49	3.43 – 5.88	**<0.001**	3.12	2.25 – 4.32	**<0.001**	6.05	5.17 – 7.07	**<0.001**
accuracy gabor [1]	1.90	1.64 – 2.20	**<0.001**	1.42	1.32 – 1.52	**<0.001**	1.58	1.43 – 1.75	**<0.001**
congruency [1]	0.94	0.90 – 0.98	**0.004**	0.94	0.89 – 0.99	**0.017**	0.90	0.86 – 0.94	**<0.001**
effector [1]	1.01	0.94 – 1.09	0.781	0.98	0.88 – 1.09	0.668			
effector order [1]	0.78	0.61 – 0.99	**0.043**	0.74	0.54 – 1.03	0.074			
rt gabor centered	0.59	0.28 – 1.26	0.175	0.07	0.04 – 0.12	**<0.001**	0.14	0.09 – 0.22	**<0.001**
accuracy gabor [1] * congruency [1]	1.04	1.00 – 1.09	**0.028**	1.00	0.96 – 1.04	0.937	1.03	0.99 – 1.07	0.185
accuracy gabor [1] * effector [1]	1.08	1.04 – 1.13	**<0.001**	1.01	0.97 – 1.06	0.479			
congruency [1] * effector [1]	0.98	0.94 – 1.01	0.221	0.98	0.94 – 1.02	0.258			
accuracy gabor [1] * effector order [1]	0.94	0.81 – 1.09	0.396	1.03	0.96 – 1.10	0.444			
congruency [1] * effector order [1]	1.03	0.99 – 1.07	0.169	1.04	0.99 – 1.10	0.107			
effector [1] * effector order [1]	0.88	0.82 – 0.96	**0.003**	0.92	0.83 – 1.02	0.110			
(accuracy gabor [1] * congruency [1]) * effector [1]	1.02	0.98 – 1.06	0.342	1.03	0.99 – 1.07	0.127			
(accuracy gabor [1] * congruency [1]) * effector order [1]	1.00	0.96 – 1.04	0.875	0.96	0.92 – 0.99	**0.024**			
(accuracy gabor [1] * effector [1]) * effector order [1]	1.08	1.04 – 1.12	**<0.001**	1.00	0.96 – 1.04	0.926			
(congruency [1] * effector [1]) * effector order [1]	1.00	0.96 – 1.04	0.924	1.03	0.98 – 1.07	0.252			
(accuracy gabor [1] * congruency [1] * effector [1]) * effector order [1]	1.01	0.97 – 1.04	0.792	0.99	0.95 – 1.03	0.460			

ERP Results (Experiment 3)

Post-stimulus activity. To investigate the way action priming may have influenced confidence judgment we analyzed whether congruency modulated EEG markers of cognitive control that are typically observed after stimulus presentation, such as the P2/N2

complex (and theta oscillations, see SI). We reasoned that the action prepared in response

to the visual cue could have conflicted with the action required to report the orientation of

the Gabor (i.e., incongruent trials). Specifically, preparing a congruent action in advance

may have facilitated response selection, while preparing an incongruent action may have

led the participant to control and inhibit unwanted choices as soon as the Gabor was

presented. This conflict would be observed rather early after the onset of the Gabor.

A cluster-based permutation test performed on a time window going from 0 ms to

350 ms post-stimulus onset (i.e. we selected a time window in which no perceptual

response could have already occurred), identified a significant temporal cluster at FCz

($t(23) = -2.6$, $p < 0.05$, Figure 4B) going from 190 to 240 ms (similar results for Fz and Cz,

see SI *Figure S7 & S8*), i.e., in a time period where the P2 component is typically observed

(Ghin et al., 2022). The amplitude of the P2 was higher for incongruent compared to

congruent trials.

Since there was a small effect of congruency on accuracy in Experiment 3, we

performed the same tests on correct trials only, which yielded to similar results (see SI,

Figure S7). We observed no modulation of congruency on the N2. Similarly, post-stimulus

theta oscillations did not vary between spatially congruent and spatially incongruent trials

(see SI for a discussion of these null results).

Post-response activity. Further analyses investigated the influence of spatial congruency

on post-response signals. Firstly, we conducted a cluster-based permutation test comparing

congruent and incongruent EEG epochs going from 0 to 500 ms post-response at Pz (other

electrodes and group of electrodes were used for these analyses, all leading to similar

results, see SI). This analysis revealed two significant temporal clusters (Figure 2A), one

spanning from 70 ms to 240 ms, and another one from 260 ms to 320 ms. Subsequently,

Figure 4. (**A, Left**) Grand average post-response ERPs at Pz, time-locked to the onset of the

perceptual response (vertical dotted line), for congruent (orange lines) and incongruent (blue lines)

trials, with low confidence (levels 1 and 2) shown in light colors and high confidence (levels 3 and

4) in dark colors. (**B, Left**) The graph depicts stimulus-locked ERP at FCz for the congruent (in

orange) and incongruent (in blue) trials. Red horizontal lines in both graphs, represent significant

time windows obtained with a cluster-based permutation test comparing congruent and incongruent

trials. The topographies presented on the right panel depict the voltage difference between

congruent and incongruent trials averaged across the time window of the temporal cluster observed

in the response-locked (**A, Middle**) and the stimulus-locked (**B, Middle**) ERPs. Note that for the

response-locked ERPs, the second cluster is depicted. The bar plots in the right panel depict the

across-participant average amplitude of post-response activity for the two temporal clusters (**A,**

Right) and post-stimulus activity at P2 (**B, Right**). Vertical bars represent the standard error of the

mean.

we categorized trials into low confidence (levels 1 and 2) and high confidence (levels 3 and 4) trials. To investigate interactions between confidence and congruency, we conducted a 2 (congruency: congruent vs. incongruent) by 2 (confidence: high vs. low) repeated-measures ANOVA on the average amplitude of each temporal cluster. The analyses of the early cluster (i.e., from 70 to 240 ms) showed a main effect of congruency ($F(23) = 10.66$, $p = 0.003$, Figure 2A), with a more negative activity for incongruent compared to congruent trials. Neither the main effect of confidence nor the interaction was significant ($F(23) = 0.03$, $p = 0.86$ and $F(23) = 0.25$, $p = 0.62$, respectively). In the second temporal cluster (i.e., from 260 to 320 ms) we observed a main effect of both congruency ($F(23) = 8.71$, $p = 0.007$) and confidence ($F(23) = 5.05$, $p = 0.03$, Figure 2A). The EEG activity was more negative for incongruent compared to congruent, and for high compared to low confidence trials. No significant congruency and confidence interaction were observed ($F(23) = 0.37$, $p = 0.55$). Since there was a small behavioral effect of congruency on accuracy, we performed the same EEG analyses on correct trials only and we observed similar results (see SI, Figure S7).

LRP results. The jackknife-based analysis revealed that the congruent LRP started slightly earlier (interpolated estimate = -184 ms) compared to the incongruent LRP (interpolated estimate = -157 ms), showing a difference in onset latency of 27 ms (SD = 7.2, $t(23) = 3.77$, $p < 0.01$, Figure 5). These findings, along with the observed behavioral effect of congruency on response times (cf. Experiments 2 and 3), suggest that the motor priming task successfully induced action preparation prior to Gabor presentation.

Response-locked LRP

Figure 5. Response-locked Lateralized Readiness Potential for the congruent (red line) and incongruent (blue line) trials prior to the onset of the perceptual response (vertical dotted black line). The colored dotted lines represent the LRP onset latency at 1/3 threshold of the peak amplitude for congruent (blue) and incongruent (red) trials extracted from a Jackknife-based procedure showing a significant difference of 27 ms in LRP onset latencies.

Discussion

Motor processes engaged during perceptual decisions have been found to affect retrospective confidence judgments. In three pre-registered experiments, we tested two hypotheses regarding how motor preparation impacts perceptual confidence. The "fluency hypothesis" suggests that the ease and smoothness of selecting and executing a motor

response may serve as a cue for confidence judgments (Fleming et al., 2015). In contrast,

the "monitoring hypothesis" proposes that confidence is influenced by action monitoring

processes; it is not the fluency of the action, but rather the degree of control and the level

of monitoring required for implementing a perceptual decision that affect confidence (cf.

Sanchez et al., 2024). To test these hypotheses, participants reported the orientation of a

Gabor stimulus (vertical or horizontal) and judged their confidence in their decision. Using

a novel motor priming paradigm, participants were prompted to prepare an action in

advance, which could be either compatible or incompatible with the action ultimately

chosen to report the Gabor orientation.

In the three experiments, spatial congruency consistently influenced confidence

judgments. Specifically, trial-by-trial analyses with mixed-effect models controlling for

accuracy and response time revealed that participants were more confident in their decision

when their response was spatially incongruent with the primed action (see Figure 3 and

Table 1). This effect was general and independent of whether we primed the same (index

fingers) or a different effector (feet in Experiment 1, and middle fingers in Experiment 2).

Accordingly, these results suggest that early action preparation, involving spatial

representation but not effector-specific representation, significantly influenced perceptual

confidence. The fact that spatial representation of motor planning impacted response time

and confidence can be explained by the partial overlap of spatial and semantic codes across

different effectors and different actions (Fournier et al., 2010). Additional evidence that

action preparation was modulated specifically by spatial congruence, rather than by

effector compatibility, comes from the fact that responses were faster in spatially congruent

compared to spatially incongruent trials (cf. Experiments 2 and 3). Furthermore, we

observed an earlier beginning of the Lateralized Readiness Potentials (LRP) in spatially

congruent compared to spatially incongruent responses. LRP activity has been typically

attributed to motor preparation activity (Schmitz et al., 2019). Accordingly, these findings

may suggest that the faster response times for congruent responses result from earlier motor

preparation, whereas incongruent responses may involve delayed motor preparation (cf.

Mordkoff & Gianaros, 2000). Overall, the findings on response times and LRP, confirms

that our priming paradigm successfully influenced the selection/preparation of perceptual

response, particularly for spatially congruent actions.

At first glance, the confidence results might seem compatible with many theories

linking confidence to the accumulation of evidence for a decision (see Desender et al.,

2021, for a recent investigation of confidence within evidence accumulation models). The

fact that participants were more accurate (in Experiment 3) and slower (in Experiments 2

and 3) for incongruent trials might suggest that they were more cautious in these trials (cf.

Vickers & Packer, 1982), and that they were accumulating more evidence before making

their decision, which would result in a higher confidence. However, this explanation is

unlikely for two main reasons. First, longer response times for incongruent trials are more

likely due to inhibitory processes at the motor planning level (as revealed by the LRP

observed in these trials in Experiment 3) than to prolonged evidence accumulation. Second,

and more importantly, longer evidence accumulation would predict an increase of

confidence for correct responses but a decrease in confidence for errors. Instead, we

observed a boost in confidence for both correct and incorrect incongruent trials, with this

effect being stronger for errors than for correct trials in Experiment 1.

The observed confidence boost in incongruent trials provides stronger support for the

monitoring hypothesis rather than the fluency hypothesis. In fact, the fluency hypothesis

would have predicted the opposite result: that actions whose selection is facilitated (i.e.,

congruent actions) lead to higher confidence judgments than incongruent actions (cf.

Fleming et al., 2015). Our findings align with our previous study showing that, even when

controlling for response time, perceptual responses preceded by partial motor activations

(either ipsilateral or contralateral) consistently yielded higher confidence (Gajdos et al.,

2019). It has been shown that partial activations are followed by an Error-Related

Negativity (Meckler et al., 2017) originating in the Supplementary Motor Area (Bonini et

al., 2014), a structure involved in action monitoring (Coull et al., 2016). Hence, these

results suggested that partial motor activations may reflect premature commitments to a

response that are effectively controlled and inhibited, thereby resulting in a greater sense

of control over one's responses and thus boosting confidence.

Similarly, in our study, perceptual responses that were incongruent with planned

actions may have been interpreted by the system as instances in which an unwanted motor

plan was successfully controlled, thereby leading to an increased sense of control and

confidence. In other words, the ability to successfully resolve response competition by

engaging monitoring mechanisms (Desender et al., 2014; Questienne et al., 2018) informs

confidence estimations.

Stimulus-locked ERPs observed in Experiment 3 provide further support for this

explanation. Our analyses revealed higher amplitude of the P2 component in incongruent

trials compared to congruent trials. The frontal P2 component is thought to reflect early

attentional resource allocation (Xie et al., 2020) associated with cognitive control (Ghin et

al., 2022), sensitive to early updating processes during task-switching (Capizzi et al.,

2015), and related to inhibitory processing, with a higher P2 for successful inhibited actions

(Senderecka et al., 2012). We argue that preparing incongruent actions resulted in a greater

involvement of early attentional resources necessary for response control, thereby

impacting confidence. Interestingly, a recent study showed that TMS pulses to the

dorsolateral prefrontal cortex (dlPFC) at 200 ms after stimulus onset influence confidence,

suggesting confidence computations begin before the perceptual decision is fully formed,

challenging post-response models where confidence is computed only after a response

(Xue et al., 2023). Our study corroborates this notion by suggesting that post-stimulus

processes as early as 200 ms correlate with retrospective confidence.

Another interesting aspect of the modulation of spatial congruency on confidence is

that this effect was also observed in Experiment 1, despite the motor priming task not

directly influencing motor execution—i.e., no effect of spatial congruency on response

times was observed. The absence of a congruency effect on response times can be

explained by the fact that, in the perceptual task of Experiment 1, a noise patch was

presented 800 ms before the Gabor stimulus, signaling to participants that the Gabor

presentation was imminent. As a result, even though the speeded RT task may have initially

prepared the system for a specific action, the 800 ms delay likely eliminated the impact of

motor priming on motor execution. This is consistent with the well-established finding that

priming effects decay over time (Van den Bussche et al., 2009). However, despite the lack

of an observable effect on motor execution, the conflicting motor representation induced

by the speeded RT task still modulated retrospective confidence. This finding suggests that

the motor signal used by the system to modulate confidence operates relatively early in the

motor processing hierarchy—specifically, at the level of early motor planning rather than

late motor preparation or motor execution. Interestingly, Fleming et al. (2015) also found

that early motor processes play a crucial role in confidence modulation, rather than late

motor processes or motor execution itself.

Although the current study supports the monitoring hypothesis, the specific feature

that triggers monitoring processes has yet to be identified. Two distinct, but not mutually

exclusive, factors may be at play. One possibility is that the effect stems from conflicting

motor plans. Alternatively, the conflict driving monitoring and changes in confidence may

arise between a motor plan and the perceptual decision the participant intends to commit

to. Further research is needed to address this issue. While the exact feature triggering

monitoring and affecting confidence is still unclear, our results more generally align with

several studies demonstrating that cues and heuristics, such as stimulus visibility cue

(Shekhar and Rahnev, 2024), response time (Kiani et al., 2014) or interoceptive states

(Allen et al, 2016) do contribute to people's sense of confidence beyond the strength of

accumulated evidence.

Our findings are in apparent contrast with recent research supporting the fluency

hypothesis. Fleming et al. (2015) showed that TMS stimulation over premotor areas linked

to the chosen perceptual response enhances confidence, whereas incongruent stimulation

decreases confidence. A critical difference between Fleming et al.'s study and ours may lie

in the presence of response conflict. In their study, stimulation of the premotor area may

have facilitated response planning without inducing implicit or explicit conflict with the

action selected to report the stimulus. Participants may not have even perceived any

difference between priming a congruent or incongruent action. In contrast, our study

involved tasks with a certain degree of action conflict, thereby more strongly engaging

monitoring and action control processes.

It could be argued that both Fleming et al.'s study and ours rely, after all, on similar

processes, such as individuals' control over their actions. Research on action fluency has

shown that subliminal motor priming reduces participants' sense of control, when the prime

is incongruent with the performed action, but it increases it when the prime is congruent

(Sidarus, Chambon, & Haggard, 2013; Chambon & Haggard, 2012; Wenke, Fleming, &

Haggard, 2010). A similar interpretation may apply to Fleming et al.'s (2015) study, where

participants may have experienced more control over fluent (congruent) responses

compared to incongruent TMS stimulations. Interestingly, recent findings suggest that

tasks involving response conflict, and thus requiring more cognitive control, increase

individuals' sense of control over their actions (Van den Bussche et al., 2020). Therefore,

one might argue that in our experiment, a greater sense of control occurred during

incongruent actions, as participants successfully resolved a conflict by controlling and

inhibiting unwanted responses. In essence, the increase in perceptual confidence observed

both in our study and in Fleming et al.'s study may rely on an increased sense of control

over one's actions. However, these experiences of control are triggered by distinct signals.

Further studies should explore the relationship between sense of control and perceptual

confidence.

Converging results regarding the influence of spatial congruency on confidence were

also observed when examining post-response brain activity. Specifically, spatial

congruency appeared to modulate post-decisional markers of response evaluation in two

separate time windows. The first time-window spanned from 70 ms to 240 ms after the

perceptual response. Activity within this period typically reflects early response

monitoring processes, such as those underlying the Error-Related Negativity (ERN;

Scheffers & Coles, 2000), a waveform observed following the detection of an incorrect

choice. In the present study, this first post-response temporal window was solely modulated

by congruency, and not by confidence. We argue that activity within this period reflects an

early retrospective evaluation of the motor response, signaling the resolution of a motor

conflict and the successful control of an unwanted motor plan. It is important to note that

we could not identify a typical ERN component in our study (see SI). The ERN is

classically observed in tasks (e.g., Simon task, Flankers task, etc.) where errors are

explicitly detected by the participants. However, instances where participants realized they

made a mistake were very rare in our experiment (see Methods). Therefore, we cannot and

do not attempt to attribute our effect specifically to the ERN.

The second time window spanned from 260 ms to 320 ms after the response, where

the Error Positivity (Pe; Pereira et al., 2020) is typically observed. The Pe is an event-

related component characterized by a positive amplitude following incorrect choices. It is

regarded as a reliable indicator of an individual's ability to monitor errors during decision-

making tasks (Davies et al., 2001). Recent studies have identified a family of ERP waves

within the time window traditionally associated with the Pe. These waves not only reflect

error monitoring but also appear to encode different levels of confidence (Boldt & Yeung,

2015). Specifically, within the time window where the classical Pe is observed, there are

graded changes in voltage amplitude, corresponding to varying levels of confidence.

Higher confidence levels are associated with more negative amplitudes, whereas lower

levels of confidence and the certainty of error tend to manifest as more positive values.

In line with these observations, the second post-response temporal cluster we observed

covaried not only with spatial congruency but also with confidence, consistent with

previous results on the Pe (Boldt & Yeung, 2015; see SI). Specifically, high-confidence

judgments were associated with more negative amplitudes compared to low-confidence

trials. Hence, we argue that this second temporal cluster reflects a metacognitive evaluation

of response accuracy. We would like to highlight that as for the ERN, we did not identify

a typical Pe, since trials in which participants acknowledged they made an error were very

rare.

Surprisingly, preparing the same or a different effector than the one used to report the

Gabor did not seem to impact response times and confidence. We believe that the absence

of this effect can be explained by the experimental design. For example, in the first

experiment, we observed a clear cost when preparing foot actions and reporting the Gabor

with the hands. Notably, accuracy decreased in this condition compared to when both the

speeded RT task and the perceptual task required hand responses. If incongruent motor

plans tend to increase confidence (because they require more monitoring), while incorrect

responses reduce confidence, then these two effects may have cancelled each other out.

Actually, in the different-effector block of the first experiment, not only did perceptual

accuracy decrease, but metacognitive accuracy also declined. This suggests that the cost of

switching from foot to hand responses impaired both perceptual and metacognitive

evaluations, as if participants were distracted by the transition from lower to upper limb

responses. In the second experiment, we aimed to eliminate the cost associated with feet

responses by replacing these effectors with the middle fingers. Similarly, we observed no

modulation of effector-specific representations on confidence or reaction times, but we

observed a clear effect of spatial congruency on these two variables. This may be due to

the stronger overlapping of motor codes for actions performed with the same hand than for

actions performed with two different hands. In other words, for example, switching from

left to right hand responses would likely generate more conflict than switching from the

index to the middle finger of the same hand. Further studies may address these difficulties

by employing, for example, hand actions and eye movements.

In summary, our results indicate that motor processes contribute, beyond perceptual

evidence, in shaping retrospective perceptual confidence. Action planning modulate

confidence through the involvement of monitoring systems (cf. also Yeung & Summerfield

2012). According to this framework, the system continuously monitors sensory and action

information, with confidence estimates influenced by how effectively the system inhibits

and controls alternative and unwanted responses (Gajdos et al., 2019; Anzulewicz et al.,

2019). Thus, action processes are not merely ancillary processes for decision

implementation but actively contribute to the formation of high-level cognitive constructs

such as retrospective confidence.

Constraints on Generality

The participants in this study were predominantly neurotypical young adults, a

demographic often associated with greater cognitive flexibility and relatively high levels

of digital literacy. Given the nature of recruitment methods, which were likely to attract

educated individuals, these findings may not fully generalize to populations with different

neurological profiles, age ranges, or educational backgrounds. Consequently, caution is

warranted in extending these results to older adults, neurodivergent individuals, or those

from less digitally engaged or lower educational backgrounds.

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1100