

Research Paper

Context-dependent Modulation of Visual Recognition
by Conscious and Unconscious PrimingSamaneh Navab Kashani¹ , Reza Khosrowabadi^{1*} , Mohammad-Reza A. Dehaqani^{2,3*}

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ABSTRACT

Introduction: Despite extensive study, how conscious and unconscious priming influence visual perception remains only partially understood. In this work, we examine their distinct effects across multiple experimental conditions within a binocular rivalry paradigm to provide a more comprehensive perspective on identity and category recognition.**Methods:** Participants were presented with word or image primes, followed by a name-picture verification task in which faces or animal bodies served as targets. Although conscious priming has been shown to facilitate identity recognition while interfering with category perception, the precise distinction between conscious and unconscious states, as well as the mechanisms underlying unconscious priming, requires further detailed analysis. Left-hemispheric processing was one of the influential factors in distinguishing conscious from unconscious priming effects.**Results:** Interestingly, we observed a negative correlation between conscious and unconscious perception during identity recognition, highlighting the condition-dependent modulation of visual perception in the priming paradigm.**Conclusion:** From the regression analyses, awareness showed the strongest predictor of priming magnitude, with recognition level playing a secondary role. More broadly, our findings demonstrate that visual perception circuits are modulated condition-dependently, underscoring how awareness and trial context jointly shape recognition processes.

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Highlights

- Conscious priming facilitated identity but impaired category recognition.
- Unconscious priming lacked identity–category dissociation.
- Identity priming showed negative conscious–unconscious correlation.
- Right visual field effects distinguished awareness states.
- Awareness was the strongest predictor of priming magnitude.

Plain Language Summary

Our brains are constantly influenced by things we have just seen—even when we are not aware of seeing them. This phenomenon is called priming. In this study, we explored how visible (conscious) and invisible (unconscious) hints affect the way people recognize images. Participants viewed brief words or pictures before being asked to identify either a specific individual (for example, a particular dog breed or person) or a broader category (such as “dog” or “animal”). Sometimes these hints were clearly visible. Other times they were hidden from awareness using a special visual technique. We found that when people were consciously aware of the hint, it helped them recognize specific identities faster—but surprisingly, it made them slower at recognizing broader categories. In contrast, when the hint was invisible, this clear pattern disappeared. Even more interesting, for identity recognition, the effects of visible and invisible hints often moved in opposite directions. What helped in the conscious condition could hinder in the unconscious one. We also discovered that awareness itself was the most important factor shaping these effects—more important than whether the hint was a word or picture, or whether it appeared on the left or right side of the screen. Why does this matter? These findings show that conscious and unconscious influences on perception are not simply weaker or stronger versions of the same process—they work differently. Understanding how awareness shapes recognition may help us better understand attention, decision-making, advertising effects, and even clinical conditions where perception or awareness is altered.

Introduction

Priming is a central mechanism in visual perception, shaping how prior exposure to stimuli influences subsequent processing at both behavioral and neural levels (Balconi, 2006; Dehaene et al., 1998; Jiang et al., 2007; Marsolek, 1999; Stein et al., 2020).

While extensive work has characterized general priming effects, how conscious and unconscious priming distinctly modulate identity versus category recognition is not well understood (Amihai et al., 2011; Chien et al., 2023; Moradi et al., 2005a). Most prior research has treated these processes in isolation, often ignoring how awareness interacts with different prime modalities, such as words versus images, or with target types, including faces and animal bodies.

Recognition at the identity level requires precise differentiation of individual exemplars, whereas category recognition relies on broader, abstract representations of

object classes (Johnson & Mervis, 1997; Rosch et al., 1976). Neuroimaging and electrophysiological studies indicate that category information is encoded rapidly in the inferior temporal cortex, whereas identity recognition depends more on feedback pathways supporting fine-grained processing (Dehaqani et al., 2016). Despite this distinction, the extent to which conscious and unconscious priming modulate these processes across awareness states remains largely unexplored.

Evidence from unconscious priming demonstrates that masked stimuli, including words and images, can facilitate subsequent perception and evoke category-specific neural activity, such as activation of the fusiform face area for faces (Breitmeyer et al., 2005; Dehaene et al., 2001; Kouider et al., 2009; Weibel et al., 2013). Emotional expressions of unseen faces can trigger amygdala responses, yet successful identity recognition often requires conscious perception (Moradi et al., 2005a; Pessoa et al., 2005). Clinical phenomena, including blindsight and unilateral neglect, further illustrate that

category-level processing can occur without awareness, whereas identity-level recognition is constrained by conscious access (Berti & Rizzolatti, 1992; Trevelyan et al., 2007). Collectively, these findings highlight a critical gap: the differential impact of conscious versus unconscious priming on identity and category recognition remains unresolved.

To address this, we employed a binocular rivalry paradigm adapted from Navab Kashani et al. (2025), which allowed precise manipulation of prime awareness by presenting stimuli to one eye while suppressing perception with a rival stimulus (Navab Kashani et al., 2025). Participants then performed a name-picture verification task using faces or animal bodies, enabling the examination of priming effects across prime and target types. Regression analyses quantified the contributions of awareness, prime type, target type, and hemispheric asymmetries, providing a comprehensive assessment of factors driving recognition performance.

Our results demonstrate that unconscious priming showed the reverse pattern for identity recognition. These findings reveal that conscious and unconscious priming differentially modulate visual recognition through distinct neural pathways. Overall, the present experiment indicates that the impact of individual factors is highly condition-dependent, with variations across experimental contexts shaping the way priming influences perception.

Materials and Methods

Study participants

A total of 22 right-handed adults (14 male; mean age=35.7 years, range=32-43), all healthy and unfamiliar with the task, took part in the experiment. Each reported normal or corrected-to-normal vision and gave written informed consent. The study complied with the Declaration of Helsinki, and ethical approval was obtained from the Shahid Beheshti University Ethics Committee.

Stimuli and apparatus

The experiment was conducted in a dimly lit room, with participants seated 64 cm from a monitor (1920×1080 resolution, 144 Hz refresh rate, 1 ms response). Visual input was delivered through a mirror stereoscope attached to a chinrest, ensuring stable alignment. Stimuli were presented against a uniform gray background.

Stimuli consisted of Persian words and grayscale images representing four classes—women, men, cats, and dogs—evaluated at two recognition levels (identity and category). Each category included a name label, a symbolic category image, and 3 identities represented both verbally and pictorially (Navab Kashani et al., 2025). Continuous flash suppression (CFS) (Tsuchiya et al., 2009) was generated in MATLAB Psychtoolbox (Brainard, 1997) using Mondrian noise patterns (colored squares flashing at 20 Hz).

Face and animal stimuli were collected from online image searches and were luminance-matched using the SHINE Toolbox. Except for primes (4° visual angle), all stimuli subtended 6° to avoid retinotopic overlap (Posner & Cohen, 1980; Posner & Cohen, 1984). A textured black-framed non-prime stimuli-and-white border extending to 6.5° (Figure 1).

Procedure and design

Awareness was manipulated via binocular rivalry combined with CFS, a method offering extended suppression and precise perceptual control compared to standard masking (Axelrod et al., 2015; Tsuchiya & Koch, 2005). In this setup, competing images presented to each eye lead to dominance of one stimulus while the other remains suppressed yet influential (Blake, 2001; Fang & He, 2005). Mondrian noise, flashing at 20 Hz, maintained suppression for up to 50 s, allowing subliminal primes to be presented in one eye. Unlike b-CFS paradigms (Jiang et al., 2007), this design assessed the impact of subliminal primes on subsequent tasks rather than suppression break. Primes were shown foveally but lateralized to one visual field to assess hemispheric asymmetries.

Experimental design

To test identity and category recognition, we employed a name-picture verification paradigm (Collin & McMullen, 2005). Participants judged whether a word label (e.g. “Bulldog” for identity or “Dog” for category) matched a subsequent picture, with match/mismatch trials balanced. Identity trials required subordinate-level recognition (e.g. “Bulldog” followed by Bulldog or Doberman), while category trials required basic-level recognition (e.g. “dog” followed by dog or cat). Mismatch conditions used within-category distractors for identity and cross-category distractors for category, avoiding overlap at the superordinate level. Primes were either images or words, with relevant primes consistent with the target and irrelevant primes drawn from unrelated categories or higher levels.

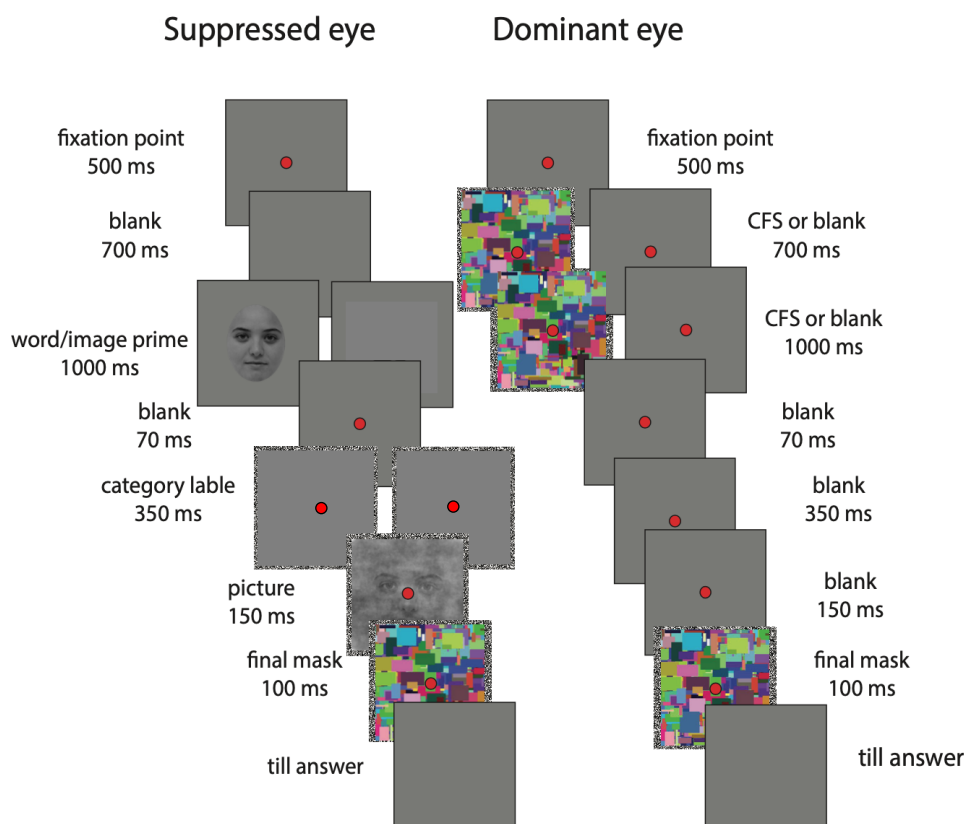


Figure 1. Experimental paradigm

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Note: Illustration of the binocular rivalry task with CFS. In this setup, one frame was delivered to the suppressed eye (left/right visual field) and the other to the dominant eye (left or right visual field), enabling comparison between conscious (no CFS) and unconscious (CFS) conditions. Primes—either words or images—were presented at the identity or category level, followed by a name–picture verification task.

Participants first underwent training without stereoscopes or primes, learning to identify categories through paper-based practice. Feedback was provided via fixation color (green=correct, red=incorrect), and only those reaching >95% accuracy continued.

The main task, conducted without feedback, required right-hand keypresses (right=match, left=mismatch) while fixating centrally. The prime and target presentations were lateralized to the left/right visual fields, respectively. In unconscious conditions, primes were suppressed by CFS noise in the contralateral field. The CFS paradigm randomly suppressed primes across eyes, effectively balancing eye dominance across conditions. Each trial began with 500 ms fixation, followed by 1700 ms of either CFS (unconscious) or blank (conscious). Primes appeared gradually for 1000 ms at low contrast (0.15–0.4%). After a 70-ms blank, a word label (350 ms) was followed by a picture (150 ms), which was immediately masked with bilateral Mondrian noise (100 ms).

Conscious primes were visible for 1000 ms without CFS (Navab Kashani et al., 2025).

In this experimental framework, 7 factors were manipulated in a factorial design, with each trial yielding a reaction time (RT) and an accuracy measure. The factors included priming (relevant vs irrelevant), awareness (conscious vs unconscious), recognition level (identity vs category), prime type (image vs word), target type (face vs animal), visual field laterality (left vs right), and matchness (match vs mismatch) that were presented in intermix trials. The target set comprised 6 exemplars—three male and three female faces, three cats, and three dogs—resulting in a total of 768 trials per participant.

Statistical analysis

Trials with incorrect responses or RTs outside 50–6000 ms were excluded (<5%). Mean RTs per participant were computed, focusing on RT because it is sensitive to priming. Planned comparisons were analyzed using

two-tailed Wilcoxon signed-rank tests, and correlations between variables were examined with the Pearson correlation coefficient (r).

Prime index (PI)

The PI was defined as the difference in mean RT between trials with a relevant prime and trials with an irrelevant prime, computed separately for identity and category tasks. If relevant-primed RT=400 ms and irrelevant=450 ms, then PI=−50 ms

Priming effect difference (PED)

To examine differential priming effects, we analyzed 16 conditions combining prime properties: laterality (left/right visual field), target type (face/animal), prime type (word/image), and recognition level (identity/category). For each condition (e.g. left-face-word-identity), we calculated eight PIs representing all combinations of these factors within that condition (e.g. left: F-I-ID, F-W-ID, A-I-ID, A-W-ID, F-I-Ca, F-W-Ca, A-I-Ca, A-W-Ca). To compare 2 conditions, we computed the Pearson correlation between the sets of eight PIs for each condition. The PED was defined as 1 minus this correlation coefficient, providing a robust measure of dissimilarity in priming effects across conditions. If the correlation coefficient between animal and face conditions=−0.2, then PED=1−(−0.2)=1.2.

Support vector regression (SVR) for predicting the prime index

To quantify the contribution of different experimental factors to RT, we employed a SVR model with a radial basis function (RBF) kernel. SVR is a robust method for handling nonlinear relationships and is particularly effective in high-dimensional spaces (Smola & Schölkopf, 2004). The response variable was the prime index, defined as the difference in RTs between the prime and the irrelevant prime conditions. The predictor variables included 5 experimental factors: laterality (left/right visual field), prime type (image/word), target type (face/animal), recognition level (identity/category), and awareness (conscious/unconscious).

Before training the SVR model, all predictor variables were standardized using z-score normalization to ensure comparability and prevent dominance by variables with larger scales:

$$X' = \frac{X - \mu_X}{\sigma_X}$$

, where X' is the standardized feature, μ_X is the mean, and σ_X is the feature's standard deviation.

The SVR model was trained using epsilon-insensitive loss, which minimizes errors within a predefined margin ϵ and optimizes the following objective function (Vapnik, 1998):

$$\min_{w,b} \frac{1}{2} \|w\|^2 + C \sum_{i=1}^n \max(0, |y_i - f(X_i)| - \epsilon)$$

, where w is the weight vector, b is the bias term, C is the regularization parameter, and $f(X)$ represents the predicted response. The model was implemented using MATLAB's fitsvm function with an RBF kernel.

Sensitivity analysis for feature contribution

To assess the contribution of each factor, we conducted sensitivity analysis by systematically varying individual predictors while holding others constant and measuring the resulting changes in model predictions. Specifically, for each predictor X_i , we generated a range of values within the observed data limits and computed the corresponding predictions:

$$\hat{Y}(X_i) = f(X_1, \dots, X_i, \dots, X_n)$$

The sensitivity score for each factor was calculated as the standard deviation of the predicted values, reflecting the extent to which changes in that predictor influenced the response variable:

$$S_i = \text{std}(\hat{Y}(X_i))$$

To facilitate comparison, sensitivity scores were normalized:

$$S'_i = \frac{S_i}{\sum_{j=1}^n S_j}$$

Higher values indicate a greater influence of the predictor on the PI. Sensitivity analysis provides an interpretable measure of feature importance without requiring explicit model assumptions (Saltelli et al., 2008).

SVR is a machine learning method that extends support vector machines (SVMs) to regression tasks by finding a function that best predicts a continuous response variable while maintaining a tolerance margin ϵ around the true values (Smola & Schölkopf, 2004). Using a kernel function, SVR can model complex, nonlinear relation-

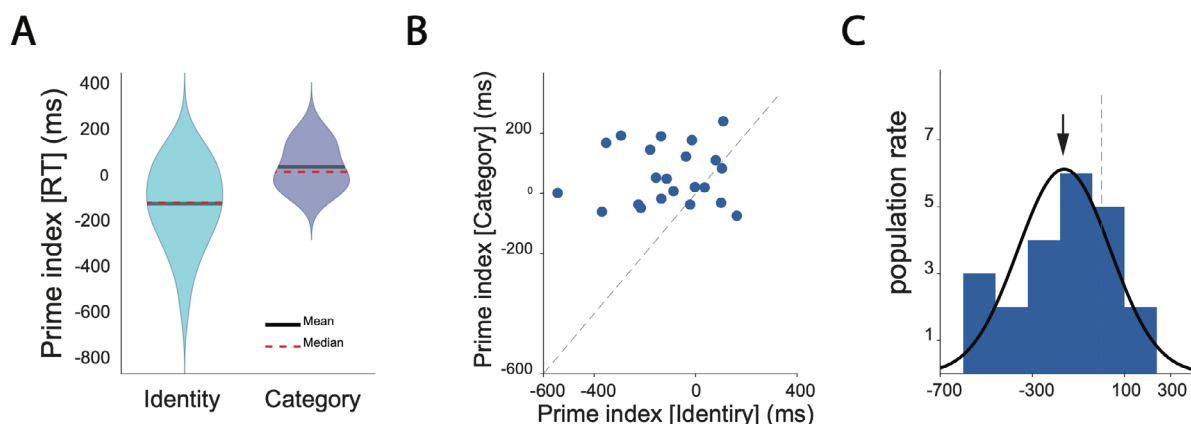


Figure 2. Conscious priming exerts distinct effects on identity and category recognition

A) The PI, defined as the mean RT difference between relevant and irrelevant prime trials, diverged by recognition level under conscious priming; B, C) The scatterplot and histogram illustrate inter-individual variability, with the majority of participants (18/22) exhibiting faster identity RTs and slower category RTs under conscious conditions.

ships between predictors and the outcome, making it particularly useful for psychophysical data where multiple interacting factors influence behavior.

Statistical significance testing

To assess whether each factor contributed significantly to the model, we performed bootstrap resampling (Tibshirani & Efron, 1993) with 500 iterations. For each bootstrap sample, we re-computed sensitivity scores, obtaining a distribution of values for each predictor. The statistical significance of each predictor's contribution was evaluated using a test against zero, assuming normality.

Additionally, we performed pairwise comparisons between predictors to determine whether their contributions differed significantly. For each pair (i, j), we computed the distribution of the difference in sensitivity scores, and the significance of these differences was assessed using the same test via a normal approximation.

Results

We investigated the differential influence of conscious and unconscious priming on identity and category recognition within a binocular rivalry paradigm (Figure 1). Participants performed a name–picture verification task in which primes—either words or images—preceded target stimuli depicting faces or animal bodies, presented in the left or right visual field. Each target was paired with 4 possible labels: identity-matching, identity-mismatching, category-matching, or category-mismatching. The design included 4 prime conditions (relevant image, irrelevant image, relevant word, and irrelevant word),

thereby manipulating both recognition level and prime relevance. This fully factorial setup enabled us to assess the specific contribution of each factor.

As shown in previous studies (Navab Kashani et al., 2025), priming exerts differential effects on identity and category recognition under conscious and unconscious conditions, as assessed using a binocular rivalry paradigm. Under conscious priming, a clear dissociation emerged: identity recognition was facilitated (Figure 2A; $PI_{id} = -104.5 \pm 38.4$ ms, $P = 0.015$), with faster RTs for relevant compared to irrelevant primes, indicating positive priming. In contrast, category recognition was impaired (Figure 2A; $PI_{ca} = 57.7 \pm 20.4$ ms, $P = 0.028$), with slower RTs relative to irrelevant primes. This divergence (Figure 2A; $\Delta PI_{(ca-id)} = 162.2 \pm 42.7$ ms, $P = 0.002$) was robust and observed in nearly all participants (Figures 3B and 3C; 18/22), highlighting opposing priming effects under conscious conditions. No such dissociation was detected under unconscious priming, suggesting that these effects are contingent on awareness.

This pattern implies that identity and category recognition rely on distinct neural mechanisms, differentially modulated by conscious perception.

Under unconscious priming, the mean PI showed little differentiation between identity and category recognition (Figure S1: Unconscious: $\Delta PI_{ca-id} = -14.2 \pm 27.0$ ms, $P = 0.520$). To further elucidate these processes, we explored additional factors—prime type (word vs image), laterality (left vs right visual field), and target stimuli (faces vs animals)—that offered insights into the underlying brain systems and their condition-specific interactions.

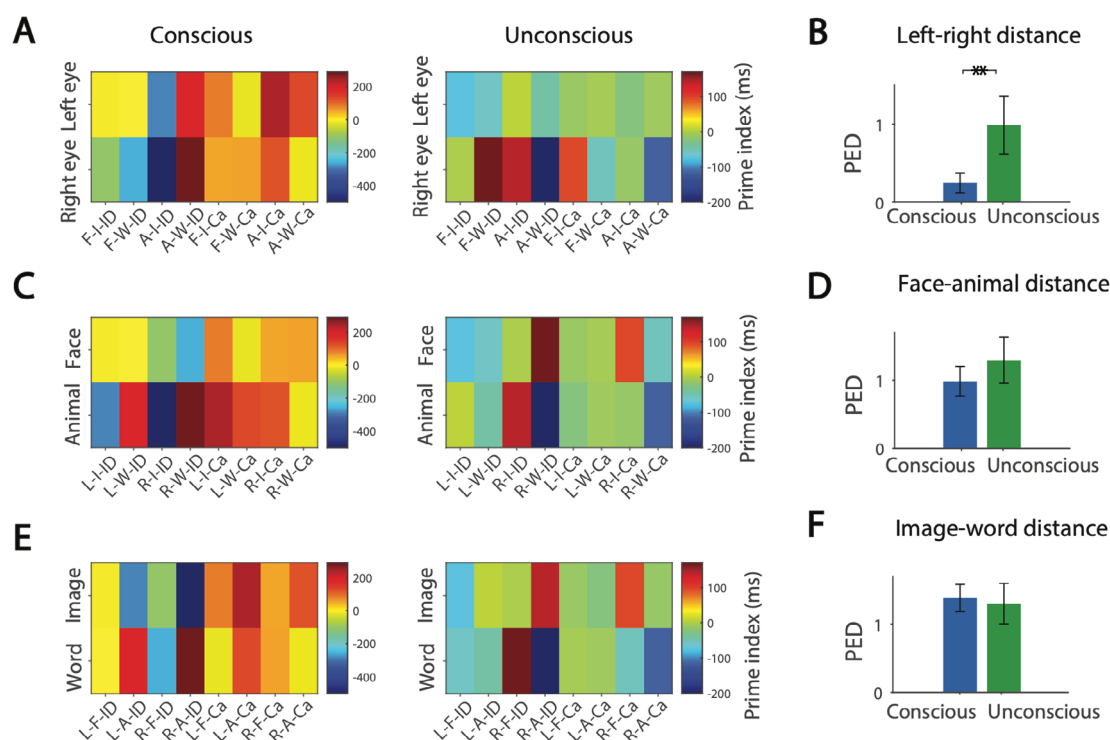


Figure 3. Priming effects across conscious and unconscious conditions

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A, C, and E) Scaled color image plots compare PI values between conscious and unconscious conditions across three dimensions: (A) visual field (L: left, R: right), (C) target type (F: face, A: animal), and (E) prime type (I: image, W: word), with recognition levels (ID: identity, Ca: category) nested within each; B, D, and F) PED, calculated as 1 minus the correlation coefficient between conscious and unconscious PIs, quantifies dissimilarity for each pairwise condition: (B) visual field, (D) target type, (F) prime type.

Note: Error bars represent SEM.

Hemispheric modulation of conscious and unconscious priming in identity and category recognition

Although overall priming effects were absent under unconscious conditions, examining hemispheric processing revealed distinct patterns between conscious and unconscious states. We assessed these states across prime type (word vs image), target type (face vs animal), and visual field (left vs right) using the PED metric—defined as 1 minus the correlation coefficient between conscious and unconscious PIs—to quantify condition-specific dissimilarities (Figure 3).

Under unconscious priming, effects were observed only when stimuli appeared in the right visual field (left hemisphere), mirroring the pattern seen in conscious conditions (Figure 3A). A significant PED difference between visual fields was detected (Figure 3A; $\text{PED}_{\text{un-con-co}} = 0.26 \pm 0.10$, $P = 0.030$), indicating that hemispheric asymmetries modulate priming differently depending on awareness. In contrast, comparisons across target stim-

uli and prime type revealed no significant differences (Figures 3C-F: face vs animal; $\Delta \text{PED}_{\text{un-con-co}} = 0.08 \pm 0.10$, $P = 0.480$; image vs word: $\Delta \text{PED}_{\text{un-con-co}} = -0.06 \pm 0.10$, $P = 0.540$).

These results indicate that unconscious priming effects are not uniformly absent but depend on specific processing pathways, with left-hemispheric dominance distinguishing conscious from unconscious processing. Unlike target type and prime type, which showed consistent effects across awareness, visual-field-dependent differences highlight a neural substrate critical for recognition, providing insight into how awareness and hemispheric specialization interact to shape priming effects.

Awareness modulates identity priming in opposite directions

To evaluate the scope and magnitude of unconscious priming, we focused on identity recognition, which involves slower processing and greater neural complexity than category recognition. We analyzed eight conditions,

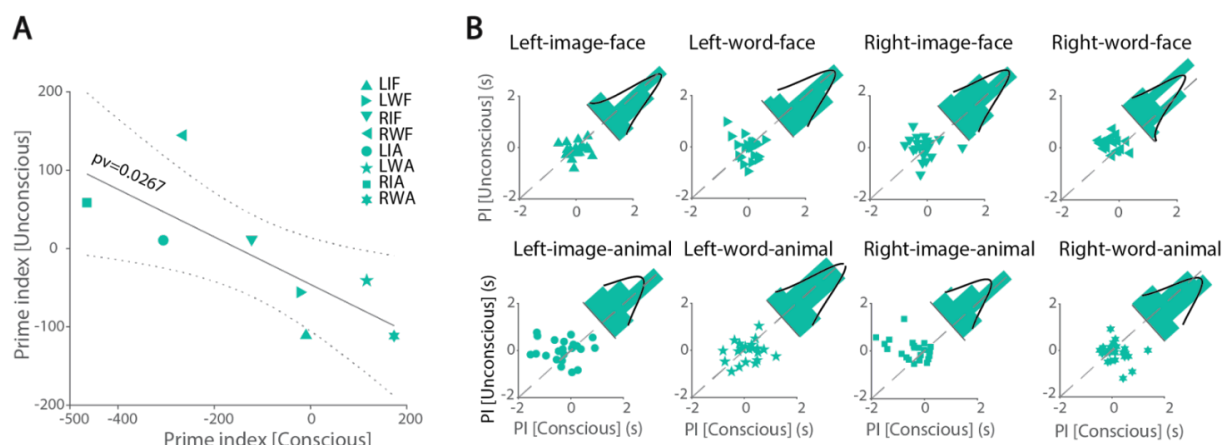


Figure 4. Divergent conscious and unconscious priming effects on identity recognition

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A) Scatter plot of PI values under conscious vs unconscious conditions at the identity level, across 8 conditions combining visual field (left/right), prime type (image/word), and target type (face/animal); B) Scatter plots and histograms depict PI (RT difference between relevant and irrelevant primes) for all participants across the 8 conditions, labeled by factor combinations (e.g. left-image-face), in conscious and unconscious states.

Note: Error bars represent SEM. The unit of time in the figure is second (s).

combining prime type (image vs word), visual field (left vs right), and target type (face vs animal) at the identity level, and assessed their PI under conscious and unconscious states (Figure 4A). A regression analysis revealed a significant negative correlation between conscious and unconscious PI values (Figure 4: $r = -0.70$, $P = 0.026$), indicating opposite effects: positive priming in conscious conditions (faster RTs) was associated with negative priming in unconscious conditions (slower RTs), and vice versa. No such relationship emerged for category recognition ($r = 0.07$, $P = 0.860$). Mean PIs and standard errors of the mean (SEMs) across these conditions are detailed in Table 1, highlighting this divergence.

This inverse pattern at the identity level—positive conscious priming shifting to negative unconscious priming, and negative priming shifting to positive priming—suggests distinct neural mechanisms underlie these states (Figure 4B). Inter-subject variability across conditions further underscores this dissociation, reinforcing the condition-dependent nature of priming effects on identity recognition. Each scatter plot displays the PI for each participant across the eight possible condition combinations derived from three factors: visual field (left/right), prime type (image/word), and target stimulus (face/animal). Each point represents a participant's mean PI for a given combination, with the x-axis showing conscious and the y-axis showing unconscious conditions. Deviations from the diagonal (unity line) indicate how awareness modulates the priming effect: points above the line reflect stronger priming under unconscious conditions, whereas points below the line reflect stronger

priming under conscious conditions. The accompanying histograms illustrate the inter-subject distributions of PI values for each condition, showing variability and consistency across participants.

Modulation of identity recognition by visual field, prime type, and target type

We examined how prime stimuli modulate RTs in identity recognition trials, focusing on visual field effects. Under conscious priming, left visual field shows did not reach significance (Figures 5A, 5B, 5C, and 5D), while right visual field (left hemisphere) presentation significantly influenced the PI across multiple conditions: image primes with faces (Figure 5E, right - image-face (RIF): $\Delta PI_{con} = -122.3 \pm 86.0$ ms, $P = 0.028$), image primes with animals (Figure 5F, right - image - animal (RIA): $\Delta PI_{con} = -464.6 \pm 128.0$ ms, $P = 0.002$), and word primes with faces (Figure 5E, Right - word - face (RWF): $\Delta PI_{con} = -264.8 \pm 61.0$ ms, $P = 0.001$), showed positive priming (faster RTs), while word primes with animals (Figure 5H, RWA: $\Delta PI_{con} = 173.6 \pm 92.5$ ms, $P = 0.094$) trended toward negative priming (slower RTs), though non-significant. This finding suggests that the prime and target types shape the direction and strength of the conscious priming effect.

On unconscious conditions, priming effects were limited to two cases: image primes with faces in the left visual field (Figure 5A, LIF: $\Delta PI_{uncon} = -110.9 \pm 52.0$ ms, $P = 0.045$) and word primes with faces in the right visual field (Figure 5G, RWF: $\Delta PI_{uncon} = 144.5 \pm 56.0$ ms,

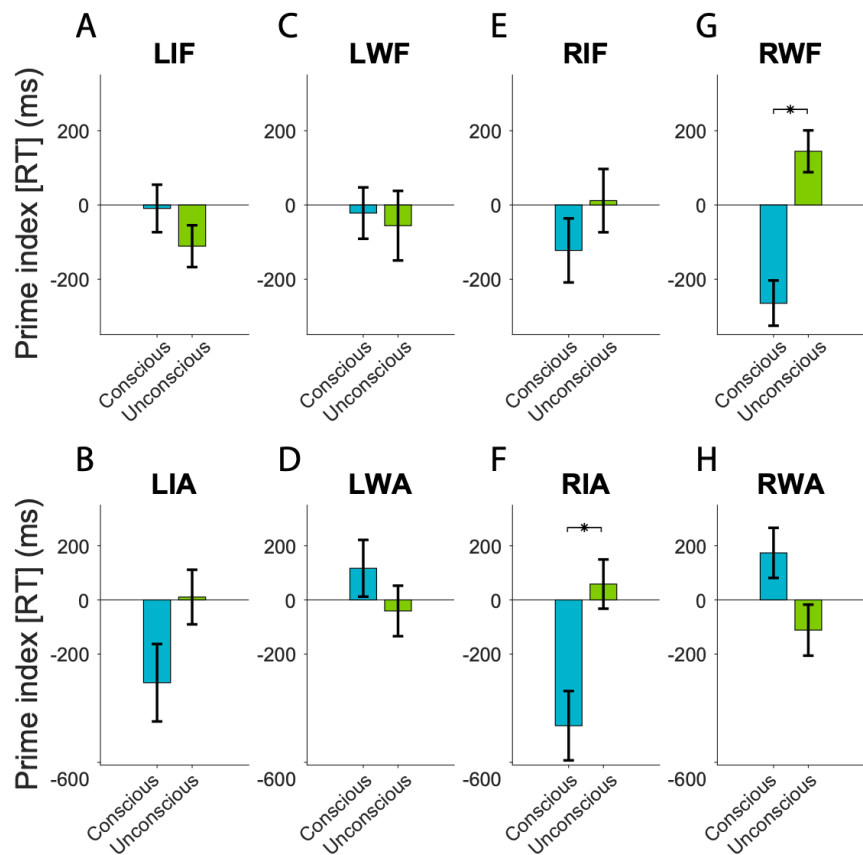


Figure 5. Conscious and unconscious priming effects on identity recognition

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A) Left - image-face, B) Left - image-face, C) Left - word - face, D) Left - word - animal, E) Right - image-face, F) Right - image - animal, G) Right - word - face, H) Right - word - animal.

Note: Bar plots depict the PI across 8 conditions – combinations of visual field (right/left), prime type (image/word), and target type (face/animal) – under conscious and unconscious states at the identity level. Error bars represent SEM.

$P=0.028$). Other conditions lacked significance (e.g. [Figure 5F](#), RIA: $\Delta PI_{uncon} = 58.5 \pm 90.9$ ms, $P > 0.05$). Notably, word primes with faces in the right visual field showed a significant conscious-unconscious divergence ([Figure 5G](#): RWF: $\Delta PI_{(uncon-con)} = 409.4 \pm 87.2$ ms, $P = 0.002$), with facilitation in conscious states and impairment in unconscious states for most participants.

These results underscore dissociable priming effects on identity recognition across awareness states. The consistent modulation of the right visual field—especially with face targets—highlights a left-hemispheric role in differentiating conscious from unconscious processing, suggesting distinct neural mechanisms underlying these effects.

Table 1. Average prime index and d' ($\pm$ SEM) for each condition on identity recognition

Conditions	LIF	LWF	RIF	RWF	LIA	LWA	RIA	RWA
Conscious	-9.34±63.9	-21.8±69	-122.38±86	-264.8±61	-305.9±143	116.8±104.7	-464.6±128	173.6±92.5
Unconscious	-110.9±52	-55.75±93.7	11.68±85	144.54±56	10.5±100.8	-40.7±93	58.5±90.9	-111.6±9

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Abbreviations: LIF: Left - image-face; LWF: Left - word - face; RIF: Right - image-face; RWF: Right - word - face; LIA: Left - image - animal; LWA: Left - word - animal; RIA: Right - image - animal; RWA: Right - word - animal

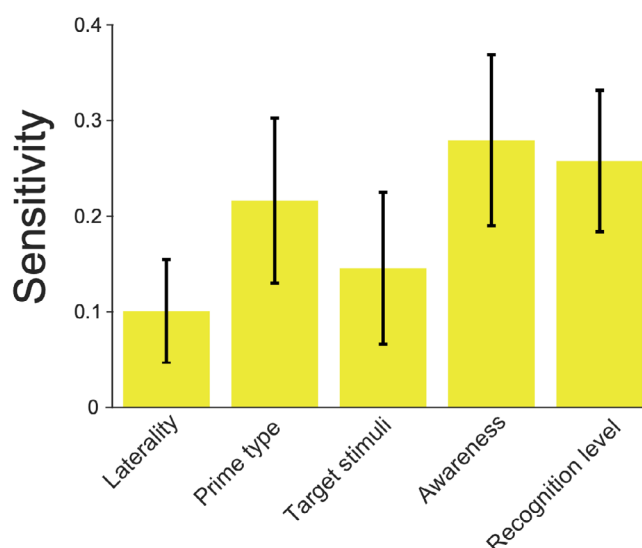


Figure 6. Feature sensitivity analysis for predicting the prime index

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Contributions of factors to prime strategies

To measure the contributions of different factors to prime strategies, we ran a nonlinear regression analysis (see Methods) and computed contributions using sensitivity scores. The sensitivity score quantifies the degree to which variations in each factor influence the response variable, with higher scores indicating greater sensitivity and a larger contribution to the prediction (Figure 6). These scores provide a clear indication of each factor's contribution to the overall model, allowing us to assess its relative importance in shaping prime strategies.

The analysis revealed distinct contributions of different experimental factors to the prediction of the prime index, measured as the RT difference between the prime and the irrelevant prime. The mean sensitivity scores ($\pm$ standard deviation) for each factor were as follows: laterality (0.1008 ± 0.0541), prime type (0.2164 ± 0.0864), target stimuli (0.1458 ± 0.0794), awareness (0.2794 ± 0.0895), and recognition level (0.2577 ± 0.0739). Among these, “awareness” exhibited the highest contribution, followed closely by recognition level, while laterality had the lowest contribution.

This bar plot shows the contributions of different experimental factors to predicting the Prime Index using a SVR model. The x-axis represents the analyzed factors: laterality (left/right visual field), prime type (image/word), target stimuli (face/animal), recognition level (identity/category), and awareness (conscious/unconscious). The y-axis shows the normalized sensitivity scores, indicating the relative influence of each factor

on the Prime Index. Bars represent the mean sensitivity scores across 500 bootstrap resampling iterations, with error bars denoting standard deviations. Statistical significance was assessed through P-values comparing each factor's contribution against zero and pairwise comparisons between factors, providing insights into their relative importance.

Statistical tests assessing whether each factor's contribution significantly differed from zero showed that all factors, except for target stimuli ($P=0.0517$; which is near significant), reached statistical significance: laterality ($P=0.0393$), prime type ($P=0.0101$), awareness ($P=0.0017$), and recognition level ($P=0.0009$). These results indicate that most factors played a meaningful role in explaining variations in the prime index.

Pairwise comparisons between factors revealed no statistically significant differences at the conventional ($\alpha=0.05$) level. However, some comparisons approached significance, including those between laterality and awareness ($P=0.0568$) and laterality and recognition level ($P=0.0578$). The remaining pairwise comparisons yielded $P>0.1$, suggesting comparable contributions among the factors.

Overall, these findings highlight the importance of awareness and recognition level in predicting the prime index, with prime type also playing a notable role. The lower contributions of laterality and target stimuli suggest a more limited influence on RT differences.

Discussion

The present study provides evidence for a context-dependent prime aftereffect, showing that conscious and unconscious priming influence visual recognition. Using a binocular rivalry paradigm (Navab Kashani et al., 2025), conscious primes led to improved identity-level recognition. In contrast, category-level recognition suffered, whereas unconscious priming failed to produce such differential effects. Moreover, we observed a left-hemisphere bias between the conscious and unconscious conditions. This lateralization may reflect the specialization of cortical areas involved in detailed visual processing and linguistic comprehension. Notably, we observed a negative correlation between conscious and unconscious perception in identity recognition tasks, underscoring that visual perception can vary across conditions within the priming paradigm.

The experimental design allows exploration of factors including global versus detail processing, the influence of image versus word analysis, functional hemispheric asymmetries, and object familiarity. This approach is founded on the idea that distinct neural substrates in the human brain are utilized for the recognition of categories and identities.

Categorization is typically faster than identity recognition (Dehaqani et al., 2016; Rosch et al., 1976) and is mainly supported by feedforward pathways in the visual system (Lamme & Roelfsema, 2000; Mohsenzadeh et al., 2018; Serre et al., 2007). This initial feedforward sweep rapidly organizes visual features but alone does not generate awareness (Jehee et al., 2007; Scholl et al., 2014; Stienen et al., 2012; Thorpe et al., 1983). While feedforward processing underlies fast, pre-attentive, unconscious vision, recurrent processing is slower and essential for attentive vision and conscious awareness (Lamme & Roelfsema, 2000; Salin & Bullier, 1995; Schmidt et al., 2011). Recurrent connections integrate detailed information over time, enabling more complex object recognition at the identity level compared to the categorization level (Hochstein & Ahissar, 2002; Jehee et al., 2007). This sustained integration supports conscious perception and contextual interpretation (Hochstein & Ahissar, 2002; Pollen, 1999). Our findings indicate that conscious priming mainly influences recurrent, detail-oriented processing, whereas feedforward global processing is less affected by either conscious or unconscious priming. When these pathways are consciously primed, repetition accelerates identity recognition, which requires deeper analysis (Dobbins et al., 2004; Tulving & Schacter, 1990). Conversely, conscious prim-

ing of faster categorization may slow processing due to increased informational load and neural fatigue (Grill-Spector et al., 2006). Thus, conscious awareness appears to selectively modulate feedforward and recurrent pathways, engaging distinct neural substrates for categorization and identification.

However, subliminal information that enters the brain unconsciously will have a different effect. Unconscious information cannot affect categorization, suggesting that the processing of categorization information in the feedforward path, which is unattended and unaware, does not allow sufficient time to analyze unconscious information. Conversely, the detailed processing of information, involving both forward and backward paths depending on the conditions, allows for the use of unconscious information to some extent during analysis.

The literature on unconscious priming presents conflicting views, with some studies demonstrating its influence on brain processing (Dehaene et al., 1998; Vuilleumier et al., 2002; Williams et al., 2004) and behavior (Jiang et al., 2007; Stein et al., 2020; Zhou et al., 2010). However, other research emphasizes the necessity of conscious awareness for effective information processing (Amihai et al., 2011; Moradi et al., 2005b). Moradi's article illustrates that unconscious image priming with face stimuli does not impact identification processing. In his experiments, stimuli were presented to the adapting eye, considering that most people are right-eyed, which is just one specific condition. A comprehensive map of conditions is needed to better understand brain function. Additionally, in conscious states, prime stimuli were presented for 4 seconds, leading to adaptation rather than a priming aftereffect (Moradi et al., 2005). Amihai et al. (2011) also argue that conscious awareness is required for processing gender and race, supporting our findings that these features are considered at the categorization level and their aftereffects vanish through unconscious priming.

These results can be interpreted within the framework of feedforward/feedback models of perception. According to this framework, perception is guided by top-down predictions that interact with bottom-up sensory input, with prediction errors driving the updating of internal models (Friston & Kiebel, 2009). Conscious primes likely instantiate strong, feedback-driven predictions about the upcoming stimulus, which, when correct at the identity level, reduce prediction error and speed recognition. However, when the prime mismatches the target at the category level, these specific predictions may interfere with rapid grouping into broader categories, producing

slower responses. In contrast, unconscious primes appear to trigger only weak feedforward activations, insufficient to generate sustained predictive feedback, which explains their lack of task-dependent influence (Kiefer & Spitzer, 2000; Kouider & Dehaene, 2007). This interpretation aligns with evidence that category-level information can be rapidly extracted via feedforward sweeps in inferior temporal cortex (Rosch et al., 1976; Grill-Spector et al., 2006), whereas conscious, recurrent processing is required for detailed identity-level recognition.

Although laterality effects are well documented in the literature, our analysis revealed that this factor contributed the least to predicting priming performance (Figure 6). This apparent discrepancy likely stems from the comprehensive nature of our model, which simultaneously examined multiple interacting variables—awareness, recognition level, prime type, and target domain. When such interdependent factors are jointly considered, the specific influence of visual field laterality becomes partially masked, reflecting shared variance with higher-level cognitive processes rather than an absence of hemispheric asymmetry.

Several limitations should be acknowledged. First, the current study relied exclusively on behavioral measures, preventing direct inference about underlying neural mechanisms. Future research combining this paradigm with EEG or functional magnetic resonance imaging could clarify the temporal and spatial dynamics of conscious and unconscious priming. Second, our stimulus set was limited to faces and animals; expanding to other stimulus domains (e.g. tools, scenes) would enhance generalizability. Third, although CFS effectively manipulated awareness, residual variability in suppression depth across participants may have influenced unconscious effects. Finally, the relatively small sample size limits the detection of subtle interactions among factors. Addressing these limitations could strengthen the mechanistic interpretation of awareness-dependent visual processing.

Conclusion

In sum, our study advances current theories of visual awareness by demonstrating that conscious and unconscious priming are not simply weaker versus stronger versions of the same mechanism, but qualitatively distinct processes with different effects on recognition. Conscious awareness enables flexible, context-dependent modulation, sometimes beneficial and sometimes detrimental, reflecting the integration of sensory input. Unconscious priming, in contrast, remains limited to

shallow, feedforward influences that lack such flexibility. These results not only deepen our understanding of how awareness and recognition level interact to shape perception,

The functioning of visual perception circuits varies across conditions, underscoring the joint influence of awareness and contextual factors on recognition processes.

Ethical Considerations

Compliance with ethical guidelines

The study complied with the Declaration of Helsinki. This study was approved by the Research Ethics Committee of Shahid Beheshti University, Tehran, Iran (Code: IR.SBU.REC.1398.047).

Data availability

The datasets used and analyzed during the current study are available from the corresponding author upon reasonable request.

Declaration of generative AI and AI-assisted technologies in the writing process

During the preparation of this work, the authors used ChatGPT (OpenAI) to assist with language editing, summarization, and refinement of highlights. After using this tool, the authors reviewed and edited the content as needed and took full responsibility for the published article.

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Authors' contributions

Conceptualization and methodology: All authors; Software and formal analysis: Samaneh Navab Kashani and Mohammad - Reza A. Dehaqani; Supervision, validation and resources: Reza Khosrowabadi and Mohammad - Reza A. Dehaqani; Investigation, data curation, visualization, and writing the original draft: Samaneh Navab Kashani; Project administration, review and editing: Mohammad - Reza A. Dehaqani.

Conflict of interest

The authors declared no conflict of interest.

References

- Amihai, I., Deouell, L., & Bentin, S. (2011). Conscious awareness is necessary for processing race and gender information from faces. *Consciousness and Cognition*, 20(2), 269–279. [DOI:10.1016/j.concog.2010.08.004] [PMID]
- Axelrod, V., Bar, M., Rees, G., & Yovel, G. (2015). Neural correlates of subliminal language processing. *Cerebral Cortex (New York, N.Y.: 1991)*, 25(8), 2160–2169. [DOI:10.1093/cercor/bhu022] [PMID]
- Balconi, M. (2006). Exploring consciousness in emotional face decoding: an event-related potential analysis. *Genetic, Social, and General Psychology Monographs*, 132(2), 129–150. [DOI:10.3200/MONO.132.2.129-150] [PMID]
- Berti, A., & Rizzolatti, G. (1992). Visual processing without awareness: evidence from unilateral neglect. *Journal of Cognitive Neuroscience*, 4(4), 345–351. [DOI:10.1162/jocn.1992.4.4.345] [PMID]
- Blake, R. (2001). A primer on binocular rivalry, including current controversies. *Brain and Mind*, 2, 5–38. [DOI:10.1023/A:1017925416289] [PMID]
- Brainard, D. H. (1997). The psychophysics toolbox. *Spatial Vision*, 10(4), 433–436. [DOI:10.1163/156856897X00357] [PMID]
- Breitmeyer, B., Ogmen, H., Ramon, J., & Chen, J. (2005). Unconscious and conscious priming by forms and their parts. *Visual Cognition*, 12(5), 720–736. [DOI:10.1080/1350628044000472] [PMID]
- Chien, S. E., Chang, W. C., Chen, Y. C., Huang, S. L., & Yeh, S. L. (2023). The limits of unconscious semantic priming. *Current Psychology*, 42(30), 26824–26835. [DOI:10.1007/s12144-022-03590-1] [PMID]
- Collin, C. A., & McMullen, P. A. (2005). Subordinate-level categorization relies on high spatial frequencies to a greater degree than basic-level categorization. *Perception & Psychophysics*, 67(2), 354–364. [DOI:10.3758/BF03206498] [PMID]
- Dehaene, S., Naccache, L., Le Clec'h, G., Koechlin, E., Muel-ler, M., & Dehaene-Lambertz, G., et al. (1998). Imaging unconscious semantic priming. *Nature*, 395(6702), 597–600. [DOI:10.1038/26967] [PMID]
- Dehaene, S., Naccache, L., Cohen, L., Bihan, D. L., Mangin, J. F., & Poline, J. B., et al. (2001). Cerebral mechanisms of word masking and unconscious repetition priming. *Nature Neuroscience*, 4(7), 752–758. [DOI:10.1038/89551] [PMID]
- Dehaqani, M. R., Vahabie, A. H., Kiani, R., Ahmadabadi, M. N., Araabi, B. N., & Esteky, H. (2016). Temporal dynamics of visual category representation in the macaque inferior temporal cortex. *Journal of Neurophysiology*, 116(2), 587–601. [DOI:10.1152/jn.00018.2016] [PMID]
- Dobbins, I. G., Schnyer, D. M., Verfaellie, M., & Schacter, D. L. (2004). Cortical activity reductions during repetition priming can result from rapid response learning. *Nature*, 428(6980), 316–319. [DOI:10.1038/nature02400] [PMID]
- Fang, F., & He, S. (2005). Cortical responses to invisible objects in the human dorsal and ventral pathways. *Nature Neuroscience*, 8(10), 1380–1385. [DOI:10.1038/nn1537] [PMID]
- Friston, K., & Kiebel, S. (2009). Predictive coding under the free-energy principle. *Philosophical Transactions of the Royal Society of London. Series B, Biological Sciences*, 364(1521), 1211–1221. [DOI:10.1098/rstb.2008.0300] [PMID]
- Grill-Spector, K., Henson, R., & Martin, A. (2006). Repetition and the brain: neural models of stimulus-specific effects. *Trends in Cognitive Sciences*, 10(1), 14–23. [DOI:10.1016/j.tics.2005.11.006] [PMID]
- Hochstein, S., & Ahissar, M. (2002). View from the top: Hierarchies and reverse hierarchies in the visual system. *Neuron*, 36(5), 791–804. [DOI:10.1016/S0896-6273(02)01091-7] [PMID]
- Jehee, J. F., Roelfsema, P. R., Deco, G., Murre, J. M., & Lamme, V. A. (2007). Interactions between higher and lower visual areas improve shape selectivity of higher level neurons-explaining crowding phenomena. *Brain Research*, 1157, 167–176. [DOI:10.1016/j.brainres.2007.03.090] [PMID]
- Jiang, Y., Costello, P., & He, S. (2007). Processing of invisible stimuli: advantage of upright faces and recognizable words in overcoming interocular suppression. *Psychological Science*, 18(4), 349–355. [DOI:10.1111/j.1467-9280.2007.01902.x] [PMID]
- Johnson, K. E., & Mervis, C. B. (1997). Effects of Varying Levels of Expertise on the Basic Level of Categorization. *Journal of Experimental Psychology. General*, 126(3), 248–277. [DOI:10.1037//0096-3445.126.3.248] [PMID]
- Kiefer, M., & Spitzer, M. (2000). Time course of conscious and unconscious semantic brain activation. *Neuroreport*, 11(11), 2401–2407. [DOI:10.1097/00001756-200008030-00013] [PMID]
- Kouider, S., & Dehaene, S. (2007). Levels of processing during non-conscious perception: A critical review of visual masking. *Philosophical Transactions of the Royal Society of London. Series B, Biological Sciences*, 362(1481), 857–875. [DOI:10.1098/rstb.2007.2093] [PMID]
- Kouider, S., Eger, E., Dolan, R., & Henson, R. N. (2009). Activity in face-responsive brain regions is modulated by invisible, attended faces: evidence from masked priming. *Cerebral Cortex (New York, N.Y.: 1991)*, 19(1), 13–23. [DOI:10.1093/cercor/bhn048] [PMID]
- Lamme, V. A., & Roelfsema, P. R. (2000). The distinct modes of vision offered by feedforward and recurrent processing. *Trends in Neurosciences*, 23(11), 571–579. [DOI:10.1016/S0166-2236(00)01657-X] [PMID]
- Marsolek, C. J. (1999). Dissociable neural subsystems underlie abstract and specific object recognition. *Psychological Science*, 10(2), 111–118. [DOI:10.1111/1467-9280.00117] [PMID]
- Mohsenzadeh, Y., Qin, S., Cichy, R. M., & Pantazis, D. (2018). Ultra-Rapid serial visual presentation reveals dynamics of feedforward and feedback processes in the ventral visual pathway. *Elife*, 7, e36329. [DOI:10.7554/eLife.36329] [PMID]
- Moradi, F., Koch, C., & Shimojo, S. (2005). Face adaptation depends on seeing the face. *Neuron*, 45(1), 169–175. [DOI:10.1016/j.neuron.2004.12.018] [PMID]
- Navab Kashani, S., Khosrowabadi, R., & Dehaqani, M. R. A. (2025). Opposite priming effects on identity vs. category recognition require conscious awareness. *bioRxiv*. [DOI:10.1101/2025.09.13.676001] [PMID]

- Pessoa, L., Padmala, S., & Morland, T. (2005). Fate of unattended fearful faces in the amygdala is determined by both attentional resources and cognitive modulation. *NeuroImage*, 28(1), 249–255. [DOI:10.1016/j.neuroimage.2005.05.048] [PMID]
- Pollen, D. A. (1999). On the neural correlates of visual perception. *Cerebral Cortex (New York, N.Y.: 1991)*, 9(1), 4–19. [DOI:10.1093/cercor/9.1.4] [PMID]
- Posner, M. I., & Cohen, Y. (1980). 14 attention and the control of movements. In *Advances in psychology* (pp. 243–258). Amsterdam: Elsevier. [DOI:10.1016/S0166-4115(08)61949-4]
- Posner, M. I., & Cohen, Y. (1984). Components of visual orienting. *Attention and Performance X: Control of Language Processes*, 32, 531–556. [Link]
- Rosch, E., Mervis, C. B., Gray, W. D., Johnson, D. M., & Boyes-Braem, P. (1976). Basic objects in natural categories. *Cognitive Psychology*, 8(3), 382–439. [DOI:10.1016/0010-0285(76)90013-X]
- Salin, P. A., & Bullier, J. (1995). Corticocortical connections in the visual system: Structure and function. *Physiological Reviews*, 75(1), 107–154. [DOI:10.1152/physrev.1995.75.1.107] [PMID]
- Saltelli, A., Ratto, M., Andres, T., Campolongo, F., Cariboni, J., & Gatelli, D., et al. (2008). *Global sensitivity analysis: The primer*. New Jersey: John Wiley & Sons. [DOI:10.1002/9780470725184]
- Schmidt, T., Haberkamp, A., Veltkamp, G. M., Weber, A., Seydell-Greenwald, A., & Schmidt, F. (2011). Visual processing in rapid-chase systems: Image processing, attention, and awareness. *Frontiers in Psychology*, 2, 169. [DOI:10.3389/fpsyg.2011.00169]
- Scholl, C. A., Jiang, X., Martin, J. G., & Riesenhuber, M. (2014). Time course of shape and category selectivity revealed by EEG rapid adaptation. *Journal of Cognitive Neuroscience*, 26(2), 408–421. [DOI:10.1162/jocn_a_00477] [PMID]
- Serre, T., Oliva, A., & Poggio, T. (2007). A feedforward architecture accounts for rapid categorization. *Proceedings of the National Academy of Sciences*, 104(15), 6424–6429. [DOI:10.1073/pnas.0700622104] [PMID]
- Smola, A. J., & Schölkopf, B. (2004). A tutorial on support vector regression. *Statistics and Computing*, 14, 199–222. [DOI:10.1023/B:STCO.0000035301.49549.88]
- Stein, T., Utz, V., & van Opstal, F. (2020). Unconscious semantic priming from pictures under backward masking and continuous flash suppression. *Consciousness and Cognition*, 78, 102864. [DOI:10.1016/j.concog.2019.102864] [PMID]
- Stienen, B. M., Schindler, K., & de Gelder, B. (2012). A computational feedforward model predicts categorization of masked emotional body language for longer, but not for shorter, latencies. *Neural computation*, 24(7), 1806–1821. [DOI:10.1162/NECO_a_00305] [PMID]
- Thorpe, S. J., Rolls, E. T., & Maddison, S. (1983). The orbitofrontal cortex: Neuronal activity in the behaving monkey. *Experimental Brain Research*, 49(1), 93–115. [DOI:10.1007/BF00235545] [PMID]
- Tibshirani, R. J., & Efron, B. (1993). An introduction to the bootstrap. *Monographs on Statistics and Applied Probability*, 57(1), 1–436. [Link]
- Trevelan, C. T., Sahraie, A., & Weiskrantz, L. (2007). Form discrimination in a case of blindsight. *Neuropsychologia*, 45(9), 2092–2103. [DOI:10.1016/j.neuropsychologia.2007.01.022] [PMID]
- Tsuchiya, N., & Koch, C. (2005). Continuous flash suppression reduces negative afterimages. *Nature Neuroscience*, 8(8), 1096–1101. [DOI:10.1038/nrn1500] [PMID]
- Tsuchiya, N., Moradi, F., Felsen, C., Yamazaki, M., & Adolphs, R. (2009). Intact rapid detection of fearful faces in the absence of the amygdala. *Nature Neuroscience*, 12(10), 1224–1225. [DOI:10.1038/nrn.2380] [PMID]
- Tulving, E., & Schacter, D. L. (1990). Priming and Human Memory Systems. *Science (New York, N.Y.)*, 247(4940), 301–306. [DOI:10.1126/science.2296719] [PMID]
- Vapnik, V. N. (1998). Statistical learning theory. Retrieved from: [Link]
- Vuilleumier, P., Henson, R. N., Driver, J., & Dolan, R. J. (2002). Multiple levels of visual object constancy revealed by event-related fMRI of repetition priming. *Nature Neuroscience*, 5(5), 491–499. [DOI:10.1038/nrn839] [PMID]
- Weibel, S., Giersch, A., Dehaene, S., & Huron, C. (2013). Unconscious task set priming with phonological and semantic tasks. *Consciousness and Cognition*, 22(2), 517–527. [DOI:10.1016/j.concog.2013.02.010] [PMID]
- Williams, M. A., Morris, A. P., McGlone, F., Abbott, D. F., & Mattingley, J. B. (2004). Amygdala responses to fearful and happy facial expressions under conditions of binocular suppression. *The Journal of Neuroscience*, 24(12), 2898–2904. [DOI:10.1523/JNEUROSCI.4977-03.2004] [PMID]
- Zhou, G., Zhang, L., Liu, J., Yang, J., & Qu, Z. (2010). Specificity of face processing without awareness. *Consciousness and Cognition*, 19(1), 408–412. [DOI:10.1016/j.concog.2009.12.009] [PMID]

Supplementary 1

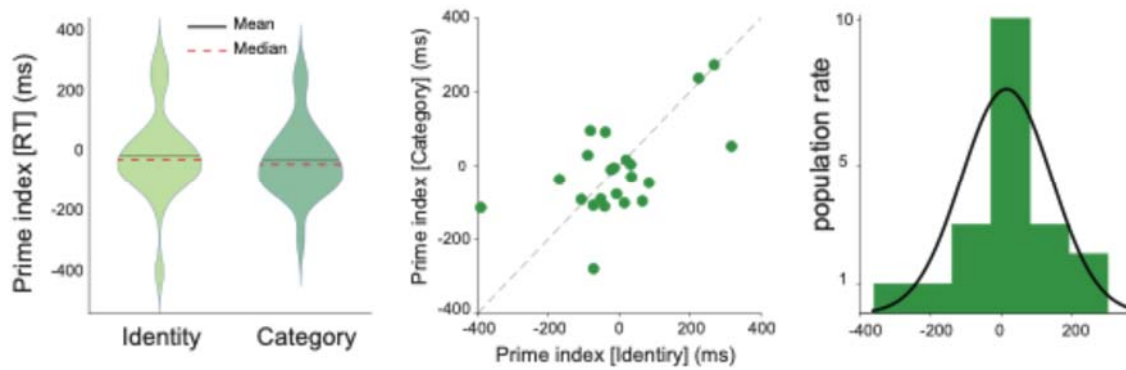


Figure S1. Unconscious priming showed little differentiation between identity and category recognition

Violin plots (left) display the distribution of reaction time prime indices (PI) for identity and category recognition under unconscious conditions. The scatter plot (middle) compares individual participants' PI values across the two recognition levels, with most points clustering near the unity line, indicating no systematic difference. The histogram (right) illustrates the population distribution of ΔPI (Category – Identity), confirming that unconscious priming effects were comparable across both recognition levels.

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