



# Motor Inhibition to Dangerous Objects: Electrophysiological Evidence for Task-dependent Aversive Affordances

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## Abstract

■ Previous work suggests that perception of an object automatically facilitates actions related to object grasping and manipulation. Recently, the notion of automaticity has been challenged by behavioral studies suggesting that dangerous objects elicit aversive affordances that interfere with encoding of an object's motor properties; however, related EEG studies have provided little support for these claims. We sought EEG evidence that would support the operation of an inhibitory mechanism that interferes with the motor encoding of dangerous objects, and we investigated whether such mechanism would be modulated by the perceived distance of an object and the goal of a given task. EEGs were recorded by 24 participants who passively perceived dangerous and neutral objects in their peripersonal, boundary, or extrapersonal space and performed either a reachability

judgment task or a categorization task. Our results showed that greater attention, reflected in the visual P1 potential, was drawn by dangerous and reachable objects. Crucially, a frontal N2 potential, associated with motor inhibition, was larger for dangerous objects only when participants performed a reachability judgment task. Furthermore, a larger parietal P3b potential for dangerous objects indicated the greater difficulty in linking a dangerous object to the appropriate response, especially when it was located in the participants' extrapersonal space. Taken together, our results show that perception of dangerous objects elicits aversive affordances in a task-dependent way and provides evidence for the operation of a neural mechanism that does not code affordances of dangerous objects automatically, but rather on the basis of contextual information. ■

## INTRODUCTION

A well-established concept in object perception research is that the passive observation of graspable objects can potentiate the possible actions that we can perform with them (Rice, Valyear, Goodale, Milner, & Culham, 2007; Grèzes, Tucker, Armony, Ellis, & Passingham, 2003; Tucker & Ellis, 1998, 2001; Ellis & Tucker, 2000; Grafton, Fadiga, Arbib, & Rizzolatti, 1997). This phenomenon is related to the concept of “affordances” (Gibson, 1979), which has been a topic of great interest in literature and has been a matter of theoretical debate (for a recent review, see Osiurak, Rossetti, & Badets, 2017). Here, we will refer to the term “affordances” to indicate the action possibilities offered to an individual from the environment, and more specifically when an individual perceives a graspable object (Borghi & Riggio, 2015; Chemero, 2003). Actions can be “afforded” or potentiated when certain object features are compatible with the motor capacities of the perceiver (Tucker & Ellis, 2004; Ellis & Tucker, 2000). For example, it has been shown that motor responses are facilitated when the object size is congruent with the shape of the hand grip (Ellis & Tucker, 2000) or

when the handle is spatially compatible with the side of the responding hand (Riggio et al., 2008; Symes, Ellis, & Tucker, 2005), even if the size and the handle position are not relevant to a given task. In recent years, the view that affordances are always activated automatically, independently from the task or context, has been challenged (for reviews, see Borghi & Barsalou, 2019; Ellis, 2018; van Elk, van Schie, & Bekkering, 2014). Much evidence has been provided, showing that activation of affordances is task and context dependent and may rely on the goals and intentions of the perceiver. Affordances are not activated in tasks that involve only processing of superficial object features, such as color (e.g., Pellicano, Iani, Borghi, Rubichi, & Nicoletti, 2010; Tipper, Paul, & Hayes, 2006). Furthermore, their activation is influenced by the context, for example, by the presence of other objects (e.g., Xu, Humphreys, & Heinke, 2015; Borghi, Flumini, Natraj, & Wheaton, 2012; Yoon, Humphreys, & Riddoch, 2010), by the scene in which they are embedded (e.g., Kalénine, Shapiro, Flumini, Borghi, & Buxbaum, 2014), and by the distance between the object and the agent (Ellis et al., 2013; Costantini, Ambrosini, Tieri, Sinigaglia, & Committeri, 2010). For example, evidence shows that affordances are activated only or to a larger extent when they are placed in a person's reachable space (Rowe,

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Haenschel, Kosilo, & Yarrow, 2017; Kalénine, Wamain, Decroix, & Coello, 2016; Cardellicchio, Sinigaglia, & Costantini, 2011; Costantini, Ambrosini, Scorolli, & Borghi, 2011; Previc, 1998).

The reachable space, also called “peripersonal” space for action, is particularly relevant for our interactions with the environment, as it represents the private area surrounding the body (Holmes & Spence, 2004; Rizzolatti, Fadiga, Fogassi, & Gallese, 1997) and delineates the immediate dimension in which we can directly act upon objects (di Pellegrino & Làdavas, 2015). By contrast, the space that is beyond this boundary, also known as “extrapersonal” space, represents the area that cannot be reached directly (Holmes & Spence, 2004; Previc, 1998). Objects that are placed in the margin of the peripersonal space rapidly attract attention, especially if they represent a threat to the individual’s safety (Graziano & Cooke, 2006). Indeed, it has been demonstrated that dangerous stimuli are detected faster and prioritized in visual selection compared to neutral ones (Zhao, 2016; Schmidt, Belopolsky, & Theeuwes, 2015; Blanchette, 2006; Smith, Cacioppo, Larsen, & Chartrand, 2003; Öhman, Flykt, & Esteves, 2001). When a dangerous object is detected in the environment, individuals need to act quickly, preparing the body to a defensive reaction, typically indicated as a flight or fight (Pichon, de Gelder, & Grezes, 2012; Brown, Hunsperger, & Rosvold, 1969).

The urgency to act in response to threats increases when the dangerous stimuli are physically closer (Pichon et al., 2012). However, according to “the threat-signal hypothesis” (Cole, Balcetis, & Dunning, 2013), dangerous objects may also lead to a perceptual bias and appear to be physically closer compared to nonthreatening ones. Similarly, threatening faces are perceived as closer in space than disgusting or neutral ones (Cole et al., 2013). Coello, Bourgeois, and Iachini (2012) showed that when an individual makes reachability judgments, a dangerous object is perceived closer when the threatening part is oriented toward the participants, compared to when it is oriented away from them.

It is believed that reachability judgments are made in relation to the action possibilities that an object offers and that they involve the mental representation of the actual reaching action and the anticipated sensory and spatial consequences (Delevoeye-Turrell, Bartolo, & Coello, 2010). This suggests that the shift of attention toward action-related features of an object may be critical to trigger the activation of affordances (Sevos, Grosselin, Brouillet, Pellet, & Massoubre, 2016; Hommel, Müsseler, Aschersleben, & Prinz, 2001). Interestingly, whereas neuroimaging (Bartolo et al., 2014; Gentile, Petkova, & Ehrsson, 2011; Delevoeye-Turrell et al., 2010; Makin, Holmes, & Zohary, 2007) and electrophysiological (Valdés-Conroy, Sebastián, Hinojosa, Román, & Santaniello, 2014; Goslin, Dixon, Fischer, Cangelosi, & Ellis, 2012; Sambo & Forster, 2009) investigations have suggested that the display of reachable objects can automatically activate

motor brain networks, other studies have challenged the notion of automaticity, showing that task goals and hand postures may have a critical modulatory influence on sensorimotor representations (Thill, Caligiore, Borghi, Ziemke, & Baldassarre, 2013). Bub and Masson (2010) showed that the compatibility effect (i.e., faster responses when the handle is aligned with the hand) emerges only when participants were required to make a reach and grasp response, but not when the task required a key press. Witt, Proffitt, and Epstein (2005) demonstrated that when participants held a tool with the intention to use it, the perceived boundary of the peripersonal space was expanded. However, when participants did not intend to reach an object, the extent of the perceived boundary of the peripersonal space was the same, with or without holding a tool. Wamain, Gabrielli, and Coello (2016) demonstrated that EEG activity over motor areas was modulated by the location of the object only when the participant was asked to make a reachability judgment, but not when performing an object discrimination task. Furthermore, it has been shown that prefrontal areas associated to top-down control can contribute to updating the neural representations of objects and contexts suitable for controlling movements so as to best pursue the person’s goals (Fogassi et al., 2005; Hamilton & Grafton, 1993). More specifically, reciprocal frontoparietal and fronto-temporal connections (Fuster, 2008; Chelazzi, Miller, Duncan, & Desimone, 1993) are critically involved in the top-down affordance processing control according to environmental contexts and attentional resources availability (Knudsen, 2007; Colby & Goldberg, 1999).

To summarize, previous research suggests that objects placed in the peripersonal space can rapidly attract our attention and that the perception of proximity might be amplified when the object is dangerous, because it represents an immediate threat to our safety. However, the processing of object motor-related information and the activation of affordances do not occur automatically when objects are reachable, but it depends on a given task goal, for example, when participants have to estimate the reachability, but not when they have to judge other object features (i.e., categorize the object). Behavioral studies on the perception of dangerous objects showed that, whereas neutral stimuli facilitate actions, eliciting faster responses, dangerous objects generate an “interference” effect that slows down the motor response, which occurs independently of the task (e.g., categorization vs. bisection) or the display of a hand prime (Anelli, Ranzini, Nicoletti, & Borghi, 2013; Anelli, Borghi, & Nicoletti, 2012). In addition, Anelli, Nicoletti, Bolzani, and Borghi (2013) investigated whether the dynamic presentation of neutral and dangerous stimuli (objects moving toward or away from the observer) would modulate the behavioral response. The results showed that responses were slower when dangerous objects moved toward the participants, suggesting that perception of dangerous objects may evoke aversive affordances, reflected in response inhibition.

However, recent EEG studies seem to indicate that this is not the case. Liu (2018), Liu et al. (2018), and Liu, Cao, Chen, and Wang (2017) investigated the ERPs in response to dangerous objects combining a motor priming paradigm (Anelli et al., 2012) with a go/no-go task. Results showed that dangerous objects elicited a larger parietal P3 (P3b) potential compared to neutral ones in the go but not in the no-go trials, which was interpreted as an indication of recruitment of additional attentional resources when perceiving dangerous objects (Isreal, Chesney, Wickens, & Donchin, 1980). In a later study, Cao, Cao, and Liu (2020) modified the perceptual salience of two stimuli in a similar motor priming paradigm combined with a shape categorization task and found a larger frontal P3 (P3a) for dangerous objects compared to safe objects, but only for objects with relatively small perceptual salience. Interestingly, the frontal N2 potential, typically associated with motor inhibition (Smith, Johnstone, & Barry, 2007; Falkenstein, Hoormann, & Hohnsbein, 1999) was similar in response to dangerous and neutral stimuli, providing no evidence that dangerous objects are automatically encoded in motor terms and elicit aversive affordances. However, all EEG studies on processing of dangerous objects (Cao et al., 2020; Liu, 2018; Liu et al., 2017, 2018) have been limited to stimuli with low ecological validity (e.g., round vs. rectangular saw blades) and a narrow range of cognitive tasks (go/no-go and shape categorization tasks).

The aim of this study was to investigate the processing of dangerous objects with a larger set of graspable stimuli and to clarify whether the location of the object and the goal of the task modulate the encoding of the object's motor properties. To test our hypotheses, we used a paradigm similar to Wamain et al. (2016) to design a controlled EEG experiment including a large set of graspable stimuli. Stimuli were rated through an on-line questionnaire by an independent sample of participants and then divided in two categories (dangerous vs. neutral). A pre-experiment session was conducted before the main experiment to determine the extent of the perceived peripersonal space for each participant. In the main experiment, the selected dangerous and neutral objects were presented in three different spaces (peripersonal, boundary, and extrapersonal) according to the subjective perceived maximum reachable point. Participants were asked to perform a reachability judgment task (RJT) and a categorization task. We predicted that the dangerous objects would elicit distinct ERPs, which would be differently modulated by the location of the objects and by the goal of the perceptual task.

More specifically, we hypothesized that the participants would pay more attention to dangerous objects especially when they are located within their peripersonal space. We predicted that this would be reflected in an enlarged amplitude of the occipital P1 potential, which is considered an index of attentional processes toward relevant stimuli attributes (Herrmann & Knight, 2001; Hillyard & Anillo-Vento, 1998; Johannes, Münte, Heinze, & Mangun,

1995). In addition, we hypothesized that a reaching action toward the object would be inhibited when the participants judge the reachability of an object (Delevoeye-Turrell et al., 2010), more so in the case of a dangerous object. Consequently, we predicted a larger amplitude of the frontal N2 potential in reachability judgments of dangerous objects, possibly when they are located close to the observer. Moreover, we predicted that activation of the link connecting the displayed object to the appropriate action toward it should be reflected in the amplitude of the parietal P3b potential (Verleger, 2020), which should be larger when the participants perceive a dangerous object (Cao et al., 2020).

## METHODS

### Participants

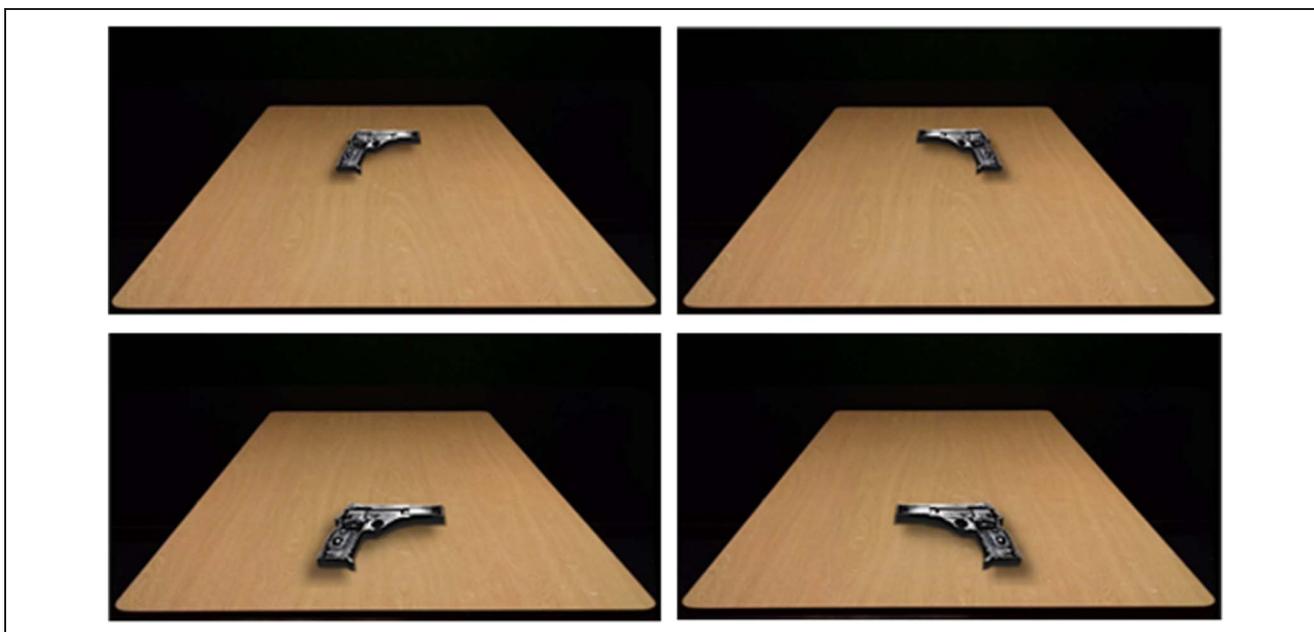
Twenty-four healthy right-handed participants (16 women and 8 men; age range = 18–28 years; mean age = 21.46 years,  $SD = 2.9$  years) took part in the experiment. The sample size was chosen according to previous EEG investigations (Cao et al., 2020; Liu et al., 2017). All participants had normal or corrected-to-normal visual acuity. The experiment was approved by the University of Stirling Ethics Committee, and all participants provided their written informed consent.

### Stimuli

Stimuli consisted of 16 color pictures of nonliving objects, half of which would be normally grasped with a precision grip and the other half with a power grip (Table 1). There were two categories (dangerous/neutral) with eight objects each. The objects were rated by an independent group of 104 participants on a 5-point Likert scale (Likert, 1932) according to harmfulness (danger/neutral), harmfulness to people (if used toward other people),

**Table 1.** Objects Used in the Main Experiment

	<i>Precision Grip</i>	<i>Power Grip</i>
Neutral objects	Pencil	Ping-Pong racket
	Dental mirror	Kettle
	Car keys	Flask
	Battery	Bulb
Dangerous objects	Syringe	Pruning saw
	Scalpel	Gun
	Fishing hook	Dagger
	Firecracker	Axe



**Figure 1.** Representation of the visual scene and the orientations of a dangerous object placed in different locations seen by the participants in the main experiment.

knowledge (familiarity), dangerousness to grasp, visual complexity, and belonging to the category of artifacts or natural objects (typicality). Paired-samples  $t$  tests revealed significant differences for harmfulness,  $t(7) = 8.73, p < .001$ ; for harmfulness to people,  $t(7) = 9.453, p < .001$ ; and dangerousness to grasp,  $t(7) = 11.789, p < .001$ , but no difference for familiarity, visual complexity, or typicality. In addition, eight pictures of plants selected on-line were used for the categorization task.

All objects were processed with Gimp 2.0 to remove the background and presented at two different orientations (i.e., graspable part to the left or to the right; see Figure 1). Each object was linearly scaled and shaded to enhance the 3-D perception. They were presented in their original shape and placed in different locations, according to individual perceived peripersonal space, in the middle of a table with a black background. The images were projected on an 86" projection screen using a projector in a dark room. The visual scene consisted of an image  $180 \times 150$  cm. All stimuli were presented at  $-35$  cm,  $-30$  cm,  $-25$  cm (peripersonal space),  $-5$  cm,  $0$  cm,  $+5$  cm (boundary space),  $+25$  cm,  $+30$  cm, and  $+35$  cm (extrapersonal space) of the perceived maximum reachable space.

## Procedure

### *Determination of Perceived Maximum Reachable Space*

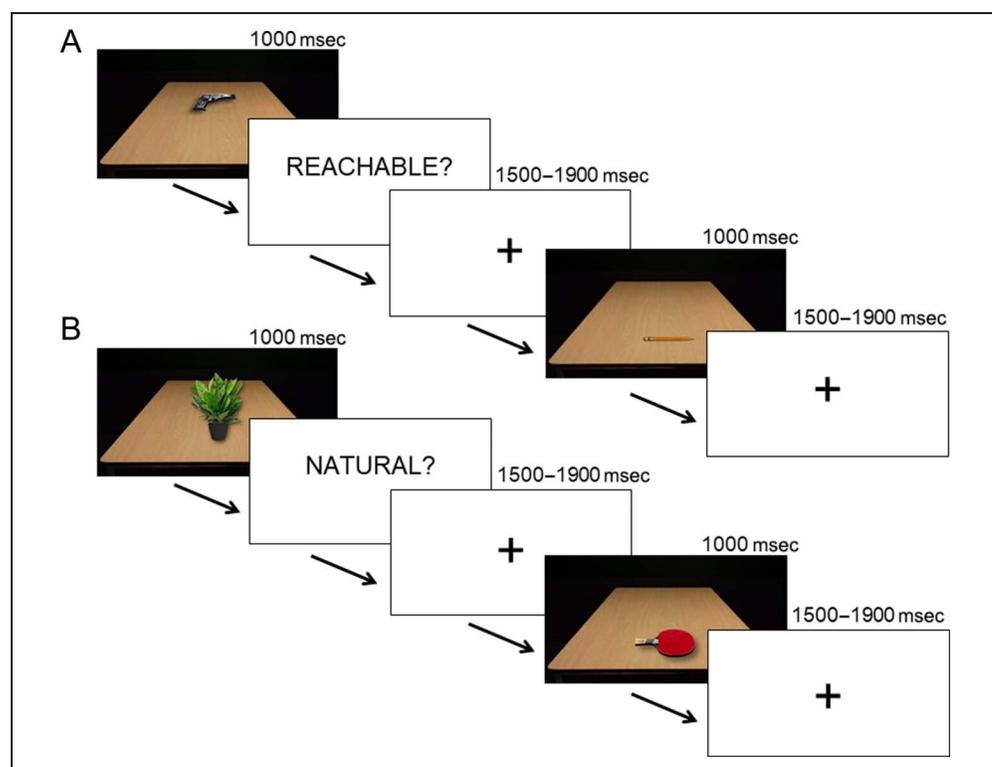
A pre-experiment session was used to determine the extent of the perceived maximum reachable space. Three different objects (a ball, a bowling pin, and a glass)

were presented on the table. Each object was linearly scaled and shaded to enhance the 3-D perception. The images were projected via E-Prime 3.0 (Psychology Tools, Inc.) on an 86-inch projection screen using a projector in the same dark room of the main experiment. The visual scene consisted of an image  $180 \times 150$  cm. The locations of the objects randomly varied between 5 cm from the edge of the table and 145 cm in steps of 5 cm (29 locations). Each object was shown 10 times in each location, which resulted in 348 trials. Participants were comfortably seated on a chair 100 cm away from the screen and asked to judge whether the object was reachable/not reachable from their position without moving or stretching the arm or the shoulder. Each image remained on the screen until a verbal response was provided. Answers were provided vocally and recorded by the experimenter. The boundary of perceived maximum reachable space was determined using a maximum likelihood method based on the second-order derivatives (quasi-Newton method) to obtain the logit regression model that best fitted the participants' reachable/unreachable space using the equation:  $y = e(\alpha + \beta X)/(1 + e(\alpha + \beta X))$ , in which  $y$  was the participant's response,  $X$  was the distance of the stimulus, and  $(\alpha/\beta)$  was the critical value of  $X$  corresponding to the transition between reachable/unreachable stimuli, thus expressing the perceived maximum reachable space (Wamain et al., 2016). The individual perceived maximal reachable space was used to select the location of the objects presented in the main experiment. The length of each participant's right arm and maximal reachable actual point (i.e., maximal point reachable on a table with the right finger) was measured.

## Main Experiment

In the main experiment, participants were first informed that they would have to perform two different tasks: a RJT and a discrimination–categorization task (DCT). Figure 2 illustrates the sequence of two trials for both RJT and the DCT. For both tasks, objects were presented centrally at different locations for 1000 msec; the interstimulus interval randomly varied between 1500–1900 msec (Wamain et al., 2016; Proverbio, 2012). The combination of category (neutral/dangerous) and location (nine locations, three for each space—peripersonal, boundary, and extrapersonal) was randomly selected for each trial. After the display of the object, a question (“Reachable?” for RJT, “Natural?” for the DCT) appeared in 20% of the trials for each block (catch trials). Participants were asked to respond as fast as possible by pressing a foot pedal either with the left or with the right foot. Questions remained on the screen until the answer was provided. In the RJT, participants indicated whether the object was reachable or not reachable from their position without moving or stretching the arm or their shoulder. Participants performed 432 trials divided in four experimental blocks. In the DCT, participants were shown also images of the plants and they were asked to indicate whether the object was natural or not. Participants performed 504 trials divided in six experimental blocks; trials in which the plants appeared (72 trials in total) were excluded from the analysis. In each task, there were 72 trials per space for neutral objects and 72 trials per space for dangerous objects. The order of the two tasks and the side of the response (left/right) was counterbalanced across participants.

**Figure 2.** Schematic representation of a sequence of two trials in (A) the RJT and (B) the DCT.



## Data Acquisition

### Behavioral Data

Behavioral data were recorded by the foot pedal box on which participants had placed their feet.

### EEG Data

EEG data were continuously recorded with Ag/AgCl electrodes from 64 scalp electrodes (Neuroscan system). The electrodes were positioned after the International 10–20 system. Vertical and horizontal eye movements were monitored using two pairs of EOG electrodes placed above and below the left eye and lateral to the external side of the eyes. EEG and EOG signals were amplified with a band-pass of 0–250 Hz.

## Data Processing and Analysis

### EEG Data

EEG data analysis was performed using BrainVision Analyzer software (Brain Products GmbH). Data were high-pass filtered at 0.05 Hz and low-pass filtered at 50 Hz. Data were rereferenced to the mean of the left and right mastoid electrodes. Ocular correction was performed using an infomax independent component analysis. Data were segmented into epochs from 500 msec before to 1500 msec after the stimulus onset. Epochs contaminated by artifacts were rejected using an automatic artifact rejection method. An epoch was rejected if the difference between the minimum and the maximum value

of a single channel exceeded 100  $\mu\text{V}$ . On average, 5.5% of epochs per condition was excluded from the analysis after artifact rejection. Data were baseline corrected (–200 msec to 0) and then averaged across participants. The ERPs were identified by visual inspection of the grand average of the different conditions during the relevant time window. ERP amplitudes were quantified by pooling the activity of neighboring electrodes within the periods of interest (for details, see Results section).

Statistical analyses of EEG data were performed by  $2 \times 2 \times 3$  ANOVAs (Greenhouse–Geisser corrected) with factors Category (dangerous vs. neutral), Task (DCT vs. RJT), and Space (peripersonal vs. boundary vs. extrapersonal). Significant interactions were further investigated via post hoc paired  $t$  tests.

## RESULTS

### Behavioral Data

In the pre-experiment data, we found that the boundary of the perceived maximal reachable space (perceived reachable space =  $49.0 \pm 7.6$  cm) corresponded to a 19.9% overestimation of the actual reachable space (reachable space =  $39.2 \pm 5.7$  cm). Although participants were required to provide a response only in the 20% of trials (catch trials),  $2 \times 3$  ANOVAs with factors Category (dangerous vs. neutral) and Space (peripersonal vs. boundary vs. extrapersonal), repeated-measures ANOVAs were conducted to assess differences between RTs separately for the RJT and the DCT. The Greenhouse–Geisser was used where the assumption of sphericity was violated and post hoc paired-samples  $t$  tests were adjusted using Bonferroni correction. In the RJT, a main effect of Space,  $F(1, 21) = 16.33, p < .001, \eta_p^2 = .438$ , revealed that participants were slower in the judgment of the boundary space

compared to the extrapersonal,  $t(21) = -6.85, p < .001$ , and to the peripersonal one,  $t(21) = 3.817, p < .001$ . In the DCT, the ANOVA did not reveal any significant main effects or interaction ( $p > .05$ ). Generally, participants were slower for dangerous compared to neutral objects in both the RJT (dangerous =  $1069.0 \pm 314$ ; neutral =  $996.8 \pm 340$ ) and the DCT (dangerous =  $1093.7 \pm 229.7$ ; neutral =  $1053.1 \pm 252.2$ ), but the comparisons did not reach the statistical significance ( $p > .05$ ).

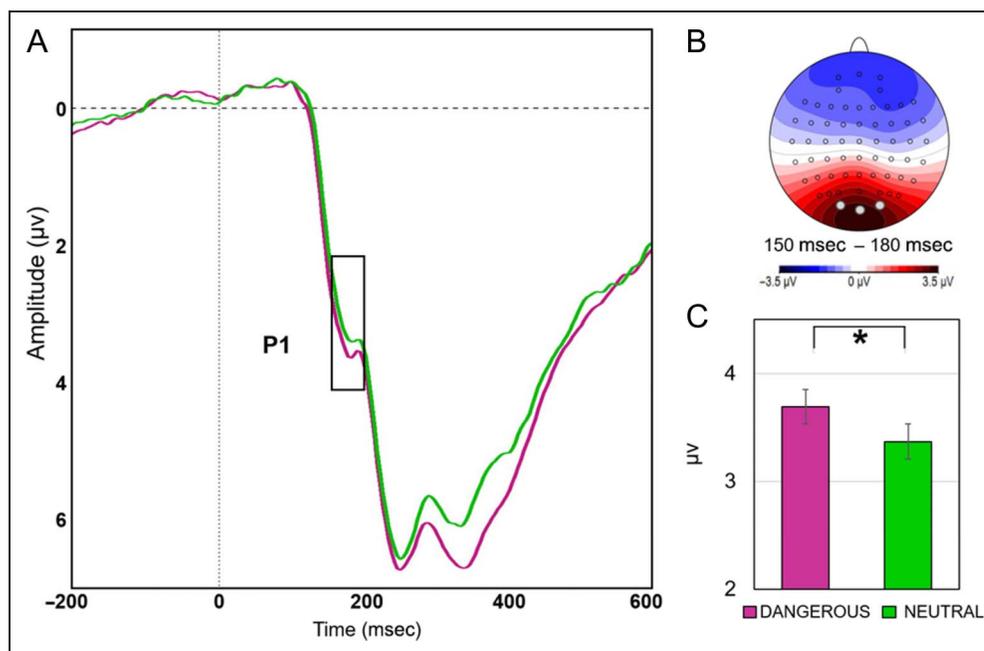
### EEG Data

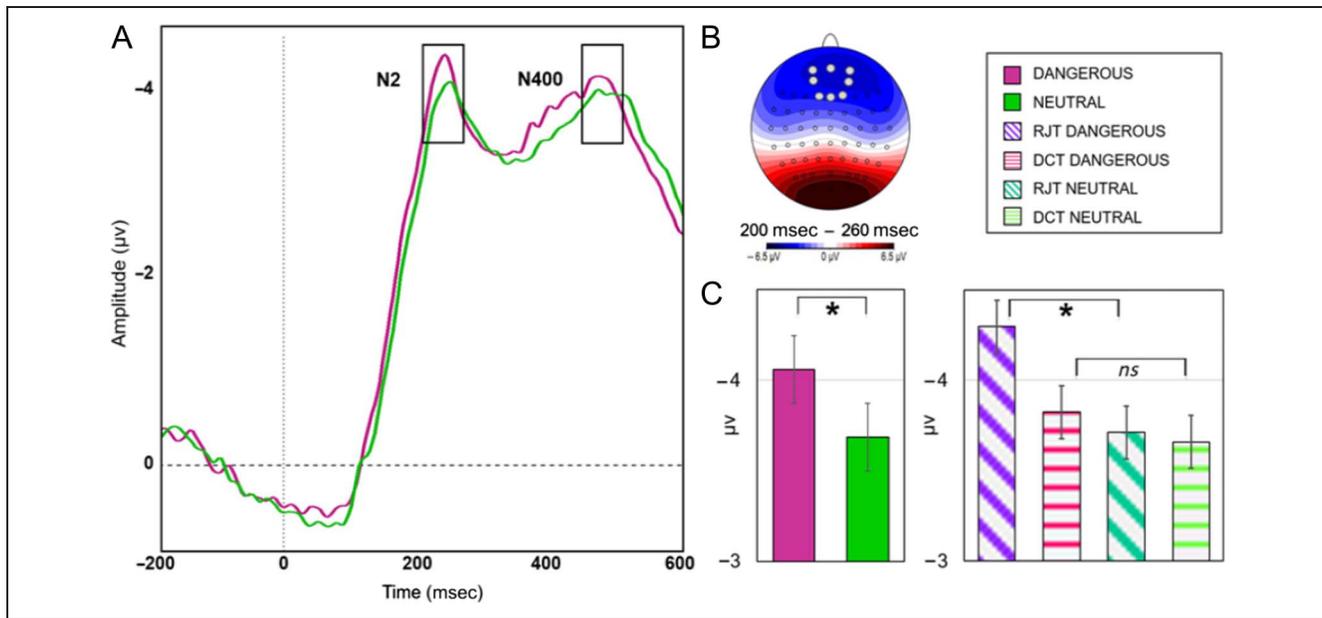
As expected, the inspection of the EEG data revealed that the onset of the stimuli elicited an occipital P1, a frontal N2, and a parietal P3b. We also identified a large frontal N400, which was also analyzed to have a complete picture of all the cognitive processes that are related to object affordances.

#### P1

The P1 was quantified as the mean amplitude of electrodes O1, O2, and Oz between 150 and 180 msec after object onset. The ANOVA revealed a main effect of Category,  $F(1, 23) = 8.61, p < .01, \eta_p^2 = .272$  (Figure 3), showing that the P1 was larger for dangerous objects compared to neutral objects. In addition, a main effect of Space,  $F(1, 23) = 3.83, p < .05, \eta_p^2 = .143$ , showed that the P1 was larger when objects were presented closer to the participant. Post hoc paired-samples  $t$  tests showed a statistically significant difference between boundary and extrapersonal space,  $t(23) = 2.76, p < .05$ , and a marginally significant difference peripersonal extrapersonal space,  $t(23) = 1.86, p = .075$ . There were no other statistically significant main effects or interactions ( $p > .05$ ).

**Figure 3.** (A) Grand average color-coded ERP waveforms. The rectangle indicates the period of interest for quantification of the P1 amplitude; time “0” indicates object onset. (B) P1 voltage scalp topography. (C) P1 amplitude as a function of category. The asterisk indicates statistical significance.





**Figure 4.** (A) Grand average color-coded ERP waveforms. The rectangle indicates the period of interest for quantification of the N2 and N400 amplitudes; time “0” indicates object onset. (B) N2 voltage scalp topography. (C) N2 amplitude as a function of Category × Task. The asterisk indicates statistical significance whereas *ns* indicates nonstatistically significant difference.

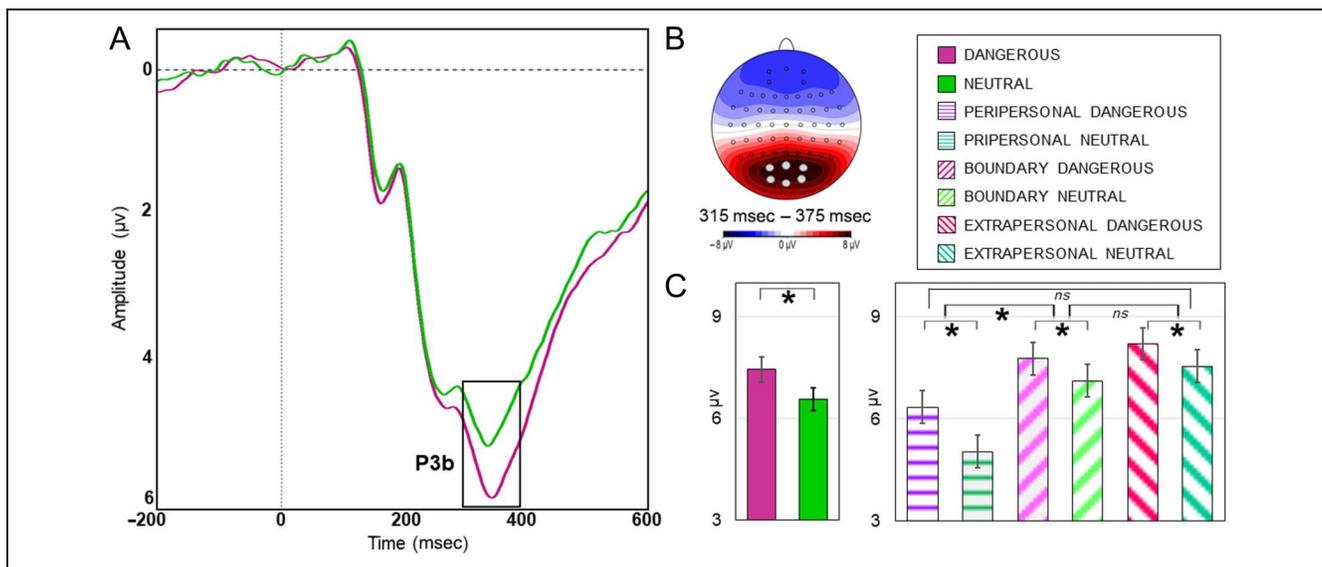
**N2**

The N2 was quantified as the mean amplitude of electrodes FPz, FP1, FP2, AF3, AF4, Fz, F1, and F2 between 200 and 260 msec after object onset. A main effect of Category,  $F(1, 23) = 5.69, p < .05, \eta_p^2 = .198$ , showed that N2 was larger for dangerous compared to neutral objects. Moreover, there was a significant Category × Task interaction,  $F(1, 23) = 6.83, p < .05, \eta_p^2 = .229$  (Figure 4). Post hoc paired-samples *t* tests showed that the N2 was larger in the RJT for dangerous compared to neutral objects,  $t(23) = -3.09, p < .01$ , whereas there was no difference in the

DCT ( $p = .503$ ). There were no other significant main effects or interactions ( $p > .05$ ).

**P3b**

The P3b was quantified as the mean amplitude of electrodes Pz, POz, P1, P2, PO3, and PO4 between 315 and 375 msec after object onset. A main effect of Category,  $F(1, 23) = 36.82, p < .001, \eta_p^2 = .616$ , revealed that the P3b was significantly larger for dangerous compared to neutral objects. In addition, there was a significant Category ×



**Figure 5.** (A) Grand average color-coded ERP waveforms. The rectangle indicates the period of interest for quantification of the P3b amplitude; time “0” indicates object onset. (B) P3b voltage scalp topography. (C) P3b amplitude as a function of Category and Category × Space. The asterisks indicate statistical significance whereas *ns* indicates nonstatistically significant difference.

Space interaction,  $F(1, 23) = 3.53, p < .05, \eta_p^2 = .133$  (Figure 5), because the difference in P3b amplitude between dangerous and neutral objects was significantly larger in the peripersonal space compared to the difference in the boundary,  $t(23) = 2.80, p < .01$ , and marginally significant to the difference in the extrapersonal space,  $t(23) = 2.05, p = .051$ .

The main effect of Space,  $F(1, 23) = 17.53, p < .001, \eta_p^2 = .433$ , indicated that the amplitude of the P3b was inversely related to perceived reachable space. More specifically, the P3b was smaller in the peripersonal space compared to the boundary,  $t(23) = -4.60, p < .001$ , or to the extrapersonal space,  $t(23) = -4.28, p < .001$ , but not in the boundary compared to the extrapersonal space,  $t(23) = -1.87, p = .074$ . Furthermore, a significant Task  $\times$  Space interaction,  $F(1, 23) = 3.69, p < .05, \eta_p^2 = .138$ , indicated that the P3b was larger in the DCT compared to the RJT when the object was placed in the peripersonal space,  $t(23) = -2.18, p < .05$ , but not in the boundary ( $p = .121$ ) or in the extrapersonal space ( $p = .727$ ). There were no other significant main effects or interactions ( $p > .05$ ).

#### N400

The N400 was quantified as the mean amplitude of electrodes FPz, FP1, FP2, AF3, AF4, Fz, F1, and F2 between 450 and 510 msec after the onset of the object. The ANOVA revealed a main effect of Space,  $F(1, 23) = 6.82, p < .05, \eta_p^2 = .229$ , showing that the N400 decreased with the perceived reachable space. The N400 was larger when the object was placed in the peripersonal space compared to the boundary,  $t(23) = 2.52, p < .05$ , or to the extrapersonal space,  $t(23) = -2.70, p < .05$ , and in the boundary compared to the extrapersonal space,  $t(23) = -2.27, p < .05$ . There were no other significant main effects or interactions ( $p > .05$ ).

## DISCUSSION

We employed high-density electroencephalography to investigate the cognitive mechanisms associated with the processing of dangerous and neutral objects in relation to their perceived distance from a passive observer. Our results show that the participants paid more attention to objects that were presented closer to them, especially to dangerous ones. Importantly, our results demonstrate that affordances of dangerous objects were task-dependent and were coded around 200 msec after object onset. Furthermore, we found evidence for higher processing demands that link the perception of dangerous objects to the representation of the relevant actions compared to neutral objects, especially when they are perceived in one's peripersonal space.

Our first hypothesis was that object location and dangerousness would modulate attentional processes. We focused our analysis on the occipital P1 potential,

which is considered an index of early visual attentional processes (Herrmann & Knight, 2001; Hillyard & Anllo-Vento, 1998; Johannes et al., 1995). The P1 was larger for objects that were presented within the observer's peripersonal space and for dangerous objects compared to neutral ones, regardless of the task. Previous studies showed that graspable objects located in the peripersonal space automatically activate attentional mechanisms that facilitate a potential interaction with proximal objects (Valdés-Conroy et al., 2014; Spence & Parise, 2010; Gallivan, Cavina-Pratesi, & Culham, 2009; Anderson, Yamagishi, & Karavia, 2002). However, in addition to object location, attention may be driven by other object characteristics, such as visually or functionally salient features (Kourtitis & Vingerhoets, 2015; Pellicano et al., 2010) and its perceived threat value. Stimuli that pose a threat to ourselves or others are detected faster than neutral stimuli (Zhao, 2016; Blanchette, 2006; Smith et al., 2003; Öhman et al., 2001) and prime our attention to enhance body responsiveness and preparation of defensive mechanisms, typically referred to as flight or fight reactions (Pichon et al., 2012; Brown et al., 1969). Accordingly, the modulation of the P1 shows that automatic allocation of attention depends on the proximity as well as the perceived dangerousness of an object regardless of the task performed.

Our second hypothesis was that a reaching action toward the object would be inhibited when the participants judge the reachability of a dangerous object. We investigated the amplitude modulation of the N2 potential, a fronto-central negativity, which is typically considered as an index of action inhibition (Huster, Enriquez-Geppert, Lavalée, Falkenstein, & Herrmann, 2013; Munro et al., 2007; Schmajuk, Liotti, Busse, & Woldorff, 2006; Kok, Ramautar, De Ruiter, Band, & Ridderinkhof, 2004; Bokura, Yamaguchi, Matsubara, & Kobayashi, 2002; Pliszka, Liotti, & Woldorff, 2000). Our analysis showed that the N2 was larger in relation to the display of dangerous objects compared to neutral ones, but it was unaffected by the proximity of the objects. This finding is in line with the notion that perception of dangerous objects may evoke aversive affordances (Anelli, Nicoletti, et al., 2013; Anelli, Ranzini, et al., 2013; Anelli et al., 2012). Importantly, this difference was significant only when participants made a reachability judgment, but not when they categorized the objects. This suggests the operation of a fast inhibitory mechanism (i.e.,  $\sim 200$  msec after object onset) that depends on the perceived dangerousness of an object. The onset of such inhibitory mechanism is consistent with previous findings (Rowe et al., 2017; Proverbio, Azzari, & Adorni, 2013; Proverbio, 2012; Proverbio, Adorni, & D'aniello, 2011). In addition, this study demonstrates for the first time that coding of object affordances for dangerous objects is not a fully automatic process, but it rather depends on contextual information. This result is similar with data on affordances evoked by neutral objects in context obtained by a large number of behavioral (reviews in Borghi & Riggio, 2015; van Elk et al., 2014)

and neurophysiological and brain imaging studies (see Fogassi et al., 2005; Thill et al., 2013, for a review).

Previous EEG research on object affordances, which showed that N2 may reflect the strength of the perception–action coupling (Wokke, Knot, Fouad, & Ridderinkhof, 2016), is larger in response to the observation of tools compared to nontools (Proverbio et al., 2011, 2013; Proverbio, 2012) and depends on the type of the grip (i.e., precision vs. power) that a person uses to handle an object (Rowe et al. 2017). Furthermore, Proverbio et al. (2011) suggested that the N2 that is elicited by the observation of tools is partially generated by the left premotor cortex and left somatosensory cortex, which is consistent with the operation of a left-lateralized “praxis” network that codes object-directed movements (e.g., Vingerhoets et al., 2012). This further corroborates our hypothesis that the N2 in this study reflects task-dependent inhibition of the motor system, which is more pertinent in the presence of dangerous objects (Anelli, Nicoletti, et al., 2013; Anelli, Ranzini, et al., 2013; Anelli et al., 2012). It should be noted that previous EEG investigations on object perception have not reported a significant effect of the object’s dangerousness on the N2 amplitude (Cao et al., 2020; Liu, 2018; Liu et al., 2017, 2018). We believe that the apparent discrepancy with the results of this study may be attributed to differences in the experimental design. These studies involved the performance of motor priming tasks, in which a dangerous or a neutral/safe object was always preceded by the display of a hand. It is plausible that the N2 did not reflect only object processing, but it was possibly influenced by a motor resonance mechanism induced by the perception of the preceding hand (Anelli, Ranzini, et al., 2013). Nevertheless, further investigation is needed to clarify the source of this apparent discrepancy.

Our third hypothesis was that the activation of the link between a perceived object and the appropriate motor response will be affected by the dangerousness of the object. To verify this, we focused on the parietal P3b potential. The P3b in this study was larger in relation to dangerous objects compared to neutral ones, and this difference was greater when the objects were located within the observer’s peripersonal space. The P3b is an endogenous cognitive potential, the functional significance of which is still a matter of debate. It has been considered to reflect context updating (Donchin & Coles, 1988) and working memory processes (Polich, 2007), although it is likely that it is indirectly associated with working memory, reflecting reactivation of stimulus–response links (Verleger 2020; Verleger, Siller, Ouyang, & Śmigasiewicz, 2017). In line with this account, which states that the P3b does not simply reflect stimulus processing mechanisms, other EEG studies suggest that it is related to response selection (Koivisto & Revonsuo, 2003; Falkenstein, Koshlykova, Kiroj, Hoormann, & Hohnsbein, 1995) and that its amplitude is enlarged by the difficulty of the task (Sawaki & Katayama, 2009; Waszak et al., 2005). Hence, our results suggest that the selection of the appropriate action toward

an object is a more cognitively demanding process for dangerous objects compared to neutral objects, especially when they are located within reachable distance. Our results extend findings from previous EEG studies on perception of dangerous objects (Cao et al., 2020; Liu, 2018; Liu et al., 2017, 2018), highlighting the importance of the reachability of a dangerous object.

Moreover, the P3b was larger when the objects were located in the observer’s extrapersonal space, suggesting the activation of a link between an object and the appropriate action was a more demanding process when the object was located outside the observer’s reach. This is consistent with previous studies that showed that activation of actions related to a graspable object is largely affected by the proximity of the object to the perceiver (Cardellicchio et al., 2011; Costantini et al., 2010). We also found a significant interaction between task and space, because the P3b was smaller in the RJT only when the object was located in the peripersonal space. This shows that linking the perception of a proximal object to the appropriate action is easier in reachability judgments compared to categorization of the object, possibly because the proximity of the object facilitates action representation. Overall, our results agree with data supporting the key role of frontoparietal and fronto-temporal connections in attention modulation and in the on-line control of visually guided movements (Knudsen, 2007; Buneo & Andersen, 2006; Andersen & Buneo, 2002; Colby & Goldberg, 1999) through the augmentation of the neural sensitivity related to the salient features of objects (Carrasco, Penpeci-Talgar, & Eckstein, 2000).

In addition to the modulation of the ERPs of main interest, we observed a frontal negativity peaking around 480 msec after stimulus onset, the amplitude of which was inversely related to the distance between object and the observer. This negativity could be considered as the N400 potential, which likely reflects the brain’s response to any type of meaningful stimulus (Kutas & Federmeier, 2000) in language, (Johnson & Hamm, 2000; Kutas & Hillyard, 1980), pictorial stimuli (Hamm, Johnson, & Kirk, 2002), mathematics (Niedeggen, Rösler, & Jost, 1999), gestures (Wu & Coulson, 2011), action–outcome relationships (Bach, Gunter, Knoblich, Prinz, & Friederici, 2009), and mismatches between objects and selected actions (Sitnikova, Kuperberg, & Holcomb, 2003). The frontal distribution of the N400 in this study is consistent with previous findings that showed that the action-related N400 has a more frontal focus compared to the language-related N400 (Amoruso et al., 2013). Previous work on object affordances suggests that recognition of action-related tool properties requires the recall of motor and semantic information, stored in a broad frontoparietal visuomotor network (Natraj, Alterman, Basunia, & Wheaton, 2018; Natraj et al., 2013; Ramayya, Glasser, & Rilling, 2010). Furthermore, Natraj et al. (2013) reported smaller N400-like amplitudes when pairs of objects were presented together with an interacting hand, which

presumably constrained the action possibilities for the observer (Natraj et al., 2018). Taking everything into consideration, it is plausible that the decreased N400 in this study when an object was located in the extrapersonal space reflects the limited action possibilities to interact with the object. This is in agreement with the view that activation of object affordances is not a purely automatic process, but rather depends on contextual information, such as the proximity of the object (Borghi & Riggio, 2015; van Elk et al., 2014).

To summarize, this study demonstrates that visual perception of a dangerous graspable object requires the engagement of greater attentional resources compared to a neutral object. Importantly, we provide evidence that aversive affordances are coded ~200 msec after the display of a dangerous object when a passive observer estimates the distance of the object on the basis of its perceived reachability. Furthermore, our results suggest that linking the perception of a dangerous object to the representation of the corresponding grasping action is a cognitively demanding process, especially when the object is located outside a person's peripersonal space. In conclusion, this study provides strong EEG evidence that challenges the notion of automaticity of object affordances, supporting the operation of a flexible mechanism that codes affordances of dangerous objects on the basis of contextual information.

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### Author Contributions

Magda Mustile: Conceptualization; Data curation; Formal analysis; Investigation; Methodology; Project administration; Resources; Software; Writing—Original draft. Flora Giocondo: Conceptualization; Data curation; Methodology; Writing—Original draft. Daniele Caligiore: Conceptualization; Methodology; Writing—Original draft; Writing—Review & editing. Anna M. Borghi: Conceptualization; Data curation; Formal analysis; Methodology; Supervision; Validation; Writing—Original draft; Writing—Review & editing. Dimitrios Kourtis: Conceptualization; Formal analysis; Methodology; Supervision; Validation; Writing—Original draft; Writing—Review & editing.

### Diversity in Citation Practices

A retrospective analysis of the citations in every article published in this journal from 2010 to 2020 has revealed a persistent pattern of gender imbalance: Although the proportions of authorship teams (categorized by estimated gender identification of first author/last author) publishing in the *Journal of Cognitive Neuroscience (JoCN)* during this period were  $M(\text{an})/M = .408$ ,  $W(\text{oman})/$

$M = .335$ ,  $M/W = .108$ , and  $W/W = .149$ , the comparable proportions for the articles that these authorship teams cited were  $M/M = .579$ ,  $W/M = .243$ ,  $M/W = .102$ , and  $W/W = .076$  (Fulvio et al., *JoCN*, 33:1, pp. 3–7). Consequently, *JoCN* encourages all authors to consider gender balance explicitly when selecting which articles to cite and gives them the opportunity to report their article's gender citation balance.

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