

Banishing the control homunculi in studies of action control and behaviour change

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For centuries, human self-control has fascinated scientists and nonscientists alike. Current theories often attribute it to an executive control system. But even though executive control receives a great deal of attention across disciplines, most aspects of it are still poorly understood. Many theories rely on an ill-defined set of 'homunculi' doing jobs like 'response inhibition' or 'updating' without explaining how they do so. Furthermore, it is not always appreciated that control takes place across different time-scales. These two issues hamper major advances. Here we focus on the mechanistic basis for the executive control of actions. We propose that at the most basic level, action control depends on three cognitive processes: signal detection, action selection, and action execution. These processes are modulated via error-correction or outcome-evaluation mechanisms, preparation, and task rules maintained in working- and long-term memory. We also consider how executive control of actions becomes automatised with practice, and how people develop a control network. Finally, we discuss how the application of this unified framework in clinical domains can increase our understanding of control deficits and provide a theoretical basis for the development of novel 'behavioural change' interventions.

Keywords: executive control, action control, behaviour change, learning

How the brain adjusts behaviour in ever-changing environments is an enduring mystery. Scientists have attributed adaptive and goal-directed behaviour to 'executive control'. This umbrella term is used for the functions of the cognitive system that allow people to regulate their behaviour according to higher-order goals or plans. This involves organising, monitoring, and altering the settings of lower-level cognitive processes such as stimulus detection and motor programming (Logan & Gordon, 2001; Miller & Cohen, 2001; Monsell & Driver, 2000; Norman & Shallice, 1986). These functions are critical in everyday life, as they allow us, for example, to resist temptations, overcome habits, or replace actions when required (e.g. when you are driving a car and a pedestrian unexpectedly crosses the street). More generally, executive control has been linked to physical and mental health, school and job success, substance dependence, personal finances, and many aspects of social behaviour (Diamond, 2013; Moffitt et al., 2011). Impairments in executive control may underlie many psychopathological disorders, including

attention deficit/hyperactivity disorder, substance abuse disorders, eating disorders, obsessive-compulsive behaviour disorders, and gambling disorders (Bechara, Noel, & Crone, 2006; Crews & Boettiger, 2009; Garavan & Stout, 2005; Nigg, 2001; Noël, Brevers, & Bechara, 2013; de Wit, 2009). The outcome of behavioural change interventions has also been linked to executive control (e.g. Nederkoorn, Jansen, Mulken, & Jansen, 2007). Thus, it is no surprise that executive control is a central component of many neurobiological models of addictions, and of impulsive and compulsive behaviours (Chamberlain & Sahakian, 2007; Crews & Boettiger, 2009; Dalley, Everitt, & Robbins, 2011; Goldstein & Volkow, 2011; Robbins, Gillan, Smith, de Wit, & Ersche, 2012).

In this article, we critically assess the current state of the executive control literature and highlight some pressing issues. We propose a unified framework of executive control, and describe how this framework can contribute to our understanding of 'behavioural change' and to the development of new behavioural change interventions that target eating behaviour, addiction, and self-control problems more generally. We focus on executive control of actions, but also consider how this work could translate to the control of thought and emotion.

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An Army Of Control Homunculi

Early research on executive control focused mostly on behavioural deficits after frontal-lobe lesions (for short reviews, see Miyake et al., 2000; Monsell & Driver, 2000). The common finding is that frontal-lobe patients experience problems with organising and regulating actions; for example, they can become impulsive, and are often unable to respond appropriately to changes in the environment. Based on such findings, it was proposed that a critical function of the frontal cortex is executive control of action and thought. After the cognitive revolution against the behaviourists in the 1950s, the concept of an 'executive controller' also became very prominent in the cognitive literature. However, in early models of cognition, control was essentially attributed to a unitary 'homunculus' who pulls the levers to regulate lower-level systems when needed (Baddeley, 1996). Around the turn of the century, many psychologists agreed that this situation was no longer tenable, because homunculus theories may explain *what* is controlled but not *how* control is exercised.

The preferred strategy to tackle the 'how' question became fractionating the executive controller and determining how distinct control functions regulate behaviour. Monsell and Driver (2000) proposed the slogan "Dissolve, deconstruct, or fractionate, the executive! Let a hundred idiots flourish!" (p.7). They argued that to know how control is exercised, we should identify the very basic processes (the 'army of idiots') that underlie control. In the last decade, great efforts have been made to deconstruct the executive controller. For example, correlational work suggests that there is both unity and diversity in executive control, with at least three distinct executive functions: switching between tasks or mental sets ('shifting'), changing and monitoring representations stored in working memory ('updating'), and suppression of irrelevant information and cancellation of inappropriate actions ('inhibition') (Miyake et al., 2000). Many studies have focused on the cognitive and neural substrates of these functions and how they interact with each other. Unfortunately, we believe that this work has not yet succeeded in banishing homunculus theories.

Too often, researchers label cognitive functions as 'executive' without questioning the mechanistic nature of the underlying processes. For example, in clinical, social, and cognitive psychology, individual or group differences in controlling actions are typically attributed to variation in the effectiveness of a single control function (e.g. 'inhibition'). Similarly, in cognitive neuroscience, prefrontal brain activation when people replace one response with another is often assumed to reflect a form of 'executive control'. However, the community seems to have fallen

into the trap of confusing tasks with mechanisms. Many processes contribute to successfully replacing an action. By referring to general constructs such as 'inhibition' (or even worse, 'executive control' or 'self-control'), we do not explain performance in complex environments—we merely re-describe it. Thus, although many researchers no longer appeal to a single control homunculus, control is often attributed to an ill-defined set of specialised 'black-box' homunculi that are assumed to do jobs like 'response inhibition' or 'updating' without explaining how they do so. We believe this theoretical strategy of focusing on general functions rather than the underlying processes is limiting progress on the control problem, because in most cases, there are no clear explanations for how the specific functions are achieved.

Furthermore, many (if not most) studies focus on action control in response to changes in the environment. However, various processes that take place on different time-scales may contribute to individual and situational differences in the efficacy of control. Preparation or pre-activation of subordinate systems that are required to detect a specific stimulus (e.g. a red light), to select a specific response (e.g. 'hit the brake pedal'), or to execute specific action (e.g. move the leg) could have a major influence; similarly, the ability to implement and maintain new rules may prove critical. Finally, action control may evolve over time. The dichotomous distinction between 'executive' and 'automatic' processes is still omnipresent in the action control literature. Automatic processes are considered to be fast, associative, emotional, effortless, and easily triggered by information in the environment, regardless of the current task goals. By contrast, executive processes are considered to be slower, effortful, rational, and goal-directed. But these may be the extremes on a continuum, and control processes that start off as deliberate and effortful can become progressively more 'automatic' through experience. By not properly acknowledging the contribution of processes such as preparation and learning, we generate an inherently limited perspective on the cognitive mechanisms behind action control.

Our proposed solution for these interlinked issues is a comprehensive theoretical framework of action control and adaptive behaviour that integrates research from different areas (see Figure 1 for a schematic representation). We will focus not only on the functions of the cognitive control system but also on the underlying cognitive processes. We define various forms of behavioural control as due to the interplay between three basic and computationally well-defined processes: signal detection, action selection, and action execution. Each process is monitored and parameters are adjusted when the outcome is

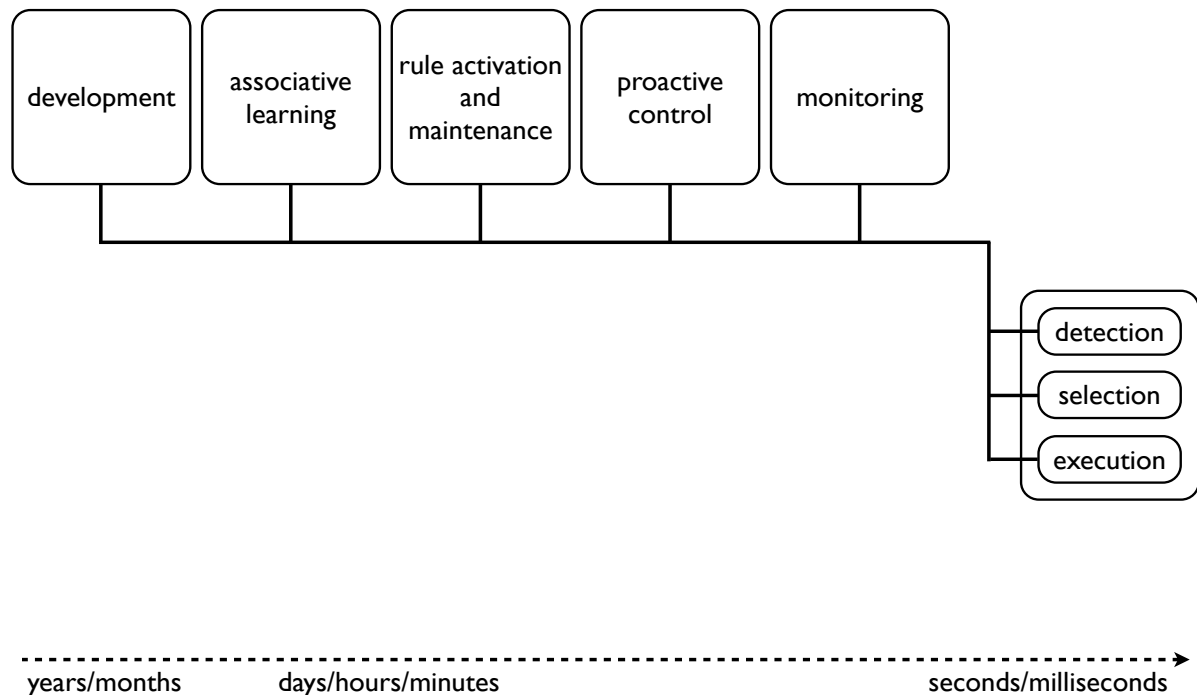


Figure 1: A schematic overview of our framework, which is inspired by Newell's Unified Theories of Cognition (Newell, 1990). We define various forms of behavioural control as an interplay between three basic and computationally well-defined processes (signal detection, action selection, and action execution), which are regulated and influenced by (sets of) processes that take place on different time scales: outcome monitoring, advance preparation, rule acquisition and maintenance, associative learning, and development. We propose that the parameters of all three basic processes (detection, selection, execution) can be influenced by these other processes. In the main text, we discuss each 'box' in more detail so as to avoid the introduction of new homunculi.

suboptimal. Furthermore, preparation will directly impact on the effectiveness of the selection and execution processes. 'Task rules', which have to be activated and maintained, will constrain the processes and adjustments. Finally, we will outline how action control and behavioural change gradually becomes automatised through practice, and more generally, how a control system can develop.

From Changes In The Environment To Changes In Behaviour

Flexible behaviour is often studied in tasks such as the stop-signal paradigm (Logan, 1994; Verbruggen & Logan, 2008c), the psychological refractory period (PRP) paradigm (Pashler, 1994; Welford, 1952), reversal learning paradigms (Izquierdo & Jentsch, 2012), Stroop tasks (Stroop, 1935), or in one of their many variants (e.g. Dodds, Morein-Zamir, & Robbins, 2011; Logan & Burkell, 1986; MacLeod, 1991; Mars, Piekema, Coles, Hulstijn, & Toni, 2007; Verbruggen, Aron, Stevens, & Chambers, 2010).

These tasks often have in common that a new action has to be selected in the context of other strong action plans (see Table 1 in Appendix for a selective overview of key paradigms). Differences in dependent variables such as response latency and error rates are usually assumed to reflect variations in the efficacy of control. For example, in most stop-signal task studies (including some of our own earlier work; e.g. Verbruggen, Liefoghe, & Vandierendonck, 2004), the stop-signal reaction time (Table 1) is assumed to reflect the duration of an executively controlled inhibition process. However, the stop-signal reaction time reflects more than the duration of an inhibitory process. Indeed, we have recently demonstrated how successful inhibition of actions depends on the detection of the stop signal and the selection of a 'stop response' (Verbruggen et al., 2010). In this section, we will further develop our framework of how humans can change actions in various situations. Based on the seminal work of Sternberg (1969) and others, we propose that at the most

basic cognitive level, action control involves three steps: signal detection, action selection, and action execution (their application is illustrated in Figure 2).

Signal detection

The first step of replacing a response is nearly always detecting the stop or change cue (e.g. a traffic light turning red or noticing an unexpected obstacle on the road). A failure to detect the signal in time could have important negative consequences. However, the contribution of detection processes to executive control of action is often neglected.

A convergence of evidence suggests that flexible behaviour depends on an interplay between two attention networks: a dorsal frontoparietal network that enables the selection of sensory stimuli, and a ventral frontoparietal network that reorients attention to important and behaviourally relevant stimuli that appear outside the focus of attention (Corbetta & Shulman, 2002; Corbetta, Patel, & Shulman, 2008). The *dorsal attention network* is thought to be involved in both stimulus-driven capture (bottom-up) and goal-directed processing (top-down) (, 2008). Precisely how these modes interact is still intensely debated (see e.g. Theeuwes, 2010, and associated commentaries). The *ventral attention network* is believed to be critical for behavioural flexibility as it allows reorienting attention from one stimulus or task towards another. Even though this network is primarily involved in stimulus-driven attention, it is activated more by weak behaviourally relevant stimuli than by salient behaviourally irrelevant stimuli (Corbetta et al., 2008). This suggests that the detection of novel signals is constrained or biased by top-down control mechanisms. For example, target detection could be controlled by an ‘attentional template’ (a representation of the relevant target or target features, such as red circle) that biases competition between sensory inputs that compete for processing resources and control of behaviour (Desimone & Duncan, 1995; Duncan, 2006). Computational work has shown how the templates could influence processing in both ventral and dorsal streams (Deco & Rolls, 2005). Pre-activation of neurons in sensory areas that code for specific stimulus features (e.g. location, colour, shape) could be the neural implementation of the templates (Deco & Rolls, 2005; Stokes & Duncan, 2013).

We believe that the literature on attention should become more integrated with the action-control literature. After all, signal detection is an important component of executive control of action: If changes in the environment are not detected quickly, this will have robust downstream effects on action control. Salinas and Stanford (2013) demonstrated that countermand-

ing (i.e. cancelling or stopping) eye movements primarily depends on the outcome of a rapid sensory detection process. Based on their computational work, they suggested that most manipulations in the countermanding task, which requires subjects to cancel eye movements (Table 1), cause changes in perceptual processes rather than inhibitory processes *per se*. The role of stimulus-detection or cue-encoding processes goes beyond action-reprogramming paradigms such as the countermanding task. Some models of interference control in paradigms such as the Eriksen flanker task (Table 1), focus on the role of spatial attention (e.g. Cohen, Servan-Schreiber, & McClelland, 1992). In the task-switching literature (Table 1), authors have argued that the cost associated with alternating between tasks is at least partly due to cue-encoding processes (Logan & Bundesen, 2003; Monsell & Mizon, 2006). Consistent with this idea, rule-switch performance in a card-sorting task in children was improved when the relevant stimulus dimensions were salient, suggesting a bottom-up attentional influence on flexible behaviour (Fisher, 2011).

Thus, we suggest that rapid detection of cues or changes in the environment is key to replacing planned or ongoing actions, and that at least some individual or situational differences in action control can be attributed to the efficacy of stimulus detection. While signal detection may seem effortless, it does require a delicate balance between selective attention and change detection: focusing on a given stimulus may lead to overly rigid behaviour while the constant reorienting of attention to novel stimuli would lead to constant distraction.

Action selection

When a change signal or cue is detected, an appropriate alternative action has to be selected. Sequential sampling models have provided a popular theoretical framework for action selection and decision-making because they explicate the various steps involved (Brown & Heathcote, 2008; Ratcliff & Smith, 2004; Smith & Ratcliff, 2004). The main assumption of these models is that action selection and decision-making involve the accumulation of noisy information about stimuli in the environment (Figure 2). Noise is present both in the environment (e.g. visibility may be reduced or the environment may be cluttered) and in the cognitive system (e.g. neurons may fire randomly and different processes may be happening at the same time), so decision-making involves collecting evidence until there is enough support for an option. More formally, accumulation of information in response counters, which keep track of the collected evidence, starts when a stimulus is detected¹. In each situation, there may be different response

options; one of them is selected when the accumulated evidence in favour of it reaches a certain threshold (Figure 2). This response option is then executed. The more noisy the information (e.g. because the stimulus is perceptually degraded), the longer it will take to reach the threshold. This will result in longer reaction times, and often, lower accuracy. This accumulation to threshold may resemble patterns of activity in certain neurons (Purcell et al., 2010; Wang, 2013). The main parameters of the selection process are the response criteria (i.e. how much information is required for a response to be selected; this is represented by the distance between the horizontal lines in Figure 2) and accumulation rate (how quickly does the information accumulate; this is represented by the slope of the tilted lines in Figure 2). Variations in these parameters can account for phenomena such as impulsive decision-making and choice errors (Ratcliff & Smith, 2004; Smith & Ratcliff, 2004), and sequential sampling models have been successfully applied to a range of decision-making tasks and to different clinical and non-clinical populations (White, Ratcliff, Vasey, & McKoon, 2010).

These sequential sampling models have been applied mostly to simple decision-making tasks in which subjects have to select a single response. But we propose that similar principles underlie the selection of actions in the context of stopping, countermanding, or replacing actions. Sequential sampling models have already been applied successfully to countermanding (Boucher, Palmeri, Logan, & Schall, 2007; Salinas & Stanford, 2013) and stop-signal tasks (Logan, Van Zandt, Verbruggen, & Wagenmakers, 2013). Boucher et al.'s model included a single go accumulator and a single stop accumulator, with two processing stages, namely 'encoding' of the countermanding signal and 'interruption' of the go process. An eye movement was successfully countermanded if stop information had accumulated quickly enough to suppress (via mutual inhibitory connections) information in the go unit and prevent it from reaching a threshold. Salinas and Stanford (2013) developed a similar rise-to-threshold model, but did not make any mechanistic assumptions about inhibitory activity; as mentioned above, they showed that perceptual processes and deceleration of information accumulation was sufficient to account for many aspects of performance. These two models had only one go accumulator. More recently, a sampling model with multiple go accumulators and a single stop accumulator has been developed to account for stopping in situations in which multiple go responses are possible (Logan et al., 2013). This model could account well for both choice and stop behaviour.

In the context of behavioural change, alternative actions must often be selected in competition with more dominant or already activated actions. Existing models could easily be modified to account for this. It has been proposed that there may be an asymmetry in mutual inhibition of units (Boucher et al., 2007) or top-down biasing of response options (Cohen, Dunbar, & McClelland, 1990). More specifically, the mutual inhibition account assumes that different response options suppress each other. In Figure 2, when evidence for the 'carrot' option accumulates, this would suppress accumulation of evidence for the 'crisps' option. Boucher et al. suggested that there may be an asymmetry in mutual inhibition, so that one response option (e.g. the carrot) may suppress the other response option (e.g. the crisp) more. The biasing account assumes that information accumulation is biased (e.g. by decreasing the distance between the starting point and the threshold; see Figure 2), making the selection of certain alternatives more likely. Others have proposed that selection of non-dominant actions is achieved by global suppression of all motor output to allow for information accumulation in the counter of the non-dominant action (Frank, 2006; Wiecki & Frank, 2013). The global suppression account overlaps with the 'circuit breaker' account of attention, which proposes that when unexpected, salient signals are detected, ongoing processes are 'automatically' interrupted to allow the cognitive system to process the new incoming information (Corbetta & Shulman, 2002; Corbetta et al., 2008).

The computational work suggests that similar selection mechanisms may be used in various situations. Cognitive neuroscience studies further support this idea. Mid-dorsolateral and ventrolateral prefrontal areas are recruited by tasks that require selection of competing actions (Bunge, 2004; Duncan & Owen, 2000), and stimulation of these areas influences action selection in different situations. We have found that magnetic stimulation of subregions within the right ventrolateral prefrontal cortex can influence attentional and action selection not only in a stop-signal task, but also in a double-response task in which participants occasionally had to execute a secondary response in addition to the originally planned response (Verbruggen et al., 2010). Similarly, Buch and colleagues demonstrated that stimulation of the ventral premotor cortex (adjacent to the posterior ventrolateral prefrontal cortex) influenced both selection and reprogramming of actions: immediately after presentation of the initial stimulus, stimulation of this area increased motor cortex excitability; however, the same stimulation reduced motor excitability when reprogramming was required (Buch, Mars, Boorman, & Rushworth, 2010). This context-dependent effect of brain stimulation is consis-

tent with the idea that similar mechanisms are involved in both programming and reprogramming actions, with the main difference being the output of the selection process (see also Mars et al., 2007). Based on these and similar findings, we propose that various forms of action control not only serve the common goal of behavioural change, they also rely on an overlapping set of selection processes (see also Mostofsky & Simmonds, 2008).

More generally, we propose that action selection and stimulus detection are governed by similar principles. The 'biased competition' account of visual attention assumes that there is competition between sources of information at many (if not all) processing stages; the main role of attention is to resolve this competition by biasing specific sources of information or specific features (Desimone & Duncan, 1995). Similar top-down bias signals can resolve competition between action options or allow the selection of non-dominant actions (see also Chun, Golomb, & Turk-Browne, 2011).

Action execution

When a response is selected or a decision has been made, the appropriate action must be executed. There can be a relatively long delay between choosing (or deciding) and acting (Schall, 2001) because a 'motor program' has to be created when an action is selected. Keele defined a motor program as 'a set of muscle commands that are structured before a movement sequence begins, and that allows the entire sequence to be carried out uninfluenced by peripheral feedback' (Keele, 1968, p.387). Creating such programs after an action is selected may contribute to the delay between choosing and acting. Consequently, the execution phase could be a final stage where individual or situational differences in action control arise. Motor control is a research area in itself, and we will not attempt to review this literature here. We will only focus briefly on three topics that are closely linked to executive control, namely the extent to which motor programs can be altered or cancelled once initiated, how they are controlled after the action is executed, and how the motor system interacts with the cognitive system.

If a motor program contains all the information needed to carry out the action, no extra control is required to complete the desired movement. This does not imply that movements can no longer be altered. In the literature on action control, researchers often make a distinction between controlled stages, which could be influenced by executive control, and ballistic stages, which must run to completion once initiated. The boundary between these two stages is called the 'point-of-no-return'. The stop-signal literature suggests that the ballistic stages must be very

brief (Verbruggen & Logan, 2009b). This idea is supported by both mathematical analyses and studies that showed that subjects could still inhibit responses that had already produced electrical activity in muscles (see Verbruggen & Logan, 2009, for a review). Thus, movements can be prepared without being executed (Schall, 2001). Not only can planned movements be cancelled, they can also be adjusted quickly if needed (e.g. Schmitz, Jenmalm, Ehrsson, & Forssberg, 2005). Thus, motor programs can still be terminated or altered quickly if new information becomes available.

Once the action has been completed, a 'reset' signal may be required to restart evidence-accumulation processes and suppress motor activity to prevent the re-execution of the same response. Indeed, in many computational models, such a reset is required to stop the system settling into a loop. In Logan and Gordon's *Executive Control of Theory of Visual Attention* (ECTVA) model, the executive system was responsible for this reset signal (Logan & Gordon, 2001). Note that there may be an overlap with the proactive suppression account discussed below, which proposes that people suppress motor activity to prevent premature responses.

In our framework, action execution is preceded by signal detection and action selection processes. This does not imply that action execution cannot interact with the preceding stages. First, an action can 'create' a signal for the next decision. To achieve certain goals, multiple movements may be required. In some situations, these movements could be 'chunked' or combined during the decision stage, when different options are selected simultaneously. But chunking may not always be possible (or preferable), so after each individual movement is completed, a new decision is required based on the imposed changes in the environment. This process would continue until the desired state or goal is achieved (Miller, Galanter, & Pribram, 1960). Second, the dual-task literature suggests that output modality has a direct influence on the decisional phase. Huestegge and Koch (2013) showed that saccades were prioritised over manual responses when participants had to make two decisions at the same time. This 'response-modality dominance pattern' could be the equivalent of the 'visual-modality dominance' pattern observed at the stimulus stage (i.e. visual modality often dominates the auditory modality when different stimuli are presented; see Huestegge & Koch, 2013). The combination of input and output modalities also influences task performance in dual-task situations (for a short review, see Huestegge & Hazeltine, 2011). For example, the dual-task cost, which is often observed in multi-task situations (Table 1), disappears after sufficient practice for some input-output combina-

tions (e.g. an auditory input and a vocal output) but not for others (e.g. visual input and verbal output). This suggests that selecting an action is not only influenced by the input, but also by the output and the input-output pairing. More generally, this shows that there may be a close link between the cognitive and motor systems, which goes beyond the cognitive system instructing the motor system which actions to perform.

Interim key points:

- Researchers should provide a more detailed account of action control because the current focus on general functions hampers theoretical and practical progress.
- We propose a framework that describes three cognitive processes underlying most forms of action control: signal detection, action selection, and action execution. Each process can be conceived as a biased competition between alternatives.
- Individual or situational differences can arise at each stage, which further highlights the need for a detailed framework.

Beyond online action control

We have outlined the core of our framework, and have illustrated how replacing an action depends on the detection of change signals, selection of an action, and the execution of that action. The detection and selection stages involve a biased competition between sources of information and response alternatives. In the following sections, we will focus on how these biases are continuously adjusted by processes that take place across different time scales. We propose that detection, selection, and execution, are influenced by monitoring, preparation, task rules maintained in memory, associative learning, and developmental changes (Figure 1). Combined, these processes lead to flexible and highly adaptive behaviour. In Figure 1, each component is depicted by a box. In the following sections, we will further unpack each box, creating our 'army of idiots' (Monsell & Driver, 2000).

Learning From Mistakes Or Unexpected Outcomes

Many things can go wrong when people try to execute a novel action. People may confuse stimuli at the detection stage, they may select the incorrect response, or they may execute the selected response incorrectly. Even when no mistakes are made, the outcome of an action may be less desirable than anticipated. Monitoring and feedback loops are a critical component of most accounts of coordinated behaviour (Ullsperger, Danielmeier, & Jocham, 2014). Within the executive control and decision-making literature, there are several detailed neuro-computational models of outcome monitoring and how this influences subsequent detection, selec-

tion, and execution processes. Three popular classes of explanation are the conflict-monitoring, error-monitoring, and outcome-evaluation accounts.

The conflict-monitoring account of Botvinick and colleagues (Botvinick, 2007; Botvinick, Braver, Barch, Carter, & Cohen, 2001) assumes that the anterior cingulate cortex, a brain area located in the medial frontal cortex, monitors for the occurrence of conflict between various response options. This brain region is often activated in situations in which prepotent responses have to be suppressed, situations in which one out of many possible but equally strong actions must be selected, situations in which errors are likely to occur, and situations with unfavourable outcomes. Based on these findings, Botvinick et al. proposed that the anterior cingulate cortex computes a 'conflict signal'. Conflict can be defined computationally as the simultaneous activation of incompatible stimulus (Egner, 2008; Verbruggen, Notebaert, Liefoghe, & Vandierendonck, 2006) or response representations (Botvinick et al., 2001). When a conflict signal is generated, task-relevant attentional or action pathways are biased, reducing the likelihood of errors or conflict on subsequent trials (Botvinick, 2007; Botvinick et al., 2001). For example, Egner and Hirsh (2005) examined control adjustments in a picture-word Stroop task. On each trial, an irrelevant word was superimposed on a task-relevant face. They found that activation in the fusiform face area, a brain area that responds strongly to face stimuli, was increased after trials on which there was competition between the face and word stimuli. This is consistent with the idea that conflict-monitoring processes bias the competition between various sources of information, enhancing detection of task-relevant stimuli and selection of task-appropriate responses.

Others have stressed the role of anterior cingulate cortex in error-driven learning and computing the likelihood of errors (Brown & Braver, 2005). Brown and Braver showed how variability in a single error-learning parameter could account for individual differences in risk aversion and in the brain's response to response conflict, error likelihood, or error consequences (Brown & Braver, 2008). Despite the differences, the conflict- and error-monitoring accounts stress that a critical aspect of optimal action control is the ability to monitor ongoing cognitive processes.

Outcome-evaluation models in the decision-making and reinforcement learning literature assume that actions can be associated with a value, which represents a prediction concerning future reward. After every action, the cognitive system compares the obtained reward with the expected reward. After reward is obtained, the action values are updated: when the reward or

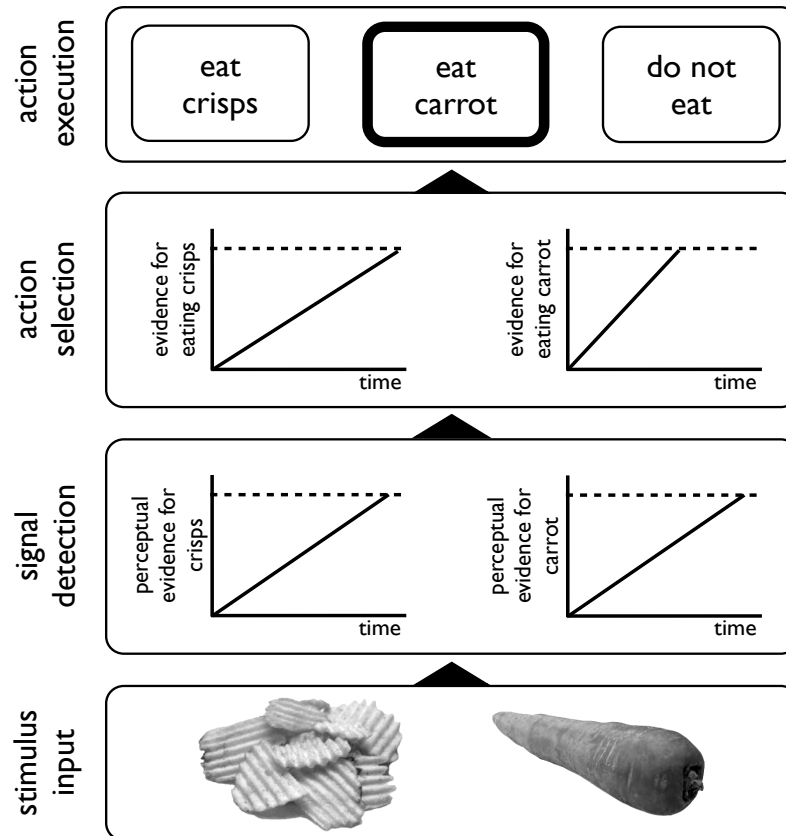


Figure 2: Action selection without a homunculus. We propose that action control involves three critical steps: signal detection, action selection, and action execution. We propose that both detection and selection can be modelled as accumulation of information towards a threshold (the dashed horizontal line). When stimuli are presented (in this example, the crisps and carrot), accumulation in the perceptual system starts, and a stimulus is perceptually encoded and attended ('signal detection') when the evidence reaches a certain threshold. When an item is encoded, evidence for possible actions starts to accumulate ('action selection'), and a response is selected when one of the thresholds is reached. Then this response is executed. In this example, the 'eat the carrot' threshold is reached first, so the person would eat the carrot. Note that for simplicity, we depict linear ballistic accumulators (Brown & Heathcote, 2008).

outcome is better than expected, the strength of the selected action is strengthened ('reinforced'); when the outcome is worse than expected, the value is decreased and the action is less likely to be selected again in similar future situations (Botvinick, Niv, & Barto, 2009; e.g. Frank & Badre, 2012). In other words, outcome-evaluation modulates action-selection biases, and this will influence how quickly information for the preferred response option will accumulate and reach the decision threshold (see above). Some have argued that value can be attached to sources of information as well, influencing stimulus detection processes (Gottlieb & Balan, 2010). Note that outcome evaluation and conflict/error-monitoring could be two sides of the same coin (Botvinick, 2007). Indeed, conflict or choice errors could be construed as aversive events, and therefore to be avoided in the future (, 2007), and activation of the anterior cingulate cortex has been linked to encoding the relationship

between an action and the reinforcement value of its outcome (Rushworth, Walton, Kennerley, & Bannerman, 2004, p.412). Ridderinkhof, Ullsperger, Crone, and Nieuwenhuis (2004) suggested that a single mechanism that signals the likelihood of obtaining a reward could account for many findings in reward-learning and conflict/error-detection literature.

Interim key points:

- Adaptive behaviour requires monitoring or evaluating the outcome of actions.
- Detection, selection and execution biases are continuously adjusted as a result of the monitoring process. This will determine how quickly a stimulus is detected and how quickly an action is selected or executed in the future.

Action Control As A Prepared Reflex

The work discussed so far may suggest that executive control is primarily 'reactive': it is only

when something changes or when something goes wrong, that the control system would 'kick in'. However, in many situations, we do not wait for unexpected events to happen. Indeed, we can adjust our behaviour 'proactively'. In contrast to the 'online' or 'reactive' control processes discussed above, proactive control refers to control processes in anticipation of an event. Proactive control can involve many things, including preparing oneself to detect a stimulus or cue, activating specific action plans, temporarily adjusting decision thresholds, and suppressing motor output to prevent premature responses (Verbruggen & Logan, 2009c). Thus, the three basic components of our framework may be influenced by preparation.

Humans can 'proactively' allocate attention. For example: in the classic Posner cuing paradigm (Posner, 1980), detection of a stimulus is enhanced by providing a central cue (e.g. an arrow) that directs attention to a specific location (e.g. the left of the screen). This has been linked to anticipatory activity in visual cortex (Kastner & Ungerleider, 2000; Luck, Chelazzi, Hillyard, & Desimone, 1997; Sylvester, Shulman, Jack, & Corbetta, 2007). Detection of stimuli or cues may also be enhanced by advance information of other features, such as shape, colour, or direction of motion (Corbetta & Shulman, 2002). Non-spatial preparatory attention is also associated with sustained activity in sensory areas (Chelazzi, Duncan, Miller, & Desimone, 1998). Such an increase in baseline activity will lead to an increased probability that the system will select the stimulus that matches the attentional template. Thus, proactively adjusting attentional settings can enhance detection of task-relevant features (especially when perceptual information is weak), and reduce interference caused by no-longer relevant features (Braver, Gray, & Burgess, 2007). Consistent with the latter idea, we have demonstrated that goal-directed cueing of the target location reduced the effect of distractors that flanked a target (Klemen, Verbruggen, Skelton, & Chambers, 2011).

Proactive action selection or movement preparation is also possible. For example, studies using a pre-cuing procedure demonstrated that individual motor actions or sets of actions can be prepared or 'primed' in advance (Rosenbaum, 1980; Rosenbaum & Kornblum, 1982). This could reduce the time required to create motor programs. Similar to attentional cueing effects, motor priming may be linked to anticipatory activation of the motor network via associations between the cue and action options. This will bias the selection and generally reduce the response time when a stimulus is presented (Meyer & Kieras, 1997). Consistent with this 'biased selection' idea, computational modelling has shown that cuing the probability of a re-

sponse or the potential payoff for a specific response, reduces the amount of information required to select the more probable response or the response associated with higher reward (Mulder, Wagenmakers, Ratcliff, Boekel, & Forstmann, 2012). Priming of a non-habitual response could also increase the probability of selecting this action in the context of other more habitual actions, or when little information is available. Note that in some situations people may proactively suppress a specific action or set of actions to prevent the premature execution of a response (Cai, Oldenkamp, & Aron, 2011; Clafey, Sheldon, Stinear, Verbruggen, & Aron, 2010; Criaud, Wardak, Ben Hamed, Ballanger, & Boulinguez, 2012; Duque, Labruna, Verset, Olivier, & Ivry, 2012). Computational work by Lo and colleagues suggests that in a countermanding task, the stopping network is activated even when no stop signal is presented (Lo, Boucher, Paré, Schall, & Wang, 2009). Thus, inhibitory motor control in stop-signal and countermanding tasks may be largely proactive in nature because it depends on control adjustments and network dynamics before a stop signal is presented (see also Wang, 2013, p.238).

Proactive control could potentially lead to a '*prepared*' or '*intention-based reflex*'. Some years ago, one of the authors (IPLM) of this article was planning to turn into a road on his bike. A car was waiting to turn into the same road on the opposite side of the street IPLM was travelling along. IPLM had priority as the car would cut across his path. IPLM made a clear signal with his extended arm just before he was about to turn. Unexpectedly, the car then immediately executed its manoeuvre, with the result that it knocked IPLM off his bike as he went around the corner. Why did this happen? We propose that when attention is proactively allocated and responses are prepared, goal-directed actions may not require much control anymore (Hommel, 2000; Logan, 1978; Meiran, Cole, & Braver, 2012); instead, actions could be activated easily by stimuli in the environment, even when they are inappropriate. Thus, when the car driver had prepared the response of turning to a high degree, IPLM's signal with the arm may have further primed the prepared reflex to the point where it exceeded threshold and was implemented as an action².

Logan (1978) demonstrated in a series of experiments that most stages in a visual-search task (including detection and response-selection) remained relatively unaffected by a concurrent load. He concluded that the preparation before the stimulus appeared was effortful, but responding was reflexive: 'the components of the task seem automatic, but the task itself is not' (Logan, 1978, p.57). Similarly, Woodman, Luck, and Schall (2007) demonstrated that a visual

working-memory load interfered with visual search only when the visual target changed from trial to trial. These findings suggest that stimulus detection, response selection, and execution may require little extra top-down support when correct task rules are properly activated. Furthermore, studies that have demonstrated that the preparation can even interfere with task-relevant or appropriate behaviour (see also the bicycle anecdote) provide further support for the 'prepared reflex' idea. Subjects are more likely to shift spatial attention to a task-irrelevant distractor when it matches a feature of the attentional template (Chun et al., 2011). Similarly, responses can be activated via instructed but unpractised stimulus-response mappings even when these mappings are task irrelevant (Cohen-Kdoshay & Meiran, 2009); however, such effects are only observed when the tasks are actually prepared and participants anticipate that they have to perform them in the near future (Liefoghe, De Houwer, & Wenke, 2013). Finally, we have found that the irrelevant distractor 'STOP' inside a go stimulus interfered with responding in stop-signal and go/no-go tasks but not in a task where participants could always respond (Verbruggen & Logan, 2009a). These findings are consistent with the prepared reflex idea: the prepared action can be triggered by irrelevant primes, even when this is not strictly required (, 2009a).

Combined, this work suggests that action control could be reflexive; paradoxically, this could even lead to a cost in some situations. But usually the main costs associated with proactive control are that this strategy requires highly reliable predictive contextual cues, it is metabolically costly, and it is capacity demanding (Braver et al., 2007). Humans usually prefer to avoid cognitive demands (Kool, McGuire, Rosen, & Botvinick, 2010), so internal costs may shift the balance between reactive and proactive control (McGuire & Botvinick, 2010). This also implies that a proactive strategy is less likely to be applied in situations with very long retention intervals between a 'warning' cue and the stimulus, as this may require too much effort. Finally, strong pre-activation of stimulus features or actions may also stand in the way of flexible behaviour in ever-changing environments. Thus, a delicate balance between proactive and reactive control is required.

The costs associated with proactive control could potentially explain some individual and situational variation. Differences in motivation (Leotti & Wager, 2010) and emotional factors (Fröber & Dreisbach, 2012) contribute to intra-individual differences in deployment of proactive control, and factors such as reward sensitivity, general intelligence, and working-memory capacity may cause inter-individual differences

(Braver, 2012; Redick & Engle, 2011). Several studies have also shown group differences. Healthy young adults seem to rely more on proactive control than both young children (Munakata, Snyder, & Chatham, 2012) and older adults (Paxton, Barch, Racine, & Braver, 2008), and proactive control seems impaired in disorders such as schizophrenia, Alzheimer, ADHD, and bipolar disorder (for review, see Braver, 2012), and in individuals who engage in delinquent and antisocial behaviours (Iselin & Decoster, 2009). These findings suggest that at least some 'control' deficits could be due to a failure to activate the control system proactively. Interestingly, training older adults on a proactive control task caused a shift from reactive to proactive control (Braver, Paxton, Locke, & Barch, 2009), suggesting that control strategies are amendable.

Interim key points:

- Executive control of actions is often proactive: the act of control takes place before the change or control signal is presented.
- When control is applied proactively, signal detection, action selection, and action execution can become a 'prepared' reflex, easily triggered by information in the environment.
- Important intra- and inter-individual differences could be due to shifts from proactive to reactive control.

Activation And Maintenance Of Action Goals And Rules

An important issue that we have not addressed so far is how the connections between input, action selection, and action execution are established. And how does the cognitive system 'know' which stimulus feature or response option to bias? The main important advance of the (mathematical) modelling framework discussed above is that ongoing processes are described in detail. However, there is still a 'homunculus' lurking: because it is the *researcher* who creates all the connections and sets up the relevant accumulators that enable a model to perform a certain task. Thus, this framework does not necessarily solve the problem of how the model could achieve this functionality in the first place. Most theoretical frameworks or models of executive control, including our framework, either explicitly or implicitly rely on 'rules' (Bunge, 2004; Logan & Gordon, 2001; Miller & Cohen, 2001; Monsell & Driver, 2000). Rules enable humans to quickly select relevant cues or stimulus features, map sensory input to action plans, and produce the motor output. Furthermore, sequential adjustments after a trial (see e.g. Hazeltine, Lightman, Schwarb, & Schumacher, 2011) and proactive control before a trial are also rule-dependent.

Thus, one could argue that rules are at the core of executive control. In this section, we will explore how rules are activated and maintained. We make an explicit distinction between a task goal and a task rule: A task goal describes what one tries to achieve, a task rule specifies how one can achieve it. A goal will activate a rule (or set of rules). We will primarily focus on the role of task rules.

A key characteristic of adaptive human behaviour is the ability to rapidly learn action rules from instructions. For example, if instructed to tap your right foot whenever you see the symbol 'x' on this page, most likely you will be able to do this without any practice (even though you have probably never done this specific task in your life). Recently, several studies have focused on the cognitive and neural mechanisms underlying this fundamental ability. For a complete overview of this instruction-based learning literature, we direct the interested reader to two recent review articles (Cole, Laurent, & Stocco, 2013; Wolfensteller & Ruge, 2012); the review of Cole et al. also addresses the issue of language in rule-learning and control (see also Oberauer, 2009). The compositional account of instruction-based learning (Cole et al., 2013) is based on five related principles: (1) *Compositionality*, which refers to the ability to reuse representations with a variety of other representations; (2) *Immediate transfer*, which refers to the ability to apply practiced rules to novel situations; (3) *Abstraction*, which refers to the ability to group specific representations; (4) *Analogy*, which refers to the ability to recognise similarities; and (5) *Compositional hierarchy*, which refers to creating a structure in which more abstract representations modulate more concrete stimulus-action representations. Of these five principles, compositionality is key, as this can offer an elegant explanation for our remarkable ability to immediately perform tasks that we have never done before. Returning to the foot-tapping example, you may never have tapped your foot when you saw an 'x' on a page, but you may have tapped your foot in response to other cues (e.g. music) and you may have searched for a specific word or letter string in a text before; by linking these representations, you are able to perform the new x-tapping task. In other words, you would reuse and recombine existing circuits, structures or representations (see also Anderson, 2010). Support for the compositional theory and other relevant findings are discussed in Cole et al. (2013).

There are large individual differences in the ability to follow new task rules. The ability may be linked to fluid intelligence (Duncan, Schramm, Thompson, & Dumontheil, 2012). Furthermore, patients with lesions to the lateral prefrontal cortex may not always be able to produce the instructed behaviour even though they can under-

stand the instructions (Cole et al., 2013). Duncan and colleagues (Duncan, Emslie, Williams, Johnson, & Freer, 1996; Duncan, Johnson, Swales, & Freer, 1997) have referred to this phenomenon as 'goal neglect'. Verbal instructions specify an abstract 'requirement' (e.g. 'if X then tap right foot'), but these requirements have to be implemented or transferred to procedural working memory (Duncan et al., 2012; Logan & Gordon, 2001; Oberauer, 2009). For example, relevant stimulus information, response options, and output modalities should become 'biased', and contexts in which the rules are relevant specified. A failure to do so would lead to goal neglect (Duncan et al., 1996; Duncan et al., 1997).

When instructions are successfully implemented, rules have to be maintained. We have argued above there is sustained activity in brain areas that process task-relevant information, which biases the selection of information. Rules likely provide the top-down signal for this bias (Miller & Cohen, 2001; Stokes & Duncan, 2013). The popular account is that rules are maintained in working memory via persistent firing of stimulus-specific neurons in the prefrontal cortex (Curtis & D'Esposito, 2003). More generally, this persistent firing would allow temporal integration of information, which is required for many functions, including working memory and the calculation of reward rate (Curtis & Lee, 2010). However, recent findings challenge this 'persistent activation' account (Postle, 2013; Stokes & Duncan, 2013). For example, Stokes and colleagues (2013) showed that the presentation of an instruction cue triggers a sequence of high-activity patterns before settling into a stable low-activity state. They proposed that, rather than sustained activity, synaptic weight changes constitute the task-dependent rules that determine how people respond to stimuli (Stokes & Duncan, 2013). One of the main challenges is to further determine how rules are maintained in long-term and short-term memory.

Finally, in many situations, people have to switch between rules. This fundamental ability is studied in the task-switching paradigm (for reviews, see Kiesel et al., 2010; Monsell, 2003; Vandierendonck, Liefoghe, & Verbruggen, 2010). Switching from one rule to another is usually associated with a performance cost. Most agree that this switch cost reflects the time needed to encode the task cues, activate the appropriate task rules, and resolve interference caused by previous trials, although the extent to which each process contributes to the overall switch cost may vary. Cue encoding and task reconfiguration are time-consuming processes, so performance generally improves when these processes can be completed before the stimulus appears (Logan & Gordon, 2001; Mayr & Kliegl, 2000; Meiran, 1996; Rogers & Monsell, 1995).

This demonstrates the close link between preparation and rule activation/maintenance. However, not everybody agrees that people always have to switch or update rules when tasks change. Logan and colleagues argued that switching between tasks could be achieved via the retrieval of learned associations between cues, stimuli, and responses (Logan & Bundesen, 2003; Schneider & Logan, 2005), although this idea remains highly controversial (e.g. Forrest, Elchlepp, Monsell, & McLaren, 2012; Jost, Mayr, & Rösler, 2008; Monsell & Mizon, 2006). In other words, they argued that 'control' could be associatively mediated.

Interim key points:

- In our framework, rules constrain performance by providing a top-down bias for each process.
- We argue that the ability to follow novel instructions and implement new rules is strongly rooted in the past: humans constantly re-use and recombine old rules that have previously governed behaviour.

Action Control As An Associatively Learned Reflex

Historically, executive control has been pitted against automatic operations. Often, functions such as response inhibition, interference control, or task switching have been classified as 'executive', whereas other processes, such as word reading in the context of a Stroop task, have been classified as 'automatic'. In this section, we discuss how 'executive' processes may also become automatic as a consequence of practice.

It is well documented that responding to a stimulus or cue can become 'automatised' over practice (Dickinson, 1985; Logan, 1988; Schneider & Shiffrin, 1977; Shiffrin & Schneider, 1977). Shiffrin and Schneider proposed that when a stimulus and a response are consistently mapped, associations are formed, allowing actions to become automatic (Schneider & Shiffrin, 1977; Shiffrin & Schneider, 1977). Similarly, Logan (1988) suggested that every time people respond to a stimulus, processing episodes are stored. These episodes consist of the stimulus (e.g. '3'), the interpretation given to a stimulus (e.g. 'odd'), the task goal ('odd/even task'), and the response ('left'), are stored. When the stimulus is repeated, previous processing episodes are retrieved, facilitating performance if the stimulus-response (S-R) mapping is consistent. Recently, we have demonstrated that more complex forms of action control could also become 'automatised'. We found that pairing a stimulus with stopping interfered with responding to these stimuli (Verbruggen & Logan, 2008a), even after a single stop presentation (Verbruggen & Logan, 2008b; Verbruggen, Logan, Liefvoeghe, &

Vandierendonck, 2008). We attributed the behavioural slowing for old stop items to the retrieval of stimulus-stop associations, which would automatically suppress responding. Similar associatively-mediated 'control' effects have been observed in other executive control tasks. The task-switching literature suggests that stimuli can become associated with 'tasks' or 'rules' (Koch & Allport, 2006; Mayr & Bryck, 2005; Waszak, Hommel, & Allport, 2003). For example, the results of Mayr and Bryck (2005) suggest that abstract spatial translation rules can become integrated with lower-level stimulus and response codes; similarly, the results of Waszak and colleagues suggest that individual stimuli can become associated with higher-order task-representations, such as 'picture naming' (Waszak et al., 2003). Finally, several studies have shown that stimuli in tasks such as the Stroop paradigm can become associated with attentional control settings (Bugg & Crump, 2012). Based on these findings, we argue that rule-based action control can also become a 'learned reflex', triggered even when it is not required or intended at a given moment (cf. Meiran et al., 2012; Tzelgov, 1997; Verbruggen & Logan, 2009a)

We suggest that there are four non-mutually exclusive ways that associative learning could influence action control: (1) conditioned attention towards (or away from) the cues, (2) associatively mediated activation of previously non-dominant responses, (3) associatively mediated activation of abstract rule representations, and (4) by changing the hedonic and/or incentive value of stimuli.

First, associative learning could influence attentional selection. For example, Le Pelley, Beesley, and Griffiths (2011) have found that subjects looked more at cues experienced as predictive of the outcomes with which they were paired, than to cues experienced as non-predictive. Similarly, Gottlieb and Balan (2010) reviewed a single-cell recording study that showed higher sustained lateral intraparietal area (LIP) activation for predictive cues in a Pavlovian task, and suggested that attentional selection is influenced by the information value of the stimuli. These results are consistent with the attentional model of associative learning proposed by Mackintosh (Mackintosh, 1975; Pearce & Mackintosh, 2010). However, Hogarth and colleagues (Hogarth, Dickinson, Austin, Brown, & Duka, 2008) have found that participants looked more at partially predictive signals in some situations, which is consistent with the Pearce-Hall model of Pavlovian learning (Pearce & Hall, 1980; Pearce & Mackintosh, 2010). Even though there is uncertainty about the direction of the effects, it is clear that attention and associative learning can interact (albeit in various ways), and the model of Mackintosh and Pearce integrates earlier ac-

counts to reflect this (, 2010). In other words, attention can become 'conditioned' (McLaren, Wills, & Graham, 2010): attention is paid to stimuli as a consequence of past associative history, rather than because of their match with our current goals. This is also supported by some event-related potential studies. For example, Wills and colleagues have demonstrated that early attentional components were modulated by associative learning (Wills, Lavric, Croft, & Hodgson, 2007). In a similar vein, ERP work reviewed by Woodman suggests that top-down biasing of visual attention is only required when targets are new, with long-term memory taking over when objects are repeated (Woodman, 2013).

Second, a non-habitual response could become habitual after sufficient practice (Dickinson, 1985; Logan, 1988; Schneider & Shiffrin, 1977; Shiffrin & Schneider, 1977). This would further reduce the need for top-down biasing; after sufficient practice, the need for top-down biasing may even disappear altogether and people would no longer have to rely on rules or PFC representations to execute an action that was initially non-dominant. This idea is supported by the work discussed above, and by the finding that neural activation in prefrontal and other control-related brain regions is reduced after practice with consistent mappings (Chein & Schneider, 2005).

Third, the studies discussed above suggest that during practice, stimuli can become associated with task goals or the task rules that bias attentional or action selection. After practice, the goal or rule representations may become activated when a stimulus is repeated, whether this is intended or not; this would then influence subordinate processing. The stimulus-rule association idea could explain why repeating an old stop stimulus activates components of the stopping network in the ventrolateral prefrontal cortex (Lenartowicz, Verbruggen, Logan, & Poldrack, 2011), or why naming the word inside a picture-word Stroop stimulus is impaired when this stimulus was previously encountered in a picture-naming task (Waszak et al., 2003). Note that such stimulus-task effects were observed even when the response (e.g. a left key press) was the same in both tasks (Koch & Allport, 2006; Waszak et al., 2003). The main difference with the previous two options is that this third option assumes that rule-like representations that bias ongoing selection processes are still involved. In other words, this third option provides a more indirect route to associative control of action. However, a possible advantage of this route is that this form of learning might generalise better to novel situations. We are currently testing this idea in our lab. Note that in the associative-learning literature, there is an ongoing debate as to whether learning associations between a stimu-

lus and an action is rule-based or based on the formation of specific stimulus-response associations (see e.g. McLaren et al., 2013a; Mitchell, De Houwer, & Lovibond, 2009). Even though this is speculative, one could hypothesise that in our framework similar learning mechanisms underlie rule-based behaviour and stimulus-response link-based behaviour. The main difference between the two is the kind of representation that is linked with the stimulus: an abstract, rule-like representation (X—if x then left'), or more concrete stimulus-response associations (X—left).

Finally, stimulus-specific learning may also have a more indirect impact on action control via a link with the outcome-evaluation processes discussed above. Veling, Holland, and van Knippenberg (2008) have shown that consistently pairing stimuli with the act of withholding a response results in devaluation of stimuli that were initially rated as positive. Similar devaluation effects have been observed in other studies (for a short overview, see Ferrey, Frischen, & Fenske, 2012; Kiss, Raymond, Westoby, Nobre, & Eimer, 2008). Furthermore, no-go training cannot only reduce the subjective hedonic value of erotic images, it may also reduce the motivational incentive of such stimuli. Ferrey et al. (2012) found that participants were less willing to invest time and effort (measured by the number of key presses participants were willing to execute) to view images similar to the ones paired with no-go cues. The link between associative learning and value is also supported by the finding that learning stimulus-go associations happens faster in a reward condition than in a punishment condition; by contrast, stimulus/no-go associations are learned faster in the punishment condition (Cavanagh, Eisenberg, Guitart-Masip, Huys, & Frank, 2013; Guitart-Masip et al., 2012). Thus, there may be a 'hard-wired' Pavlovian bias that couples reward with approach ('go') and punishment with avoidance ('nogo') (Cavanagh et al., 2013; Guitart-Masip et al., 2012). Note that the value of items could also be modulated associatively via associations between stimuli. Wimmer and Shohamy (2012) demonstrated that the delivery of reward for a specific item can spread to associated items stored in long-term memory. In other words, the value of unrewarded items was modulated via associations with rewarded items. This phenomenon could explain how people can quickly decide between items that they have never seen before.

In combination, the work above suggests how changing behaviour can become automatised. However, work on extinction learning indicates that some associatively mediated forms of action control may be context-dependent (in contrast to rule-based action control which seems context-independent; see above). In

Pavlovian learning, extinction occurs when a stimulus that was originally paired with an event is repeatedly presented alone; in instrumental learning, extinction occurs when an action that was originally paired with a reward, is no longer reinforced. In both cases, learned behaviour typically declines, but the originally learned behaviour often returns when the context changes (Bouton & Woods, 2008). This suggests that extinction learning is context-dependent. Thus, even though automatised action control, it does come with certain limitations.

Interim key points:

- Action control can become a 'learned' reflex: replacing dominant actions initially requires top-down bias but could gradually become automatised, with the need for top-down bias disappearing altogether.
- Associative learning can influence action control by modulating each processing step in our framework (i.e. signal detection, action selection, action execution).
- We hypothesise that similar learning mechanisms underlie both rule-based and stimulus-response link-based behaviour.

The Development Of An Action Control Network

We argued above that a key characteristic of flexible human behaviour is the ability to implement new rules quickly, and we proposed that this feat can be achieved by reusing or recombining existing representations or rules (i.e. the 'compositionality' idea). But in our quest to abolish the control homunculus from theories of action control, we need to address one final issue: how does the 'control repertoire', or the set of basic rules and control processes, initially develop?

Learning of rules and abstract representations.

Little research has been done on how rules for complex actions are initially acquired (Collins & Frank, 2013). Basic reinforcement learning accounts can explain how people acquire simple stimulus-action rules. A central assumption of these accounts is that simple rules are learned via exploration of the environment: when a stimulus is presented, one can try different courses of action (e.g. pressing a button on a new piece of equipment) and subsequently monitor the outcome of the chosen actions. Each time a particular action in response to the presentation of a stimulus leads to a positive outcome, the strength of the action increases, and eventually, simple rule-like structures develop. As argued above, stimuli and responses can also become 'paired' via error-driven and Hebbian associative learning mechanisms. Error-correction learning

mechanisms try to reduce the discrepancy between the predicted outcome and the actual outcome (McLaren et al., 2013b), whereas Hebb's learning rule states that 'cells that fire together bind together'. However, basic reinforcement and associative learning accounts struggle to explain more complex goal-directed behaviour in environments in which multiple stimuli or stimulus features (e.g. colour or shape) can be attended to and in which many actions can be selected.

One of the harder questions in psychology is how, starting with a set of basic associative- or reinforcement-learning processes, it might be possible to deploy them so as to arrive at a system capable of propositional reasoning. In other words, how can we go from associations to rules? There have been some successful attempts to integrate basic learning and rule acquisition. For example, Rougier and colleagues (2005) developed a neurologically inspired computational model of 'rule' learning. The model was trained to respond to multi-dimensional stimuli. In each block, only one dimension was relevant (e.g. colour). Across trials, the specific features within a dimension changed (e.g. red, green, yellow), but activity in the prefrontal cortex was maintained due to a gating mechanism (Hazy, Frank, & O'reilly, 2007). As a result, the PFC system developed patterns of activity that encoded abstract representations of the relevant stimulus-dimension (e.g. 'colour'). These abstract rule-like representations subsequently guided behaviour by providing 'top-down' excitatory support for the relevant stimulus dimension in the subordinate processing levels (cf. biased competition). The biasing was possible because links between the abstract representations and the processing layers were built during training. Thus, the model produced flexible rule-like behaviour without 'biologically problematic symbolic processing computations' (Rougier et al., 2005, p. 7343). After sufficient training, the model was also able to respond correctly to stimuli it had not seen before. This generalisation correlated strongly with development of abstract representations. By contrast, models without the PFC system, such as the associative simple recurrent network (SRN) model (Elman, 1990), had to learn specific stimulus-response combinations, and these did not generalise well to novel situations.

The latter result is consistent with findings of Spiegel and McLaren (2006). Humans and the recurrent network model were trained on a task in which they had to respond to the location of circles on a computer screen. In the experimental group, trial sequences always had a specific structure (e.g. ABB(varying numbers of Cs)BBA; the letters represent three possible locations of the circles). Over a series of experiments, it was

demonstrated that the network model used all the structure available to develop simple rule-like representations. This resulted in faster and more accurate responses. These representations allowed some generalisation to novel situations (hence they were rule-like). However, generalisation was imperfect because the model was sensitive to seemingly inconsequential departures from the initial structure. This was very similar to the behaviour of humans in these experiments when they were unaware of the contingencies in play. However, under some conditions, humans were able to induce the rule as programmed by the experimenters (e.g. 'always as many Bs before as after the Cs'), and in these instances generalisation was near perfect. The recurrent network model was never able to do this. This suggests that rule learning in humans comes about as the result of a more complex system with many interacting parts. Nevertheless, the finding that a simple associative network is able to develop basic rules (albeit imperfectly) further supports the idea that basic associative or reinforcement mechanisms may indeed underlie rule learning. Consistent with this, Ramamoorthy and Verguts (2012) recently developed a computational model of instruction following which relied on basic Hebbian learning processes (see above) in prefrontal cortex (see also e.g. Deco & Rolls, 2005).

If we assume that we begin with reward- or error-driven associative learning processes, then we believe that in order to be capable of developing rule-like representations, these processes will need to be embedded in an architecture that must, at a minimum, be complex (by which we mean multi-layer or more than one layer of weights), non-linear (so that the multiple layers are not simply equivalent to a single layer; see Minsky & Parert, 1969) and recursive (so that the system can, in principle, compute any computable function). Obviously the learning algorithm used will have to be capable of operating within this framework (for an example of such an algorithm, see Rumelhart, Hinton, & Williams, 1986). In essence, then, we propose that associative processes within a sufficiently rich and complex architecture can deliver the possibility of rule-based (symbolic) computation. But even if all these requirements are met there is still much to be done. It will be the interaction of that system with the world that will allow this development to take place. The architecture and learning algorithms, which have evolved throughout human evolution, merely confer the potential for rule-based processing; the potential has to be realised in the course of experience, and so the transition from association to cognition is also a developmental issue.

In sum, we believe that a key to behavioural flexibility is the development of abstract repre-

sentations via basic learning mechanisms. These representations will guide or 'contextualise' stimulus detection, action selection, and action execution (Badre, Kayser, & D'Esposito, 2010; Collins & Frank, 2013; see also e.g. Frank & Badre, 2012), and allow generalisation to novel situations (see also Forrest et al., 2012). Even though these models were used to simulate relatively straightforward rule-based behaviour, the general principles are likely to apply to more complex situations (Rougier et al., 2005). In complex environments, the agent may make 'temporal abstractions': grouping together a set of interrelated actions ('options' or 'policies') (Botvinick, 2012; Botvinick et al., 2009). These policies can be learned and selected via the same reinforcement-based learning mechanisms discussed above. When a policy is selected (e.g. 'making coffee'), the more 'primitive' motor actions are produced based on the acquired lower-level stimulus-action associations (see e.g. Botvinick, 2012, for an accessible discussion).

Development: Building a network for the future.

Major changes in rule-based action control take place during childhood. Indeed, the ability of children to regulate their behaviour improves remarkably from infancy through adolescence (for recent reviews, see e.g. Bunge & Crone, 2009; Bunge, Mackey, & Whitaker, 2009; Diamond, 2013). Such changes have been linked to development of executive control functions and the protracted development of the prefrontal control network (Bunge & O'Hare, 2012).

Developmental changes in rule-based action control throughout early and late development can be linked to a shift from concrete stimulus-action associations to abstract rule-like representations (Bunge & Zelazo, 2006; Munakata et al., 2012). Initially, young children would learn simple stimulus-action associations via exploration ('if I push this button, a light turns on'), automatic imitation, or deliberate demonstration by others. These associations then become the 'building blocks' for the rule-based control network, and shape the development of more abstract representations that constrain and regulate other ongoing processes. Indeed, Rougier et al. (2005) found that concrete stimulus-response representations (in posterior brain systems) had to stabilise before abstract rule-like representations could be extracted.

Several studies support the transition account. For example, young children are influenced more by specific stimulus-response associations than adults when switching between tasks (Crone, Bunge, van der Molen, & Ridderinkhof, 2006). Furthermore, Kharitonova and Munakata (2011) have demonstrated that in 3-year-old children, flexible rule use in a card-sort-

ing test correlated with performance in an abstraction test that required children to select the odd-one-out based on an overarching category. They suggested that this link could be explained by a common underlying working memory mechanism that supports rule-like abstraction and perceptually-based abstraction (see also Collins & Frank, 2013). In other words, abstraction underlies flexibility.

In addition to changes in the ability to develop abstract rule-like representations, children may also develop an ability to generate 'temporal abstractions'. Botvinick et al. (2009) noted that throughout development, action control becomes more hierarchical, with simple actions or rules becoming integrated within larger wholes or structures. Similarly, Bunge and Zelazo (2006) reviewed a series of studies suggesting that development of cognitive control was associated with an increased ability to represent hierarchies of rules in which higher-order rules (cf. 'policies') operate on lower-order rules.

To conclude, Thompson-Schill et al. (2009) proposed that protracted development of the executive prefrontal network is necessary to allow children to learn linguistic and social conventions. Here, we propose that learning necessarily precedes executive control because learning has to provide the building blocks for a 'control repertoire' based on abstraction first.

Interim key points:

- Our framework places learning of increasingly abstract representations at the heart of executive control.
- Only through constant interaction with their environment can people build up a control repertoire that will underlie all forms of rule-based behaviour.
- This repertoire continues to develop throughout the life-span.

Implications For Behavioural Change

Clinical disorders often have many origins; alterations of cognitive processes may be one of them. Therefore, we believe that our framework can be applied in clinical domains to increase our understanding of certain control deficits and provide a theoretical basis for the development of novel 'behavioural change' interventions.

Just as in the cognitive and neuroscience domain, attribution of control to convenient control homunculi is still very present in the clinical and more applied domains. Most clinical and neurobiological models that rely on executive control lack a precise description of the underlying cognitive components and mechanisms. We have argued that a failure to change behaviour could have multiple origins. Thus, merely describing a deficit or phenomenon as a deficit of 'inhibition' or 'executive control' does not provide

an explanation, and discourages discussion of alternative theories. For instance, poor stopping performance in adults with ADHD may be partly due to inattention (Bekker et al., 2005). Many studies have observed stopping deficits in children and adults with ADHD, which has led researchers to suggest that poor response inhibition is central to their deficit (Lijffijt, Kenemans, Verbaten, & van Engeland, 2005; Lipszyc & Schachar, 2010; Nigg, 2001). However, Bekker et al. found using event-related potentials (ERPs) that an early attention-related component (the N1, which is a negative-going ERP component observed 80-120 ms after the presentation of an auditory stop signal) was larger for successful stop trials than for unsuccessful stop trials in the control group. This finding suggests that perceptual attention contributes to stopping. This difference in N1 was absent in adults with ADHD, which suggests that stopping deficits in adults with ADHD are not entirely due to deficiencies in inhibition but also to deficiencies in stimulus detection. Similarly, Brown and Braver (2008) have argued that the failure to suppress risky and inappropriate behaviour in addictions could stem from a failure to adjust performance after learning (for a similar idea, see e.g. Garavan & Stout, 2005). These studies indicate that focusing on basic processes provides a more detailed account of control deficits in e.g. behavioural and substance addictions. This may lead not only to important new insights in the aetiology of various disorders characterised as deficits in changing behaviour, but also to the development of strategies for treating these conditions. Indeed, a common critique is that the effective mechanisms of most behavioural treatments are still underspecified (Toneatto & Ladoceur, 2003). Therefore, providing a detailed account of action control deficits could stimulate the development of new theory-driven behavioural treatments. For example, it could lead to personalised interventions: Person A and Person B may both have 'stopping' deficits with different origins, so the intervention for Person A could e.g. focus on biasing attention (e.g. in adults with ADHD; see e.g. Bekker et al., 2005), whereas the intervention for Person B could e.g. focus on performance monitoring and control adjustments (in e.g. substance abusers; see e.g. Garavan & Stout, 2005).

The work on proactive control suggests that preparation could lead to a prepared reflex, making action control less susceptible to the negative effects of concurrent load (Logan, 1978). This is consistent with findings in the 'implementation intention' literature (Gollwitzer, 1999; Gollwitzer & Sheeran, 2006; Gollwitzer, Gawrilow, & Oettingen, 2010). Implementation intentions refer to the linking of critical situations or cues to specific actions (e.g. 'Whenever I see a red light on a

food item, I will not buy it'). This could lead to a prepared reflex; indeed, Gollwitzer noted that after implementation intentions are formed, 'action initiation becomes swift, efficient, and does not require conscious intent' (Gollwitzer, 1999, p. 495). Others have argued that forming implementation intentions leads to increased monitoring for cues (see Rummel, Einstein, & Rampey, 2012, for a discussion), but this is still consistent with the proactive control idea discussed above. Importantly, implementation intentions, and consequently, proactive control, could have a positive impact on behaviour (Gollwitzer & Sheeran, 2006; Gollwitzer et al., 2010). For example, they reduce the negative impact of stress on rule-based action control (Scholz et al., 2009), presumably because less reactive control is required. They may also strengthen the effects of commercial weight loss programs (Luszczynska, Sobczyk, & Abraham, 2007) and reduce binge-drinking (Hagger et al., 2012). Thus, an avenue for future research is how proactive control can be used in treatments, bearing in mind that there are certain costs associated with it (as discussed above).

It is also important to understand how people develop and use new rules. For example, supermarkets in the UK recently started using a traffic-light labelling system to indicate sugar, fat, salt, and calorie-contents of food items. But how do people use this new system to replace their favourite (but unhealthy) food item with a more healthy option? The work on rule learning, and in particular, generalisation and abstraction, discussed above could provide some clues. For example, it suggests that new rules that are based on previously acquired rules (e.g. red light = stop) might be learned more quickly (and consequently, be more effective). The ability to form abstract rules may also lead to generalisation of control across domains. Recent work from our lab suggests that preparing to stop a motor response can result in more cautious decision-making in a gambling task (Verbruggen, Adams, & Chambers, 2012). We are currently exploring the mechanisms behind this transfer, but it is possible that this effect is caused by proactive control: Having the abstract stop rule activated could have biased monetary decision-making. More generally, we agree with Munakata et al. (2012), among others, that abstraction may explain executive-control training effects in children. Executive control training may work better in children with low self-control than in adults with low self-control (Berkman, Graham, & Fisher, 2012). A better understanding of how rules are developed would lead to more effective training. Even though this is highly speculative, building abstract rule-like representations may also provide an explanation for some more idiosyncratic transfer effects, such as the positive

effect of avoiding sweets or regularly squeezing a handgrip in a two-week training period on stop performance afterwards (Muraven, 2010), or the differential effect of open vs. skilled sports on stopping (Wang et al., 2013), or other inhibitory spill-over effects (e.g. Berkman, Burklund, & Lieberman, 2009). As proposed by the compositional account, people may recycle or recombine rules that they used in other situations; building up a control repertoire in one domain could therefore lead to improved performance in other apparently non-related domains as long as the acquired representations are abstract enough.

Finally, associatively-mediated action control could open the avenue for new treatments. Several recent studies have already shown that learning to stop motor responses towards food- or alcohol-related stimuli influences food and alcohol intake both inside and outside the lab. For example, several studies have found that consistent pairing of food-related pictures to stopping in a go/no-go or stop-signal-paradigm reduced subsequent food consumption (Houben, 2011; Houben & Jansen, 2011; Lawrence, Verbruggen, Adams, & Chambers, 2013; Veling, Aarts, & Papies, 2011; Veling, Aarts, & Stroebe, 2012). Furthermore, a similar procedure with alcohol-related stimuli reduced the hedonic value of alcohol and the subsequent weekly alcohol intake of heavy drinking students (Houben, Havermans, Nederkoorn, & Jansen, 2012). Wiers et al. found that a similar avoidance training task influenced treatment outcomes in alcoholics a year later (Wiers, Eberl, Rinck, Becker, & Lindenmeyer, 2011). Finally, Jones and Field found that stimulus-specific stop training reduced alcohol-intake in the laboratory but not self-reported drinking in the week after training (Jones & Field, 2013). In combination, these results suggest that go/no-go, avoidance or stop-signal training can influence approach behaviour towards a range of stimuli, possibly by changing attitudes towards these stimuli or by creating non-respond (avoid) associations. People may also associatively learn to direct their attention either towards or away from stimuli. Recent meta-analyses suggest that attentional-bias modification could be used in treatments for anxiety (Hakamata et al., 2010), although the effect may be more modest than initially suggested (Hallion & Ruscio, 2011). Several studies have also examined attentional-bias modification in addiction. This could involve training people to redirect attention away from drug-related cues towards more neutral cues (Wiers, Gladwin, Hofmann, Salemink, & Ridderinkhof, 2013). The effectiveness of this training on addiction is still unclear. For example, a single session of attentional bias modification did not influence subjective craving or behavioural measures of tobacco seeking in cigarette smokers (Field, Duka, Tyler,

& Schoenmakers, 2009). Approach/avoidance or stop/no-go training may be more effective because several aspects of inappropriate behaviour can be influenced at the same time. Indeed, avoidance or stop/no-go training could influence hedonistic value, motivational behaviour (approach vs. avoidance), and possibly even attention towards the stimuli. Preliminary data from studies inspired by the framework presented here suggest that subjects may learn associations between the go stimulus and the stop signal, enhancing detection of the latter. However, much more research is needed to examine how well various forms inhibitory and executive-control training can influence behaviour outside the lab (see also e.g. Jones, Christiansen, Nederkoorn, Houben, & Field, 2013). This work will also have to address the context-dependence issue (Bouton & Woods, 2008).

Final Thoughts And Conclusions

We have discussed how action control can be attributed to the interplay between three basic cognitive processes: signal detection, action selection, and action execution. These processes are constantly adjusted and biased via abstract representations that develop slowly but that can be generalised to different contexts. These representations support flexible behaviour. At the same time, more concrete stimulus-action outcome associations are learned, which can result in automatization of actions that were initially regulated by the 'executive' abstraction-based system. We have attempted to unpack each component of our framework. It is possible that future research will demonstrate that some components or processes may have to be broken down further, leading to an even more nested system. We agree with Dennett that the only way to 'discharge fancy homunculi from one's scheme [is] by organizing armies of such idiots to do the work' (quoted in Monsell & Driver, 2000, p.7). We believe that in order to understand how control is achieved, boxes have to be broken down until we understand how complex behaviour arises from a basic set of cognitive processes that can be implemented by our neural system. One may object against this 'deconstruction idea' on the basis of parsimony: a model with fewer components may seem more parsimonious. However, this parsimony would only be achieved by attributing multiple powers to specific controllers or control functions (Monsell & Driver, 2000), making the seemingly more parsimonious account equally complex.

Relation with other frameworks and models

As indicated in the text, our framework builds upon existing accounts of attention, control, working memory, and learning (e.g. Chein &

Schneider, 2012; Deco & Rolls, 2005; Desimone & Duncan, 1995; Logan & Gordon, 2001; Miller & Cohen, 2001; Rougier et al., 2005). Consistent with these accounts, we postulate that the main role of the executive control system is to bias competition in subordinate processes via rules maintained in working memory (either in an active or silent mode). But we also propose that once the rules are implemented, the control system can take a back seat in many, if not most, situations, and action control may eventually become automatized. This overall framework is consistent with the 'Learning and Control' framework of Chein and Schneider (2012), who proposed that there are three systems: a *meta-cognitive system*, which supports rule learning, monitoring, and task sequencing (cf. hierarchical control); a *cognitive control system*, which supports attention and action control; and a *representation system*, which supports associative learning. They also suggested that through practice, the associative system will take over from the metacognitive and control systems, which is consistent with our 'learned reflex' idea.

We believe that the main strength of our framework is that it integrates various theories and models, links findings, and points out similarities and differences between domains. This integration is a crucial step to enhance our understanding of executive control and behavioural change.

Beyond action control

Many principles of our action control framework may translate to control in other domains, including emotion and control of thought. Several lines of evidence suggest a certain overlap between control of action and control of thought and emotion. Action control and control of emotion and thought may also be coupled because changes in internal states (such as thoughts or an emotional reaction) could trigger changes in ongoing actions. Corbetta et al.'s review suggests that the ventral attention network, which is required for action control, might be involved in switching between aspects of 'internal' processing, such as memory retrieval or self-referential thoughts (Corbetta et al., 2008). Furthermore, brain areas that are important for action control, such as the right inferior frontal gyrus and the right middle frontal gyrus, may also regulate emotional memories (Depue, Curran, & Banich, 2007) or unwanted thoughts (Benoit & Anderson, 2012; Depue, Burgess, Willcutt, Ruzic, & Banich, 2010). The overlap between action control and emotion regulation is further supported by correlations between rumination, inhibition and task switching (Whitmer & Banich, 2007). These and other similar findings have led several researchers to propose that similar control mechanisms may be required to regulate various as-

pects of human behaviour (Aron, Robbins, & Poldrack, 2004; Banich et al., 2009; Cohen & Lieberman, 2010). Although speculative, this overlap could again be partly due to involvement of the abstract rule-like representations in various domains ('do not think', 'do not respond', ...). More generally, we believe that the main difference between domains may be in the content controlled, not in the mechanisms by which control is achieved (see also Logan et al., 2013).

The work of Depue, Banich, and others suggest that emotion and executive control may influence each other. This link is further discussed by Pessoa (2009), who proposed a dual-competition framework to account for the effects of low- and high-threat emotional information on executive control. This framework can be integrated with our action control framework. More specifically, based on Pessoa's framework we hypothesise that emotional content that is low in threat primarily interferes with attentional selection; by contrast, high-threat information would interfere with action selection as well. We believe that this highlights one of the major benefits of our processing framework: by focusing on the specific mechanisms rather than the general functions, a richer and more detailed picture emerges.

In a similar vein, one could use our framework to examine how motivation, mood, stress and other state-dependent factors influence action control and flexible behaviour. For example, animal studies have shown that the prefrontal cortex is modulated by neurotransmitter systems mediating stress and arousal (Arnsten, 2009; Robbins, 2007). The link between stress and action control is also demonstrated by the finding that people with addictions are prone to fail to suppress drug-seeking behaviour in stressful situations (Sinha, 2008). We hypothesise that acute stress could influence action control in at least three different ways: it could lead to a narrowed focus of attention (Chajut & Algom, 2003), it could discourage selection of alternative actions (Goschke, 2000), or it could shift the balance between rule-based and associatively-mediated action control (Schwabe, Dickinson, & Wolf, 2011). Given the impact of state-dependent factors on behavioural change, a better understanding of which processes are influenced by factors such as stress can again lead to the development of new interventions and improvements in existing ones.

Conclusion

To conclude, we hope that this article will inspire research on action control, behavioural flexibility, and behavioural change to focus more on specific cognitive processes and representations, how these can be learned throughout development and adulthood, and how these are

maintained. We believe that this will lead to better models of executive control of action and behavioural change, which can inspire the development of new and more effective theory-driven interventions. Ultimately, this approach will allow us to banish homunculi from our theories.

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Footnotes

1. The accumulation process is analogous to drawing a series of balls from an urn (Usher & McClelland, 2001). For example, the urn can contain balls with 'left' or 'right' written on them. At each time step in the response-selection process, a ball would be drawn, and a count is added to the appropriate counter (e.g. when a 'left' ball is selected, a count is added to the left counter). A response is selected when one of the counters reaches a threshold. When the response threshold is 10, then a response is selected when 10 'left' balls or 10 'right' balls have been drawn from the urn.

2. A quasi-experiment was subsequently conducted. The 'prepared reflex' idea was tested by either signalling or not signalling at this junction when a car was waiting to turn on another 20 days, 10 signal, 10 no signal in alternation (this took about half a year). The results were that on 50% of 'signal' occasions the car immediately turned. This never happened on 'no-signal' occasions.

References

- Anderson, M. L. (2010). Neural reuse: A fundamental organizational principle of the brain. *The Behavioral and Brain Sciences*, 33, 245-66; discussion 266-313. doi:10.1017/S0140525X10000853
- Arnsten, A. F. (2009). Stress signalling pathways that impair prefrontal cortex structure and function. *Nature Reviews. Neuroscience*, 10, 410-22. doi: 10.1038/nrn2648

- Aron, A. R., Robbins, T. W., & Poldrack, R. A. (2004). Inhibition and the right inferior frontal cortex. *Trends in Cognitive Sciences*, *8*, 170-7. doi:10.1016/j.tics.2004.02.010
- Baddeley, A. (1996). Exploring the central executive. *Quarterly Journal of Experimental Psychology* (2006), *49A*, 5-28.
- Badre, D., Kayser, A. S., & D'Esposito, M. (2010). Frontal cortex and the discovery of abstract action rules. *Neuron*, *66*, 315-26. doi:10.1016/j.neuron.2010.03.025
- Banich, M. T., Mackiewicz, K. L., Depue, B. E., Whitmer, A. J., Miller, G. A., & Heller, W. (2009). Cognitive control mechanisms, emotion and memory: A neural perspective with implications for psychopathology. *Neuroscience and Biobehavioral Reviews*, *33*, 613-30. doi:10.1016/j.neubiorev.2008.09.010
- Bechara, A., Noel, X., & Crone, E. A. (2006). Loss of willpower: Abnormal neural mechanisms of impulse control and decision making in addiction. *Handbook of Implicit Cognition and Addiction*, 215-232.
- Bekker, E. M., Overtoom, C. C., Kooij, J. J., Buitelaar, J. K., Verbaten, M. N., & Kenemans, J. L. (2005). Disentangling deficits in adults with attention-deficit/hyperactivity disorder. *Archives of General Psychiatry*, *62*, 1129-36. doi:10.1001/archpsyc.62.10.1129
- Benoit, R. G., & Anderson, M. C. (2012). Opposing mechanisms support the voluntary forgetting of unwanted memories. *Neuron*, *76*, 450-60. doi:10.1016/j.neuron.2012.07.025
- Berkman, E. T., Burkland, L., & Lieberman, M. D. (2009). Inhibitory spillover: Intentional motor inhibition produces incidental limbic inhibition via right inferior frontal cortex. *NeuroImage*, *47*, 705-12. doi:10.1016/j.neuroimage.2009.04.084
- Berkman, E. T., Graham, A. M., & Fisher, P. A. (2012). Training self-control: A domain-general translational neuroscience approach. *Child Development Perspectives*, n/a. doi:10.1111/j.1750-8606.2012.00248.x
- Botvinick, M. M. (2007). Conflict monitoring and decision making: Reconciling two perspectives on anterior cingulate function. *Cognitive, Affective & Behavioral Neuroscience*, *7*, 356-66.
- Botvinick, M. M. (2012). Hierarchical reinforcement learning and decision making. *Current Opinion in Neurobiology*, *22*, 956-62. doi:10.1016/j.conb.2012.05.008
- Botvinick, M. M., Braver, T. S., Barch, D. M., Carter, C. S., & Cohen, J. D. (2001). Conflict monitoring and cognitive control. *Psychological Review*, *108*, 624-652.
- Botvinick, M. M., Niv, Y., & Barto, A. C. (2009). Hierarchically organized behavior and its neural foundations: A reinforcement learning perspective. *Cognition*, *113*, 262-80. doi:10.1016/j.cognition.2008.08.011
- Boucher, L., Palmeri, T. J., Logan, G. D., & Schall, J. D. (2007). Inhibitory control in mind and brain: An interactive race model of countermanding saccades. *Psychological Review*, *114*, 376-97. doi:10.1037/0033-295X.114.2.376
- Bouton, M. E., & Woods, A. M. (2008). 1.09 extinction: Behavioral mechanisms and their implications. *Learning and Memory: A Comprehensive Reference*, *1*, 151.
- Braver, T. S. (2012). The variable nature of cognitive control: A dual mechanisms framework. *Trends in Cognitive Sciences*, *16*, 106-13. doi:10.1016/j.tics.2011.12.010
- Braver, T. S., Gray, J. R., & Burgess, G. C. (2007). Explaining the many varieties of working memory variation: Dual mechanisms of cognitive control. *Variation in Working Memory*, 76-106.
- Braver, T. S., Paxton, J. L., Locke, H. S., & Barch, D. M. (2009). Flexible neural mechanisms of cognitive control within human prefrontal cortex. *Proceedings of the National Academy of Sciences of the United States of America*, *106*, 7351-6. doi:10.1073/pnas.0808187106
- Brown, J. W., & Braver, T. S. (2005). Learned predictions of error likelihood in the anterior cingulate cortex. *Science*, *307*, 1118-21. doi:10.1126/science.1105783
- Brown, J. W., & Braver, T. S. (2008). A computational model of risk, conflict, and individual difference effects in the anterior cingulate cortex. *Brain Research*, *1202*, 99-108. doi:10.1016/j.brainres.2007.06.080
- Brown, S. D., & Heathcote, A. (2008). The simplest complete model of choice response time: Linear ballistic accumulation. *Cognitive Psychology*, *57*, 153-78. doi:10.1016/j.cogpsych.2007.12.002
- Buch, E. R., Mars, R. B., Boorman, E. D., & Rushworth, M. F. (2010). A network centered on ventral premotor cortex exerts both facilitatory and inhibitory control over primary motor cortex during action reprogramming. *The Journal of Neuroscience*, *30*, 1395-401. doi:10.1523/JNEUROSCI.4882-09.2010
- Bugg, J. M., & Crump, M. J. (2012). In support of a distinction between voluntary and stimulus-driven control: A review of the literature on proportion congruent effects. *Frontiers in Psychology*, *3*, 367. doi:10.3389/fpsyg.2012.00367
- Bunge, S. A. (2004). How we use rules to select actions: A review of evidence from cognitive neuroscience. *Cognitive, Affective & Behavioral Neuroscience*, *4*, 564-79.
- Bunge, S. A., & Crone, E. A. (2009). Neural correlates of the development of cognitive control. *Neuroimaging in Developmental Clinical Neuroscience*, 22-37.
- Bunge, S. A., & O'Hare, E. D. (2012). *The developing human brain*. A Frontiers Research Topic. 2012.
- Bunge, S. A., & Zelazo, P. D. (2006). A brain-based account of the development of rule use in childhood. *Current Directions in Psychological Science*, *15*, 118-121.
- Bunge, S. A., Mackey, A. P., & Whitaker, K. J. (2009). Brain changes underlying the development of cognitive control and reasoning. In Gazzinga (Ed.), *The cognitive neurosciences, 4th edition*. Cambridge: MA: MIT Press.
- Cai, W., Oldenkamp, C. L., & Aron, A. R. (2011). A proactive mechanism for selective suppression of response tendencies. *The Journal of Neuroscience*, *31*, 5965-9. doi:10.1523/JNEUROSCI.6292-10.2011
- Cavanagh, J. F., Eisenberg, I., Guitart-Masip, M., Huys, Q., & Frank, M. J. (2013). Frontal theta overrides pavlovian learning biases. *The Journal of Neuroscience*, *33*, 8541-8548. doi:10.1523/JNEUROSCI.5754-12.2013
- Chajut, E., & Algom, D. (2003). Selective attention improves under stress: Implications for theories of

social cognition. *Journal of Personality and Social Psychology*, *85*, 231. doi:10.1037/0022-3514.85.2.231

Chamberlain, S. R., & Sahakian, B. J. (2007). The neuropsychiatry of impulsivity. *Current Opinion in Psychiatry*, *20*, 255-261.

Chein, J. M., & Schneider, W. (2012). The brain's learning and control architecture. *Current Directions in Psychological Science*, *21*, 78-84. doi:10.1177/0963721411434977

Chein, J. M., & Schneider, W. (2005). Neuroimaging studies of practice-related change: FMRI and meta-analytic evidence of a domain-general control network for learning. *Brain Research. Cognitive Brain Research*, *25*, 607-23. doi:10.1016/j.cogbrainres.2005.08.013

Chelazzi, L., Duncan, J., Miller, E. K., & Desimone, R. (1998). Responses of neurons in inferior temporal cortex during memory-guided visual search. *Journal of Neurophysiology*, *80*, 2918-40.

Chun, M. M., Golomb, J. D., & Turk-Browne, N. B. (2011). A taxonomy of external and internal attention. *Annual Review of Psychology*, *62*, 73-101. doi:10.1146/annurev.psych.093008.100427

Claffey, M. P., Sheldon, S., Stinear, C. M., Verbruggen, F., & Aron, A. R. (2010). Having a goal to stop action is associated with advance control of specific motor representations. *Neuropsychologia*, *48*, 541-8. doi:10.1016/j.neuropsychologia.2009.10.015

Cohen, J. D., Dunbar, K., & McClelland, J. L. (1990). On the control of automatic processes: A parallel distributed processing account of the stroop effect. *Psychological Review*, *97*, 332-61.

Cohen, J. D., Servan-Schreiber, D., & McClelland, J. L. (1992). A parallel distributed processing approach to automaticity. *The American Journal of Psychology*, *105*, 239-69.

Cohen, J. R., & Lieberman, M. D. (2010). The common neural basis of exerting self-control in multiple domains. In *Self control in society, mind, and brain* (Vol. 1, pp. 141-162). Oxford Scholarship Online Monographs.

Cohen-Kdoshay, O., & Meiran, N. (2009). The representation of instructions operates like a prepared reflex: Flanker compatibility effects found in first trial following S-R instructions. *Experimental Psychology*, *56*, 128-33. doi:10.1027/1618-3169.56.2.128

Cole, M. W., Laurent, P., & Stocco, A. (2013). Rapid instructed task learning: A new window into the human brain's unique capacity for flexible cognitive control. *Cognitive, Affective & Behavioral Neuroscience*, *13*, 1-22. doi:10.3758/s13415-012-0125-7

Collins, A. G., & Frank, M. J. (2013). Cognitive control over learning: Creating, clustering, and generalizing task-set structure. *Psychological Review*, *120*, 190-229. doi:10.1037/a0030852

Corbetta, M., & Shulman, G. L. (2002). Control of goal-directed and stimulus-driven attention in the brain. *Nature Reviews. Neuroscience*, *3*, 201-15. doi:10.1038/nrn755

Corbetta, M., Patel, G., & Shulman, G. L. (2008). The reorienting system of the human brain: From environment to theory of mind. *Neuron*, *58*, 306-24. doi:10.1016/j.neuron.2008.04.017

Crews, F. T., & Boettiger, C. A. (2009). Impulsivity, frontal lobes and risk for addiction. *Pharmacology, Biochemistry, and Behavior*, *93*, 237-47. doi:10.1016/j.pbb.2009.04.018

Criaud, M., Wardak, C., Ben Hamed, S., Ballanger, B., & Boulinguez, P. (2012). Proactive inhibitory control of response as the default state of executive control. *Frontiers in Psychology*, *3*, 59. doi:10.3389/fpsyg.2012.00059

Crone, E. A., Bunge, S. A., van der Molen, M. W., & Ridderinkhof, K. R. (2006). Switching between tasks and responses: A developmental study. *Developmental Science*, *9*, 278-87. doi:10.1111/j.1467-7687.2006.00490.x

Curtis, C. E., & D'Esposito, M. (2003). Persistent activity in the prefrontal cortex during working memory. *Trends in Cognitive Sciences*, *7*, 415-423. doi:10.1016/S1364-6613(03)00197-9

Curtis, C. E., & Lee, D. (2010). Beyond working memory: The role of persistent activity in decision making. *Trends in Cognitive Sciences*, *14*, 216-22. doi:10.1016/j.tics.2010.03.006

Dalley, J. W., Everitt, B. J., & Robbins, T. W. (2011). Impulsivity, compulsivity, and top-down cognitive control. *Neuron*, *69*, 680-94. doi:10.1016/j.neuron.2011.01.020

Deco, G., & Rolls, E. T. (2005). Attention, short-term memory, and action selection: A unifying theory. *Progress in Neurobiology*, *76*, 236-56. doi:10.1016/j.pneurobio.2005.08.004

Depue, B. E., Burgess, G. C., Willcutt, E. G., Ruzic, L., & Banich, M. T. (2010). Inhibitory control of memory retrieval and motor processing associated with the right lateral prefrontal cortex: Evidence from deficits in individuals with ADHD. *Neuropsychologia*. doi:10.1016/j.neuropsychologia.2010.09.013

Depue, B. E., Curran, T., & Banich, M. T. (2007). Prefrontal regions orchestrate suppression of emotional memories via a two-phase process. *Science*, *317*, 215-9. doi:10.1126/science.1139560

Desimone, R., & Duncan, J. (1995). Neural mechanisms of selective visual attention. *Annual Review of Neuroscience*, *18*, 193-222.

Diamond, A. (2013). Executive functions. *Annual Review of Psychology*, *64*, 135-68. doi:10.1146/annurev-psych-113011-143750

Dickinson, A. (1985). Actions and habits: The development of behavioural autonomy. *Philosophical Transactions of the Royal Society of London. B, Biological Sciences*, *308*, 67-78.

Dodds, C. M., Morein-Zamir, S., & Robbins, T. W. (2011). Dissociating inhibition, attention, and response control in the frontoparietal network using functional magnetic resonance imaging. *Cerebral Cortex*, *21*, 1155-65. doi:10.1093/cercor/bhq187

Duncan, J. (2006). EPS mid-career award 2004: Brain mechanisms of attention. *Quarterly Journal of Experimental Psychology* (2006), *59*, 2-27. doi:10.1080/17470210500260674

Duncan, J., & Owen, A. M. (2000). Common regions of the human frontal lobe recruited by diverse cognitive demands. *Trends in Neurosciences*, *23*, 475-83.

Duncan, J., Emslie, H., Williams, P., Johnson, R., & Freer, C. (1996). Intelligence and the frontal lobe: The organization of goal-directed behavior. *Cognitive Psychology*, *30*, 257-303. doi:10.1006/cogp.1996.0008

Duncan, J., Johnson, R., Swales, M., & Freer, C. (1997). Frontal lobe deficits after head injury: Unity

and diversity of function. *Cognitive Neuropsychology*, *14*, 713-741.

Duncan, J., Schramm, M., Thompson, R., & Dumontheil, I. (2012). Task rules, working memory, and fluid intelligence. *Psychonomic Bulletin & Review*, *19*, 864-70. doi:10.3758/s13423-012-0225-y

Duque, J., Labruna, L., Verset, S., Olivier, E., & Ivry, R. B. (2012). Dissociating the role of prefrontal and premotor cortices in controlling inhibitory mechanisms during motor preparation. *The Journal of Neuroscience*, *32*, 806-16. doi:10.1523/JNEUROSCI.4299-12.2012

Egner, T. (2008). Multiple conflict-driven control mechanisms in the human brain. *Trends in Cognitive Sciences*, *12*, 374-80. doi:10.1016/j.tics.2008.07.001

Egner, T., & Hirsch, J. (2005). Cognitive control mechanisms resolve conflict through cortical amplification of task-relevant information. *Nature Neuroscience*, *8*, 1784-90. doi:10.1038/nn1594

Elman, J. L. (1990). Finding structure in time. *Cognitive Science*, *14*, 179-211.

Ferrey, A. E., Frischen, A., & Fenske, M. J. (2012). Hot or not: Response inhibition reduces the hedonic value and motivational incentive of sexual stimuli. *Frontiers in Psychology*, *3*, 575. doi:10.3389/fpsyg.2012.00575

Field, M., Duka, T., Tyler, E., & Schoenmakers, T. (2009). Attentional bias modification in tobacco smokers. *Nicotine & Tobacco Research*, *11*, 812-22. doi:10.1093/ntr/ntp067

Fisher, A. V. (2011). Automatic shifts of attention in the dimensional change card sort task: Subtle changes in task materials lead to flexible switching. *Journal of Experimental Child Psychology*, *108*, 211-9. doi:10.1016/j.jecp.2010.07.001

Forrest, C. L., Elchlepp, H., Monsell, S., & McLaren, I. P. L. (2012). Task switching without knowledge of the tasks. *Proceedings Annual Meeting of the Cognitive Science Society, Sapporo, Japan*.

Frank, M. J. (2006). Hold your horses: A dynamic computational role for the subthalamic nucleus in decision making. *Neural Netw*, *19*, 1120-36. doi:10.1016/j.neunet.2006.03.006

Frank, M. J., & Badre, D. (2012). Mechanisms of hierarchical reinforcement learning in corticostriatal circuits 1: Computational analysis. *Cerebral Cortex*, *22*, 509-26. doi:10.1093/cercor/bhr114

Fröber, K., & Dreisbach, G. (2012). How positive affect modulates proactive control: Reduced usage of informative cues under positive affect with low arousal. *Frontiers in Psychology*, *3*, 265. doi:10.3389/fpsyg.2012.00265

Garavan, H., & Stout, J. C. (2005). Neurocognitive insights into substance abuse. *Trends in Cognitive Sciences*, *9*, 195-201. doi:10.1016/j.tics.2005.02.008

Goldstein, R. Z., & Volkow, N. D. (2011). Dysfunction of the prefrontal cortex in addiction: Neuroimaging findings and clinical implications. *Nature Reviews Neuroscience*, *12*, 652-69. doi:10.1038/nrn3119

Gollwitzer, P. M. (1999). Implementation intentions: Strong effects of simple plans. *American Psychologist*, *54*, 493-503.

Gollwitzer, P. M., & Sheeran, P. (2006). Implementation intentions and goal achievement: A meta-analysis of effects and processes. *Advances in Experimental Social Psychology*, *38*, 69-119.

Gollwitzer, P. M., Gawrilow, C., & Oettingen, G. (2010). The power of planning: Self-control by effective goal-striving. *Self Control in Society, Mind, and Brain*, 279-296.

Goschke, T. (2000). Intentional reconfiguration and involuntary persistence in task-set switching. In S. Monsell & J. Driver (Eds.), *Control of cognitive processes: Attention and performance XVIII* (pp. 331-355). Cambridge, MA: MIT Press.

Gottlieb, J., & Balan, P. (2010). Attention as a decision in information space. *Trends in Cognitive Sciences*, *14*, 240-8. doi:10.1016/j.tics.2010.03.001

Guitart-Masip, M., Huys, Q. J., Fuentemilla, L., Dayan, P., Duzel, E., & Dolan, R. J. (2012). Go and no-go learning in reward and punishment: Interactions between affect and effect. *NeuroImage*, *62*, 154-66. doi:10.1016/j.neuroimage.2012.04.024

Hagger, M. S., Lonsdale, A., Koka, A., Hein, V., Pasi, H., Lintunen, T., & Chatzisarantis, N. L. (2012). An intervention to reduce alcohol consumption in undergraduate students using implementation intentions and mental simulations: A cross-national study. *International Journal of Behavioral Medicine*, *19*, 82-96. doi:10.1007/s12529-011-9163-8

Hakamata, Y., Lissek, S., Bar-Haim, Y., Britton, J. C., Fox, N. A., Leibenluft, E., . . . Pine, D. S. (2010). Attention bias modification treatment: A meta-analysis toward the establishment of novel treatment for anxiety. *Biological Psychiatry*, *68*, 982-90. doi:10.1016/j.biopsych.2010.07.021

Hallion, L. S., & Ruscio, A. M. (2011). A meta-analysis of the effect of cognitive bias modification on anxiety and depression. *Psychological Bulletin*, *137*, 940-58. doi:10.1037/a0024355

Hazeltine, E., Lightman, E., Schwarb, H., & Schumacher, E. H. (2011). The boundaries of sequential modulations: Evidence for set-level control. *Journal of Experimental Psychology: Human Perception and Performance*, *37*, 1898-914. doi:10.1037/a0024662

Hazy, T. E., Frank, M. J., & O'Reilly, R. C. (2007). Towards an executive without a homunculus: Computational models of the prefrontal cortex/basal ganglia system. *Philosophical Transactions of the Royal Society of London. Series B, Biological Sciences*, *362*, 1601-13. doi:10.1098/rstb.2007.2055

Hogarth, L., Dickinson, A., Austin, A., Brown, C., & Duka, T. (2008). Attention and expectation in human predictive learning: The role of uncertainty. *Quarterly Journal of Experimental Psychology (2006)*, *61*, 1658-68. doi:10.1080/17470210701643439

Hommel, B. (2000). The prepared reflex: Automaticity and control in stimulus-response translation. *Control of Cognitive Processes: Attention and Performance XVIII*, 247-273.

Houben, K. (2011). Overcoming the urge to splurge: Influencing eating behavior by manipulating inhibitory control. *Journal of Behavior Therapy and Experimental Psychiatry*, *42*, 384-388. doi:10.1016/j.jbtep.2011.02.008

Houben, K., & Jansen, A. (2011). Training inhibitory control. A recipe for resisting sweet temptations. *Appetite*, *56*, 345-9. doi:10.1016/j.appet.2010.12.017

Houben, K., Havermans, R. C., Nederkoorn, C., & Jansen, A. (2012). Beer à no-go: Learning to stop responding to alcohol cues reduces alcohol intake via reduced affective associations rather than increased

response inhibition. *Addiction*, *107*, 1280-7. doi:10.1111/j.1360-0443.2012.03827.x

Huestegge, L., & Hazeltine, E. (2011). Cross-modal action: Modality matters. *Psychological Research*, *75*, 445-51. doi:10.1007/s00426-011-0373-0

Huestegge, L., & Koch, I. (2013). Constraints in task-set control: Modality dominance patterns among effector systems. *Journal of Experimental Psychology: General*, *142*, 633-7. doi:10.1037/a0030156

Iselin, A. M., & Decoster, J. (2009). Reactive and proactive control in incarcerated and community adolescents and young adults. *Cognitive Development*, *24*, 192-206. doi:10.1016/j.cogdev.2008.07.001

Izquierdo, A., & Jentsch, J. D. (2012). Reversal learning as a measure of impulsive and compulsive behavior in addictions. *Psychopharmacology*, *219*, 607-20. doi:10.1007/s00213-011-2579-7

Jones, A., & Field, M. (2013). The effects of cue-specific inhibition training on alcohol consumption in heavy social drinkers. *Experimental and Clinical Psychopharmacology*, *21*, 8-16. doi:10.1037/a0030683

Jones, A., Christiansen, P., Nederkoorn, C., Houben, K., & Field, M. (2013). Fluctuating disinhibition: Implications for the understanding and treatment of alcohol and other substance use disorders. *Frontiers in Psychiatry*, *4*, 140. doi:10.3389/fpsyt.2013.00140

Jost, K., Mayr, U., & Rösler, F. (2008). Is task switching nothing but cue priming? Evidence from erps. *Cognitive, Affective & Behavioral Neuroscience*, *8*, 74-84. Retrieved from PubMed.

Kastner, S., & Ungerleider, L. G. (2000). Mechanisms of visual attention in the human cortex. *Annual Review of Neuroscience*, *23*, 315-41. doi:10.1146/annurev.neuro.23.1.315

Keele, S. W. (1968). Movement control in skilled motor performance. *Psychological Bulletin*, *70*, 387-403.

Kharitonova, M., & Munakata, Y. (2011). The role of representations in executive function: Investigating a developmental link between flexibility and abstraction. *Frontiers in Psychology*, *2*, 347. doi:10.3389/fpsyg.2011.00347

Kiesel, A., Steinhauser, M., Wendt, M., Falkenstein, M., Jost, K., Philipp, A. M., & Koch, I. (2010). Control and interference in task switching—a review. *Psychological Bulletin*, *136*, 849-74. doi:10.1037/a0019842

Kiss, M., Raymond, J. E., Westoby, N., Nobre, A. C., & Eimer, M. (2008). Response inhibition is linked to emotional devaluation: Behavioural and electrophysiological evidence. *Frontiers in Human Neuroscience*, *2*, 13. doi:10.3389/neuro.09.013.2008

Klemen, J., Verbruggen, F., Skelton, C., & Chambers, C. D. (2011). Enhancement of perceptual representations by endogenous attention biases competition in response selection. *Attention, Perception & Psychophysics*. doi:10.3758/s13414-011-0188-5

Koch, I., & Allport, A. (2006). Cue-based preparation and stimulus-based priming of tasks in task switching. *Memory & Cognition*, *34*, 433-444.

Kool, W., McGuire, J. T., Rosen, Z. B., & Botvinick, M. M. (2010). Decision making and the avoidance of cognitive demand. *Journal of Experimental Psychology: General*, *139*, 665-82. doi:10.1037/a0020198

Lawrence, N. S., Verbruggen, F., Adams, R. C., & Chambers, C. D. (2013). Stopping to food cues reduces snack food intake: Effects of cue specificity, control conditions and individual differences. *In Preparation*.

Lenartowicz, A., Verbruggen, F., Logan, G. D., & Poldrack, R. A. (2011). Inhibition-related activation in the right inferior frontal gyrus in the absence of inhibitory cues. *Journal of Cognitive Neuroscience*, *23*, 3388-99. doi:10.1162/jocn_a_00031

Leotti, L. A., & Wager, T. D. (2010). Motivational influences on response inhibition measures. *Journal of Experimental Psychology: Human Perception and Performance*, *36*, 430-47. doi:10.1037/a0016802

Le Pelley, M. E., Beesley, T., & Griffiths, O. (2011). Overt attention and predictiveness in human contingency learning. *Journal of Experimental Psychology: Animal Behavior Processes*, *37*, 220-9. doi:10.1037/a0021384

Liefoghe, B., De Houwer, J., & Wenke, D. (2013). Instruction-based response activation depends on task preparation. *Psychonomic Bulletin & Review*. doi:10.3758/s13423-013-0374-7

Lijffijt, M., Kenemans, J. L., Verbaten, M. N., & van Engeland, H. (2005). A meta-analytic review of stopping performance in attention-deficit/hyperactivity disorder: Deficient inhibitory motor control? *Journal of Abnormal Psychology*, *114*, 216-22. doi:10.1037/0021-843X.114.2.216

Lipszyc, J., & Schachar, R. (2010). Inhibitory control and psychopathology: A meta-analysis of studies using the stop signal task. *Journal of the International Neuropsychological Society*, *1*-13. doi:10.1017/S1355617710000895

Lo, C. C., Boucher, L., Paré, M., Schall, J. D., & Wang, X. J. (2009). Proactive inhibitory control and attractor dynamics in countermanding action: A spiking neural circuit model. *The Journal of Neuroscience*, *29*, 9059-71. doi:10.1523/JNEUROSCI.6164-08.2009

Logan, G. D. (1978). Attention in character-classification tasks: Evidence for the automaticity of component stages. *Journal of Experimental Psychology: General*, *107*, 32-63.

Logan, G. D. (1988). Toward an instance theory of automatization. *Psychological Review*, *95*, 492-527.

Logan, G. D. (1994). On the ability to inhibit thought and action: A users' guide to the stop signal paradigm. In D. Dagenbach & T. H. Carr (Eds.), *Inhibitory processes in attention, memory, and language* (pp. 189-239). Academic Press.

Logan, G. D., & Bundesen, C. (2003). Clever homunculus: Is there an endogenous act of control in the explicit task-cuing procedure? *Journal of Experimental Psychology: Human Perception and Performance*, *29*, 575-599. doi:10.1037/0096-1523.29.3.575

Logan, G. D., & Burkell, J. (1986). Dependence and independence in responding to double stimulation - a comparison of stop, change, and dual-task paradigms. *Journal of Experimental Psychology: Human Perception and Performance*, *12*, 549-563.

Logan, G. D., & Cowan, W. B. (1984). On the ability to inhibit thought and action: A theory of an act of control. *Psychological Review*, *91*, 295-327.

Logan, G. D., & Gordon, R. D. (2001). Executive control of visual attention in dual-task situations. *Psychological Review*, *108*, 393-434. doi:10.1037/0033-295X.108.2.393

- Logan, G. D., Van Zandt, T., Verbruggen, F., & Wagenmakers, E. J. (2013). On the ability to inhibit thought and action: General and special theories of an act of control. *Psychological Review*, *in press*.
- Luck, S. J., Chelazzi, L., Hillyard, S. A., & Desimone, R. (1997). Neural mechanisms of spatial selective attention in areas V1, V2, and V4 of macaque visual cortex. *Journal of Neurophysiology*, *77*, 24-42.
- Luszczynska, A., Sobczyk, A., & Abraham, C. (2007). Planning to lose weight: Randomized controlled trial of an implementation intention prompt to enhance weight reduction among overweight and obese women. *Health Psychology*, *26*, 507-12. doi:10.1037/0278-6133.26.4.507
- Mackintosh, N. J. (1975). A theory of attention: Variations in the associability of stimuli with reinforcement. *Psychological Review*, *82*, 276.
- MacLeod, C. M. (1991). Half a century of research on the stroop effect: An integrative review. *Psychological Bulletin*, *109*, 163-203.
- Mars, R. B., Piekema, C., Coles, M. G., Hulstijn, W., & Toni, I. (2007). On the programming and reprogramming of actions. *Cerebral Cortex*, *17*, 2972-9. doi:10.1093/cercor/bhm022
- Mayr, U., & Bryck, R. L. (2005). Sticky rules: Integration between abstract rules and specific actions. *Journal of Experimental Psychology: Learning, Memory, and Cognition*, *31*, 337-50. doi:10.1037/0278-7393.31.2.337
- Mayr, U., & Kliegl, R. (2000). Task-set switching and long-term memory retrieval. *Journal of Experimental Psychology: Learning Memory and Cognition*, *26*, 1124-1140.
- McGuire, J. T., & Botvinick, M. M. (2010). Prefrontal cortex, cognitive control, and the registration of decision costs. *Proceedings of the National Academy of Sciences of the United States of America*, *107*, 7922-6. doi:10.1073/pnas.0910662107
- McLaren, I. P., Forrest, C. L., McLaren, R. P., Jones, F. W., Aitken, M. R., & Mackintosh, N. J. (2013a). Associations and propositions: The case for a dual-process account of learning in humans. *Neurobiology of Learning and Memory*. doi:10.1016/j.nlm.2013.09.014
- McLaren, I. P., Forrest, C. L., McLaren, R. P., Jones, F. W., Aitken, M. R., & Mackintosh, N. J. (2013b). Associations and propositions: The case for a dual-process account of learning in humans. *Neurobiology of Learning and Memory*. doi:10.1016/j.nlm.2013.09.014
- McLaren, I. P., Wills, A. J., & Graham, S. (2010). Attention and perceptual learning. In C. Mitchell & M. Le Pelley (Eds.), *Attention and associative learning: From brain to behaviour* (p. 131). Oxford: Oxford University Press.
- Meiran, N. (1996). Reconfiguration of processing mode prior to task performance. *Journal of Experimental Psychology: Learning Memory and Cognition*, *22*, 1423-1442.
- Meiran, N., Cole, M. W., & Braver, T. S. (2012). When planning results in loss of control: Intention-based reflexivity and working-memory. *Frontiers in Human Neuroscience*, *6*, 104. doi:10.3389/fnhum.2012.00104
- Meyer, D. E., & Kieras, D. E. (1997). A computational theory of executive cognitive processes and multiple-task performance: Part 1. Basic mechanisms. *Psychological Review*, *104*, 3-65.
- Miller, E. K., & Cohen, J. D. (2001). An integrative theory of prefrontal cortex function. *Annual Review of Neuroscience*, *24*, 167-202.
- Miller, G. A., Galanter, E., & Pribram, K. H. (1960). *Plans and structure of behavior*. New York: Holt, Rinehart & Winston.
- Minsky, M., & Parert, S. (1969). *Perceptrons*. In Cambridge, MA: MIT press.
- Mitchell, C. J., De Houwer, J., & Lovibond, P. F. (2009). The propositional nature of human associative learning. *The Behavioral and Brain Sciences*, *32*, 183-98; discussion 198-246. doi:10.1017/S0140525X09000855
- Miyake, A., Friedman, N. P., Emerson, M. J., Witzki, A. H., Howerter, A., & Wager, T. D. (2000). The unity and diversity of executive functions and their contributions to complex "frontal lobe" tasks: A latent variable analysis. *Cognitive Psychology*, *41*, 49-100. doi:10.1006/cogp.1999.0734
- Moffitt, T. E., Arseneault, L., Belsky, D., Dickson, N., Hancox, R. J., Harrington, H., . . . Caspi, A. (2011). A gradient of childhood self-control predicts health, wealth, and public safety. *Proceedings of the National Academy of Sciences of the United States of America*, *108*, 2693-8. doi:10.1073/pnas.1010076108
- Monsell, S. (2003). Task switching. *Trends in Cognitive Sciences*, *7*, 134-140.
- Monsell, S., & Driver, J. (2000). Banishing the control homunculus. In S. Monsell & J. Driver (Eds.), *Control of cognitive processes: Attention and performance XVIII* (pp. 3-32). Cambridge, MA: MIT Press.
- Monsell, S., & Mizon, G. A. (2006). Can the task-cueing paradigm measure an endogenous task-set reconfiguration process? *Journal of Experimental Psychology: Human Perception and Performance*, *32*, 493-516. doi:10.1037/0096-1523.32.3.493
- Mostofsky, S. H., & Simmonds, D. J. (2008). Response inhibition and response selection: Two sides of the same coin. *Journal of Cognitive Neuroscience*, *20*, 751-761.
- Mulder, M. J., Wagenmakers, E. J., Ratcliff, R., Boekel, W., & Forstmann, B. U. (2012). Bias in the brain: A diffusion model analysis of prior probability and potential payoff. *The Journal of Neuroscience*, *32*, 2335-43. doi:10.1523/JNEUROSCI.4156-11.2012
- Munakata, Y., Snyder, H. R., & Chatham, C. H. (2012). Developing cognitive control: Three key transitions. *Current Directions in Psychological Science*, *21*, 71-77. doi:10.1177/0963721412436807
- Muraven, M. (2010). Building self-control strength: Practicing self-control leads to improved self-control performance. *Journal of Experimental Social Psychology*, *46*, 465-468. doi:10.1016/j.jesp.2009.12.011
- Nederkoorn, C., Jansen, E., Mulkens, S., & Jansen, A. (2007). Impulsivity predicts treatment outcome in obese children. *Behaviour Research and Therapy*, *45*, 1071-5. doi:10.1016/j.brat.2006.05.009
- Newell, A. (1990). *Unified theories of cognition*. Harvard University Press
- Nigg, J. T. (2001). Is ADHD a disinhibitory disorder? *Psychological Bulletin*, *127*, 571-98.
- Noël, X., Brevers, D., & Bechara, A. (2013). A neurocognitive approach to understanding the neurobiology of addiction. *Current Opinion in Neurobiology*. doi:10.1016/j.conb.2013.01.018

- Norman, D. A., & Shallice, T. (1986). Attention to action: Willed and automatic control of behavior. In R. J. Davidson, G. E. Schwartz, & D. Shapiro (Eds.), *Consciousness and self-regulation: Advances in research and theory* (Vol. 4, pp. 1-18). New York: Plenum.
- Oberauer, K. (2009). Design for a working memory. *Psychology of Learning and Motivation*, 45-100.
- Pashler, H. (1994). Dual-task interference in simple tasks: Data and theory. *Psychological Bulletin*, 116, 220-44.
- Paxton, J. L., Barch, D. M., Racine, C. A., & Braver, T. S. (2008). Cognitive control, goal maintenance, and prefrontal function in healthy aging. *Cerebral Cortex*, 18, 1010-28. doi:10.1093/cercor/bhm135
- Pearce, J. M., & Hall, G. (1980). A model for pavlovian learning: Variations in the effectiveness of conditioned but not of unconditioned stimuli. *Psychological Review*, 87, 532.
- Pearce, J. M., & Mackintosh, N. J. (2010). Two theories of attention: A review and a possible integration. *Attention and Associative Learning: From Brain to Behaviour*, 11-39.
- Pessoa, L. (2009). How do emotion and motivation direct executive control? *Trends in Cognitive Sciences*, 13, 160-6. doi:10.1016/j.tics.2009.01.006
- Poldrack, R. A., Kittur, A., Kalar, D., Miller, E., Seppa, C., Gil, Y., . . . Bilder, R. M. (2011). The cognitive atlas: Toward a knowledge foundation for cognitive neuroscience. *Frontiers in Neuroinformatics*, 5, 17. doi:10.3389/fninf.2011.00017
- Posner, M. I. (1980). Orienting of attention. *Quarterly Journal of Experimental Psychology*, 32, 3-25.
- Postle, B. R. (2013). Activation and information in working memory research. In D. R. Addis, M. Barense, & A. Duarte (Eds.), *The wiley-blackwell handbook on the cognitive neuroscience of memory*.
- Purcell, B. A., Heitz, R. P., Cohen, J. Y., Schall, J. D., Logan, G. D., & Palmeri, T. J. (2010). Neurally constrained modeling of perceptual decision making. *Psychological Review*, 117, 1113-43. doi:10.1037/a0020311
- Ramamoorthy, A., & Verguts, T. (2012). Word and deed: A computational model of instruction following. *Brain Research*, 1439, 54-65. doi:10.1016/j.brainres.2011.12.025
- Ratcliff, R., & Smith, P. L. (2004). A comparison of sequential sampling models for two-choice reaction time. *Psychological Review*, 111, 333-67. doi:10.1037/0033-295X.111.2.333
- Redick, T. S., & Engle, R. W. (2011). Integrating working memory capacity and context-processing views of cognitive control. *Quarterly Journal of Experimental Psychology* (2006), 64, 1048-55. doi:10.1080/17470218.2011.577226
- Ridderinkhof, K. R., Ullsperger, M., Crone, E. A., & Nieuwenhuis, S. (2004). The role of the medial frontal cortex in cognitive control. *Science*, 306, 443-7. doi:10.1126/science.1100301
- Robbins, T. W. (2007). Shifting and stopping: Fronto-striatal substrates, neurochemical modulation and clinical implications. *Philosophical Transactions of the Royal Society of London. Series B, Biological Sciences*, 362, 917-32. doi:10.1098/rstb.2007.2097
- Robbins, T. W., Gillan, C. M., Smith, D. G., de Wit, S., & Ersche, K. D. (2012). Neurocognitive endophenotypes of impulsivity and compulsivity: Towards dimensional psychiatry. *Trends in Cognitive Sciences*, 16, 81-91. doi:10.1016/j.tics.2011.11.009
- Rogers, R. D., & Monsell, S. (1995). Costs of a predictable switch between simple cognitive tasks. *Journal of Experimental Psychology. General*, 124, 207-231.
- Rosenbaum, D. A. (1980). Human movement initiation: Specification of arm, direction, and extent. *Journal of Experimental Psychology. General*, 109, 444-74.
- Rosenbaum, D. A., & Kornblum, S. (1982). A priming method for investigating the selection of motor responses. *Acta Psychologica*, 51, 223-243.
- Rougier, N. P., Noelle, D. C., Braver, T. S., Cohen, J. D., & O'Reilly, R. C. (2005). Prefrontal cortex and flexible cognitive control: Rules without symbols. *Proceedings of the National Academy of Sciences of the United States of America*, 102, 7338-43. doi:10.1073/pnas.0502455102
- Rumelhart, D. E., Hinton, G. E., & Williams, R. J. (1986). Learning representations by back-propagating errors. *Nature*, 329, 534-536.
- Rummel, J., Einstein, G. O., & Rampey, H. (2012). Implementation-intention encoding in a prospective memory task enhances spontaneous retrieval of intentions. *Memory (Hove, England)*, 20, 803-17. doi:10.1080/09658211.2012.707214
- Rushworth, M. F., Walton, M. E., Kennerley, S. W., & Bannerman, D. M. (2004). Action sets and decisions in the medial frontal cortex. *Trends in Cognitive Sciences*, 8, 410-7. doi:10.1016/j.tics.2004.07.009
- Salinas, E., & Stanford, T. R. (2013). The countermanding task revisited: Fast stimulus detection is a key determinant of psychophysical performance. *The Journal of Neuroscience*, 33, 5668-85. doi:10.1523/JNEUROSCI.3977-12.2013
- Schall, J. D. (2001). Neural basis of deciding, choosing and acting. *Nature Reviews. Neuroscience*, 2, 33-42. doi:10.1038/35049054
- Schmitz, C., Jenmalm, P., Ehrsson, H. H., & Forssberg, H. (2005). Brain activity during predictable and unpredictable weight changes when lifting objects. *Journal of Neurophysiology*, 93, 1498-509. doi:10.1152/jn.00230.2004
- Schneider, D. W., & Logan, G. D. (2005). Modeling task switching without switching tasks: A short-term priming account of explicitly cued performance. *Journal of Experimental Psychology. General*, 134, 343-67. doi:10.1037/0096-3445.134.3.343
- Schneider, W., & Shiffrin, R. M. (1977). Controlled and automatic human information processing. I. Detection, search and attention. *Psychological Review*, 84, 1-66.
- Scholz, U., La Marca, R., Nater, U. M., Aberle, I., Ehlert, U., Hornung, R., . . . Kliegel, M. (2009). Go no-go performance under psychosocial stress: Beneficial effects of implementation intentions. *Neurobiology of Learning and Memory*, 91, 89-92. doi:10.1016/j.nlm.2008.09.002
- Schwabe, L., Dickinson, A., & Wolf, O. T. (2011). Stress, habits, and drug addiction: A psychoneuroendocrinological perspective. *Experimental and Clinical Psychopharmacology*, 19, 53-63. doi:10.1037/a0022212
- Shiffrin, R. M., & Schneider, W. (1977). Controlled and automatic human information processing: II. Per-

ceptual learning, automatic attending, and a general theory. *Psychological Review*, *84*, 127-190.

Sinha, R. (2008). Chronic stress, drug use, and vulnerability to addiction. *Annals of the New York Academy of Sciences*, *1141*, 105-30. doi:10.1196/annals.1441.030

Smith, P. L., & Ratcliff, R. (2004). Psychology and neurobiology of simple decisions. *Trends in Neurosciences*, *27*, 161-8. doi:10.1016/j.tins.2004.01.006

Spiegel, R., & McLaren, I. P. (2006). Associative sequence learning in humans. *Journal of Experimental Psychology. Animal Behavior Processes*, *32*, 150-63. doi:10.1037/0097-7403.32.2.150

Sternberg, S. (1969). The discovery of processing stages: Extensions of donders' method. *Acta Psychologica*, *30*, 276-315.

Stokes, M., & Duncan, J. (2013). Dynamic brain states for preparatory attention and working memory. In S. Kastner & A. C. Nobre (Eds.), *Oxford's handbook of attention*. Oxford, UK: Oxford University Press.

Stokes, M. G., Kusunoki, M., Sigala, N., Nili, H., Gaffan, D., & Duncan, J. (2013). Dynamic coding for cognitive control in prefrontal cortex. *Neuron*, *78*, 364-75. doi:10.1016/j.neuron.2013.01.039

Stroop, J. R. (1935). Studies of interference in serial verbal reactions. *Journal of Experimental Psychology*, *18*(643-662)

Sylvester, C. M., Shulman, G. L., Jack, A. I., & Corbetta, M. (2007). Asymmetry of anticipatory activity in visual cortex predicts the locus of attention and perception. *The Journal of Neuroscience*, *27*, 14424-33. doi:10.1523/JNEUROSCI.3759-07.2007

Theeuwes, J. (2010). Top-down and bottom-up control of visual selection. *Acta Psychologica*, *135*, 77-99. doi:10.1016/j.actpsy.2010.02.006

Thompson-Schill, S. L., Ramscar, M., & Chrysikou, E. G. (2009). Cognition without control-when a little frontal lobe goes a long way. *Current Directions in Psychological Science*, *18*, 259.

Toneatto, T., & Ladoceur, R. (2003). Treatment of pathological gambling: A critical review of the literature. *Psychology of Addictive Behaviors*, *17*, 284-92. doi:10.1037/0893-164X.17.4.284

Tzelgov, J. (1997). Specifying the relations between automaticity and consciousness: A theoretical note. *Consciousness and Cognition*, *6*, 441-51.

Ullsperger, M., Danielmeier, C., & Jochem, G. (2014). Neurophysiology of performance monitoring and adaptive behavior. *Physiological Reviews*, *94*, 35-79. doi:10.1152/physrev.00041.2012

Usher, M., & McClelland, J. L. (2001). The time course of perceptual choice: The leaky, competing accumulator model. *Psychological Review*, *108*, 550-592.

Vandierendonck, A., Liefoghe, B., & Verbruggen, F. (2010). Task switching: Interplay of reconfiguration and interference control. *Psychological Bulletin*, *136*, 601-26. doi:10.1037/a0019791

Veling, H., Aarts, H., & Papies, E. K. (2011). Using stop signals to inhibit chronic dieters' responses toward palatable foods. *Behaviour Research and Therapy*, *49*, 771-80. doi:10.1016/j.brat.2011.08.005

Veling, H., Aarts, H., & Stroebe, W. (2012). Using stop signals to reduce impulsive choices for palatable unhealthy foods. *British Journal of Health Psychology*. doi:10.1111/j.2044-8287.2012.02092.x

Veling, H., Holland, R. W., & van Knippenberg, A. (2008). When approach motivation and behavioral inhibition collide: Behavior regulation through stimulus devaluation. *Journal of Experimental Social Psychology*, *44*, 1013-1019. doi:10.1016/j.jesp.2008.03.004

Verbruggen, F., & Logan, G. D. (2008a). Automatic and controlled response inhibition: Associative learning in the go/no-go and stop-signal paradigms. *Journal of Experimental Psychology. General*, *137*, 649-72. doi:10.1037/a0013170

Verbruggen, F., & Logan, G. D. (2008b). Long-term aftereffects of response inhibition: Memory retrieval, task goals, and cognitive control. *Journal of Experimental Psychology: Human Perception and Performance*, *34*, 1229-35. doi:10.1037/0096-1523.34.5.1229

Verbruggen, F., & Logan, G. D. (2008c). Response inhibition in the stop-signal paradigm. *Trends in Cognitive Sciences*, *12*, 418-24. doi:10.1016/j.tics.2008.07.005

Verbruggen, F., & Logan, G. D. (2009a). Automaticity of cognitive control: Goal priming in response-inhibition paradigms. *Journal of Experimental Psychology: Learning, Memory, and Cognition*, *35*, 1381-8. doi:10.1037/a0016645

Verbruggen, F., & Logan, G. D. (2009b). Models of response inhibition in the stop-signal and stop-change paradigms. *Neuroscience and Biobehavioral Reviews*, *33*, 647-61. doi:10.1016/j.neubiorev.2008.08.014

Verbruggen, F., & Logan, G. D. (2009c). Proactive adjustments of response strategies in the stop-signal paradigm. *Journal of Experimental Psychology: Human Perception and Performance*, *35*, 835-54. doi:10.1037/a0012726

Verbruggen, F., Adams, R., & Chambers, C. D. (2012). Proactive motor control reduces monetary risk taking in gambling. *Psychological Science*, *23*, 805-15. doi:10.1177/0956797611434538

Verbruggen, F., Aron, A. R., Stevens, M. A., & Chambers, C. D. (2010). Theta burst stimulation dissociates attention and action updating in human inferior frontal cortex. *Proceedings of the National Academy of Sciences of the United States of America*, *107*, 13966-71. doi:10.1073/pnas.1001957107

Verbruggen, F., Liefoghe, B., & Vandierendonck, A. (2004). The interaction between stop signal inhibition and distractor interference in the flanker and stroop task. *Acta Psychologica*, *116*, 21-37. doi:10.1016/j.actpsy.2003.12.011

Verbruggen, F., Logan, G. D., Liefoghe, B., & Vandierendonck, A. (2008). Short-term aftereffects of response inhibition: Repetition priming or between-trial control adjustments? *Journal of Experimental Psychology: Human Perception and Performance*, *34*, 413-26. doi:10.1037/0096-1523.34.2.413

Verbruggen, F., Notebaert, W., Liefoghe, B., & Vandierendonck, A. (2006). Stimulus- and response-conflict-induced cognitive control in the flanker task. *Psychonomic Bulletin & Review*, *13*, 328-333.

Wang, C. H., Chang, C. C., Liang, Y. M., Shih, C. M., Chiu, W. S., Tseng, P., . . . Juan, C. H. (2013). Open vs. Closed skill sports and the modulation of inhibitory control. *PLoS One*, *8*, e55773. doi:10.1371/journal.pone.0055773

Wang, X. -J. (2013). The prefrontal cortex as a quintessential "cognitive-type" neural circuit. In D. T.

Stuss & R. T. Knight (Eds.), *Principles of frontal lobe function* (Second Edition ed., pp. 226-248). Cambridge University Press.

Waszak, F., Hommel, B., & Allport, A. (2003). Task-switching and long-term priming: Role of episodic stimulus-task bindings in task-shift costs. *Cognitive Psychology*, *46*, 361-413. doi:10.1016/S0010-0285

Welford, A. T. (1952). The "psychological refractory period" and the timing of high speed performance: A review and a theory. *British Journal of Psychology*, *43*, 2-19.

White, C. N., Ratcliff, R., Vasey, M. W., & McKoon, G. (2010). Using diffusion models to understand clinical disorders. *Journal of Mathematical Psychology*, *54*, 39-52. doi:10.1016/j.jmp.2010.01.004

Whitmer, A. J., & Banich, M. T. (2007). Inhibition versus switching deficits in different forms of rumination. *Psychological Science*, *18*, 546-53. doi:10.1111/j.1467-9280.2007.01936.x

Wiecki, T. V., & Frank, M. J. (2013). A computational model of inhibitory control in frontal cortex and basal ganglia. *Psychological Review*, *120*, 329-55. doi:10.1037/a0031542

Wiers, R. W., Gladwin, E., Hofmann, W., Salemink, E., & Ridderinkhof, R. (2013). Cognitive bias modification and cognitive control training in addiction and related psychopathology: Mechanisms, clinical perspectives, and ways forward. *Clinical Psychological Science*, *2*, 192-212. doi:10.1177/2167702612466547

Wiers, R. W., Eberl, C., Rinck, M., Becker, E. S., & Lindenmeyer, J. (2011). Retraining automatic action tendencies changes alcoholic patients' approach bias for alcohol and improves treatment outcome. *Psychological Science*, *22*, 490-7. doi:10.1177/0956797611400615

Wills, A. J., Lavric, A., Croft, G. S., & Hodgson, T. L. (2007). Predictive learning, prediction errors, and attention: Evidence from event-related potentials and eye tracking. *Journal of Cognitive Neuroscience*, *19*, 843-54. doi:10.1162/jocn.2007.19.5.843

Wimmer, G. E., & Shohamy, D. (2012). Preference by association: How memory mechanisms in the hippocampus bias decisions. *Science*, *338*, 270-3. doi:10.1126/science.1223252

de Wit, H. (2009). Impulsivity as a determinant and consequence of drug use: A review of underlying processes. *Addiction Biology*, *14*, 22-31. doi:10.1111/j.1369-1600.2008.00129.x

Wolfensteller, U., & Ruge, H. (2012). Frontostriatal mechanisms in instruction-based learning as a hallmark of flexible goal-directed behavior. *Frontiers in Psychology*, *3*, 192. doi:10.3389/fpsyg.2012.00192

Woodman, G. F. (2013). Viewing the dynamics and control of visual attention through the lens of electrophysiology. *Vision Research*, *80*, 7-18. doi:10.1016/j.visres.2013.01.003

Woodman, G. F., Luck, S. J., & Schall, J. D. (2007). The role of working memory representations in the control of attention. *Cerebral Cortex (New York, N.Y. : 1991)*, *17 Suppl 1*, i118-24. doi:10.1093/cercor/bhm065

Appendix

Table 1: Overview of popular paradigms to study action control and behavioural flexibility. Definitions are based on the Cognitive Atlas project (Poldrack et al., 2011). For more information about this project and other tasks, visit <http://www.cognitiveatlas.org/>. Note that this project also aims to increase the focus on the underlying processes. In the electronic version of this article, the task name is a hyperlink to the corresponding Cognitive Atlas page.

Task name	Manipulation	Main dependent variable(s)
Stop-signal task	<p>Participants usually perform a choice reaction time in which they have to respond as quickly as possible to a particular stimulus feature (e.g. colour, shape, identity, or location). On a minority of the trials, the go stimulus is followed by an additional signal (e.g. an auditory tone or a visual cue), which instructs participants to withhold their planned response.</p> <p>In the countermanding task, participants have to cancel a saccade towards a target when a fixation cross reappears.</p> <p>In the stop-change variant, participants have to cancel the planned manual response and execute an alternative response instead.</p>	The stop-signal reaction time (SSRT), which is the estimated covert latency of stopping (Logan & Cowan, 1984). Longer SSRTs are usually interpreted to reflect poorer inhibitory control.
Go/no-go task	Participants are instructed to respond as quickly as possible to go stimuli (e.g. letters) but to refrain from responding when a no-go stimulus is presented (e.g. a digit). Go events typically occur with higher frequency than no-go events.	The probability of responding on a no-go trial.
Psychological refractory period (PRP) task	Participants are presented with two stimuli to which they have to respond. The interval between the two is usually so brief that the second stimulus appears before the response to the first one is finished.	Response latency of the second response (RT2), often as a function of the delay between the two stimuli (SOA). The PRP effect refers to the decrease in RT2 as SOA increases.
Stroop task and variants	In the Stroop task, colour words are presented in various ink colours. Participants are instructed to respond to the ink colour and ignore the words. In incongruent stimuli, colour names and ink colours are non-matching. Related tasks include the picture-word naming task , in which words appear inside pictures of objects.	The congruency effect, which refers to the difference between incongruent and congruent or neutral (e.g. 'OOO' written in red) stimuli.
The Eriksen flanker task	A task in which participants view target stimuli to which they must make a simple lexical response. These stimuli are surrounded by flankers. Distracting flankers are typically associated with an opposite response ('incongruent'), whereas facilitating flankers are typically associated with the same response as the target stimulus ('congruent').	The congruency effect, which refers to the difference between incongruent and congruent items.

[Task-switching paradigm](#)

Participants frequently alternate between two or more tasks (e.g. naming the colour or identifying the shape of a stimulus). Which task they have to perform is often indicated by a cue (e.g. the task name or the location of the stimulus) or by a sequence they have to remember.

The difference between task-switch trials and task-repeat trials. Usually, switching from one task to another is slower and more error-prone than repeating the same task.

[\(Wisconsin\) Card Sorting Test](#)

The participant is presented with stimulus cards containing shapes. The cards differ in colour of the shapes, number of the shapes, and the form of the shapes. The participant is asked to sort these cards into two piles. The participant is not told what stimulus dimension to use in order to sort the cards, but feedback is provided to tell the participant if a particular match is correct. During the test, the sorting rules are changed and the participant must discover the new sorting rule in order to be successful.

The total number of categories achieved and the number of perseveration errors after a rule switch.

[Response-reversal learning](#)

Participants first learn to respond to stimuli based on feedback, followed by a reversal of the stimulus-action mapping. Participants have to overcome the old (habitual) response, and instead, execute an alternative novel response.

Proportion of correct responses before and after the reversal stage.
