**Why motor imagery isn’t really motoric: Towards a reconceptualization in terms of effect-based action control.**

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**Abstract**Overt and imagined action seem inextricably linked. Both have similar timing, activate shared brain circuits, and motor imagery influences overt action and vice versa. Motor imagery is therefore often assumed to recruit the same motor processes that govern action execution, and which allow one to play through or simulate actions offline. Here, we advance a very different conceptualization. Accordingly, the links between imagery and overt action do not arise because action imagery is intrinsically motoric, but because action planning is intrinsically imaginistic and occurs in terms of the perceptual effects one want to achieve. Seen like this, the term 'motor imagery' is a misnomer of what is more appropriately portrayed as 'effect imagery'. In this article, we review the long-standing arguments for effect-based accounts of action, which are often ignored in motor imagery research. We show that such views provide a straightforward account of motor imagery. We review the evidence for imagery-execution overlaps through this new lens and argue that they indeed emerge because every action we execute is planned, initiated and controlled through an imagery-like process. We highlight findings that this new view can now explain and point out open questions.

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Overt and imagined action seem inextricably linked. Before undertaking a difficult motor task, people often experience themselves imaging what they intend to do, and the form this imagination takes (e.g., imagining intended outcomes or motor behaviours) affects their success and subsequent learning (e.g., Land, Frank, & Schack, 2014; Woolfolk et al., 1985ab). Sometimes, people even imagine behaviours they will execute at a much later time and in a different environment, for example, when they mentally play through the actions of their sport from the privacy of their home. Again, this form of motor imagery – sometimes termed mental practice, mental training or motor imagery training (Schack, Essig, Frank, & Koester, 2014; for definitions and conceptualizations, see Morris, Spittle, & Watt, 2005) –affects later performance (for meta-analysis, Driskell, Copper, & Moran, 1994; Simonsmeier, Andronie, Buecker, & Frank, 2021; Toth, McNeill, Hayes, Moran, & Campbell, 2020) and is recommended by most professional coaches (Mayer & Hermann, 2019). Purely mental practice can even increase measured muscle strength, from simple finger contractions to leg pressing and triceps extension, albeit not to the same extent as physical practice (Yue & Cole, 1992, for recent replications and review see, Paravlik, Slimani, Tod, Marusic, Milanovic & Pisot, 2018; Reiser, Büsch, & Munzert, 2011; Smith, Collins, & Holmes, 2003).

Studies from experimental psychology and cognitive neuroscience support this coupling of overt and imagined action. There are tight correspondences between the timing of imagined and overt actions (Decety, Jeannerod, Prablanc, 1989; Wohlschläger & Wohlschläger, 1998; for a critical review, see Guillot & Collet, 2005), between the activated brain structures in parietal and premotor cortices (for reviews, see Lotze & Halsband, 2006; Hétu et al., 2013; O’Shea & Moran, 2017), and between the lawful regularities that govern the kinematics of both overt and imagined action (e.g., Fitt’s law, Decety & Jeannerod, 1995; two-thirds power law, Karklinsky & Flash, 2015; Papaxanthis, Paizis, White, Pozzo & Stucchi, 2012). Moreover, several studies show that motor imagery can engender (sub-threshold) activation in the muscles used in the imagined behaviour (Jacobson, 1931; 1932; Shaw, 1938; Munzert & Krüger, 2018; Lutz, 2003; Guillot et al., 2007; Guillot, Lebon, Rouffet, 2010), and, conversely, that executing motor actions makes imagining the same actions easier and imagining different actions harder (e.g., Wohlschläger, 1996; 2001; Callow et al., 2006; Guillot, Moschberger & Collet, 2013; for a broader review of the effects of such “dynamic motor imagery”, see Guillot, under review). The link from imagined to overt behaviour is so strong that it provides the basis for several (stage) magical phenomena. In Chevreul’s pendulum and the Ouija board, for example, seemingly supernatural motions happen simply because participants’ imagined motions are, unbeknownst to them, translated into subliminal hand and finger movements that are made visible by the devices (Chevreul, 1833; Easton & Shor, 1975, 1976, 1977; Wegner et al., 1998; Cantergi et al., 2021).

A standard explanation for these findings is that imagery of action is an intrinsically motoric process. This view assumes that motor imagery, in a form of *neural reuse* (e.g., Anderson, 2010), draws upon the same neuronal networks and cognitive processes that underlie action execution itself (Jeannerod, 1994; Jeannerod & Decety, 1995). As a potential mechanism, it has been proposed that the brain predicts – via forward models – the sensory consequences that each of its motor commands will produce, so that it can anticipate the visual, tactile, and proprioceptive sensations that will soon be registered (e.g., Miall & Wolpert, 1996; Sperry, 1950). During overt action, such predictions may allow the actor to filter out predicted sensations (e.g., Reichenbach, Franklin, Zatka-Haas & Diedrichsen, 2014) or to correct for movement errors before they happen (e.g., Desmurget & Grafton, 2000; Shadmehr, Smith & Krakauer, 2010). During imagery, the same forward models could be used offline, triggered perhaps by sub-threshold motor commands, and allow one to mentally play through how different actions will unfold, without the signals ever reaching the muscles (e.g., Jeannerod, 1994; Jeannerod & Decety, 1995; Kilteni, Andersson, Houborg & Ehrsson, 2018).

In these proposals, motor imagery is often described as “neural simulation of action” (e.g., Jeannerod, 2001), “covert execution” (e.g., Scheil, Kleinsorge & Liefooghe, 2020), and imagined actions are taken as “real actions, except for the fact that they are not executed” (Jeannerod, 2001, p. 103). In essence, these accounts hold that people can imagine their actions because the motoric structures of the brain can, in some form, *pretend* that the imagined actions are currently executed, and project their perceptual consequences into the imagination, so that one can watch them unfold in front of one’s mind’s eye. Imagination therefore has the same timing, is governed by the same regularities, and activates largely overlapping brain structures as overt action.

1. **A different view on motor imagery**

The previous section describes the “standard” conception of motor imagery. In this article, we would like to advance a markedly different conceptualization, which turns the proposed relationship on its head, and which – as we argue further below – provides a closer match to the extant data. On this view, the links between imagery and overt action do not arise because action imagery is intrinsically motoric, but – conversely – because the mechanisms people use to control their voluntary behaviour are intrinsically imaginistic. In other words, the observed overlaps emerge not because motor imagery recruits motor-based resources but because every action we execute is planned, initiated and controlled through imagery (e.g., Colton, Bach, Whalley & Mitchell, 2018; Hommel, 2009; Hommel, Müsseler, Aschersleben, & Prinz, 2001; Janczyk & Kunde, 2020; Pfister, 2019; Pfister, Janczyk, Wirth, Dignath, & Kunde, 2014; Prinz, 1997; Pfister, 2019; Shin et al., 2010).

This proposal is not new. In fact, it is the classic solution of the ideomotor theorists (e.g., Carpenter, 1852; James, 1890; Harleß, 1861; Lotze, 1852) to the puzzle of how people can achieve voluntary control over their body movements, given that they have very little actual insight into the actual working of their motor apparatus. Even the simplest act of reaching and grasping requires complex coordination of a multitude of muscles, most of which a person is not aware of, including those in one’s back that prevent one from falling forward when extending one’s arm. The reader could ask themselves, for example, *where* in fact the muscles they use to control their finger movements are located, *what* produces the sounds when they snap their fingers, or *how* they make a bicycle go left or right. Surprisingly, most people answer these questions incorrectly, even though they have performed the actions many times (see Footnote1 for answers). This “executive ignorance” (Turvey, 1977) into the motor activities that make up our daily lives was neatly summarized by William James: “we are only conversant with the outward results of our volition, and not with the hidden inner machinery of nerves and muscles which are what primarily sets it at work” (1890, p. 499).

Ideomotor theorists argue that imagery is the trick that people use to gain control of a motor apparatus that is essentially a black box to them. The idea is that people do not try to – and actually cannot – directly control their “hidden machinery of nerves of muscles” (James, 1890, p. 499). They can only ever bring to mind a mental image of the “outward results” that they want to achieve, which then activates the motor patterns which will bring this result about. The mechanisms that are assumed to underlie this transformation of imagery into action are surprisingly simple and can be accounted for by established laws of associative learning. Accordingly, human agents learn to link, through a lifetime process of self-observation, the different efferent/motor activities that they produce, at first accidentally (‘motor babbling’), with the perceptual effects these activities reliably cause (James, 1890; Prinz, 1997; Hommel & Elsner, 2001). In short, they learn how their body movements look, feel, and sound, and how they affect the environment. The main argument is that as soon as a behaviour and its likely effects are robustly associated, the mere *intention* to produce any perceivable effect activates the motor pattern to which it was associated by previous experience. Thus people can purposefully control their black box motor apparatus, merely by thinking of the perceptual effects they wish to achieve: We imagine what it looks like to move our fingers, we bring to mind the sound of our fingers snapping, or where we want our bicycle to go next – and by the previously formed associations the corresponding efferent activities are recollected, without us ever needing to know how they internally realized: imagery of the intended “outward results” is enough to elicit the motor behaviours itself.

While such proposals of effect-based action control have been only rarely connected to the phenomenon of motor imagery, they form the core of modern accounts of effect-based action control (for reviews, see Shin, Proctor, & Capaldi, 2010, Pfister, 2019), and are captured by recent ideas of action control through internal (inverse and forward) models (Wolpert, 1997) and predictive coding/active inference (e.g., Adams, Shipp, & Friston, 2013). They are supported by a large body of evidence. A full review is not possible here (see Pfister, 2019), but studies have shown that any manipulation that helps people bring to mind the effects of their actions indeed activates the motor behaviours themselves, in line with the idea that both are closely associated. For example, participants execute actions more quickly when primed, just before execution, with an image of the finger (or other body part) movements they should make (e.g., Brass, Bekkering, & Prinz, 2001; Bach, Peatfield, & Tipper, 2007), even if this prime is only anticipated but not perceived (Kunde, Koch, & Hoffmann, 2004; see Badets, Koch, & Philipp, 2016 for review). In some cases, this priming is enough to inadvertently cause participants to execute actions they have been asked to withhold (Colton et al., 2019). Even distal perceptual consequences of one’s actions (like a sound elicited by a button press) prime the motor behaviours that usually bring them about, provided they have been robustly associated before (e.g., Hommel & Elsner, 2001; Ziessler, Nattkemper & Vogt, 2012).

Neuroimaging research is also consistent with the proposed tight coupling of motor and effect-related representations. In the last decades, it has become clear that, throughout the cortical hierarchy, neuronal populations often code for both, efferent (motor) activities and the afferent (perceptual) states they produce, to the extent that it has become increasingly difficult to delineate purely afferent or efferent regions. Regions in early and late visual cortex, for example, previously thought to play purely perceptual roles, have been found to be involved in action planning, and specifically to encode the effects one wants to cause with one’s actions (Kühn, Keizer, Rombouts, & Hommel, 2011; van Steenbergen et al., 2017; Zimmermann et al., 2018; Zimmermann, Verhaegen, de Lange, & Toni, 2016). A similar “common coding” (Prinz, 1997) of perceptual and motor components also exists in the parietal lobe (e.g., Oosterhof, Tipper & Downing, 2012; Monaco, Malfatti, Culham, Cattaneo & Turella, 2020) and the premotor cortex, where neurons have been identified that code actions equally when they are executed and when their effects are perceived, both visually and auditorily (i.e., “mirror neurons”, di Pellegrino, Fadiga, Fogassi, Gallese & Rizzolatti, 1992; Gallese, Fadiga, Fogassi & Rizzolatti, 1996; Kohler, et al., 2002). Even the primary motor cortex, often implicated in motor output exclusively, seems to play perceptual roles by representing the anticipated proprioceptive/kinaesthetic consequences of one’s behaviour (Naito, 2004; Gandolla et al., 2014; de Lange, Rahnev, Donner & Lau, 2013). Indeed, it has been argued that the motor commands it sends to the spinal cord may be, in fact, nothing else than proprioceptively coded goal states for one’s limbs (e.g., Adams, Shipp, & Friston, 2013).

Together therefore, there is ample evidence from both experimental psychology and neuroscience that actions are represented in terms of their intended effects, which then activate the motor behaviours to which they are associated. Recent proposals have extended this simple associative architecture towards alternative mechanisms for acquiring motor-perceptual links (e.g., propositional relationships, Sun et al. 2022), describe how they can account for complex, hierarchical or multi-step actions (e.g., Moeller & Pfister, 2022; Moeller & Frings, 2019; Kachergis et al., 2014), and how motor behaviour can be dynamically adjusted in response to error, when actual motor output diverges from the intended goal states (Kunde et al., 2017; Adams et al., 2013, Wolpert, 1997).

If these ideas of effect-based action control are taken seriously, they lead to a subtle but – in our minds – illuminating re-conceptualization of motor imagery. Motor imagery, in such frameworks, does not rely on neural re-use of execution-related (efferent) structures in the brain, but instead reflects specifically the *perceptual* process through which people plan, initiate and control their action: the bringing to mind of the goal states – the effects – an actor wants to achieve: how they want their actions to look, sound and feel, when they are carried out. The difference to overt action is only that, during motor imagery, this imaginistic process is decoupled from the motor apparatus, to prevent the imagined action effects from inadvertently causing overt behaviour.

Several – not mutually exclusive – alternatives exist how this decoupling may work. One possibility is that efferent activities are somehow inhibited (e.g., Guillot, Di Rienzo, MacIntyre, Moran, & Collet, 2012; Rieger, Dahm, & Koch, 2017; Di Rienzo et al., 2014; Berthoz, 1996). Another possibility is that the activation threshold to trigger associated motor pattern is deliberately upregulated to prevent an automatic outflow of efferent activity (see also, Berthoz, 1996). Both alternatives would make it possible for people to plan/imagine actions freely, without being in danger of inadvertently releasing the associated motor behaviours. In fact, this increase of the execution threshold might be the very reason for why agents can become aware of their action imagery at all. During most everyday activities, people move their body without experiencing the mental images they use to control these movements, suggesting that even weak, subliminal action images can drive motor behaviour effectively (for evidence, see Kunde, 2004; Linser & Goschke, 2007). From an ideomotor perspective, the upregulation of the motor threshold may therefore be precisely what makes motor imagery possible. It allows people to imagine their actions strongly enough to be consciously experienced while still not eliciting overt behaviour.

A recent series of experiments (Colton et al., 2018) tested the idea that imagined actions are planned actions that are activated just below the execution threshold. Participants were asked to imagine – but not execute – different sequences of finger movements. In some trials, we unexpectedly strengthened this imagination through a visual cue that showed the effects of the action they currently imagined (e.g., a specific finger depressing), in the hope that this surprising additional activation would drive the action super-threshold and cause its involuntary release. This is exactly what was found. When imagery and visual cues were congruent and could therefore combine, participants found themselves sometimes executing actions they were asked to withhold. If, however, the visual cue did not match what was imagined, the likelihood of an accidental action slip was reduced compared to baseline (for similar evidence, see Kunde et al., 2004; Maslovat, Chua & Hodges, 2013).

Other studies show that overt action be effectively influenced even when the relevant images are week and remain subliminal, for example, when very briefly presented before participants make a relevant motor response (e.g., Kunde, 2004; Linser & Goschke, 2007). Importantly, a recent study showed that vividly imagined actions have stronger effects on subsequent motor behaviour than even actions that have been explicitly prepared for action (Toovey, Seiss & Sterr, 2021), consistent with the idea that motor imagery involves a hyper-activation of the movement images that people use to control their behaviour so that they can be consciously experienced, while at the same time being prevent from causing motor output.

This conceptualization of motor imagery as (hyper-activated) effect imagery differs from conventional accounts in that it does not require information from output-related components of the motor apparatus. Instead, it conceptualizes motor imagery simply as the planning-related processes that otherwise shape’s people overt motor behaviour (see also Glover and Baran, 2017; Jeannerod, 1995; Toovey, Seiss & Sterr, 2021) by specifying its goals as well as those more proximal components that are important for action success (e.g., movements speeds, specific trajectories around goals, etc.)Footnote 2. In other words, human agents can imagine actions because they have done them over and over in their daily life and have internalized how they look, sound, and feel. From then on, they can then simply recall this perceptual knowledge to produce a vivid experience of the actions they want to imagine.

Importantly, while this account does not *require* a contribution of motoric knowledge, it does not imply that imagery *cannot* make use of it. In a radical ideomotor view, efferent (“motor”) activities are bidirectionally associated to the effects they will cause (cf. Hommel et al., 2001; Hommel and Elsner, 2001; Hecht, Vogt & Prinz, 2001). Via these links, imagining or intending a specific effect can bring about the associated motor behaviours, but – once they are selected – the activated motor behaviours can also activate the effects they will most likely cause. This is functionally important because the effects used to *select* a motor behaviour and those it will *cause* do not need to be identical; any action will bring with itself changes that have not been explicitly intended. For example, one may intend to grasp a coffee mug, by anticipating the required visual and proprioceptive components of the movement trajectory towards the goal, as well as the specific points of contact with the mug, but without anticipating the tactile experience of one’s pullover’s sleeve moving across the forearm, when carrying out that movement.

Through these links, any motor behaviour that is activated may therefore enrich or further constrain imagery with what is most likely to happen when it is executed. This associative activation of the effects an action will *also* cause might be called a “prediction” and is functionally equivalent to the output of a forward model (cf. Kawato, 1999). Such a bi-directional associative architecture therefore fully implements the main functional mechanism of conventional accounts of imagery, through which (subliminal) motor activation can project ahead which perceptual effects it will most likely cause. The difference is that, in effect-based accounts, this motor-based prediction is not the default mode. Instead, it is considered as an additional, optional step, that one can draw upon only after motor activities have been accessed through the intended perceptual changes in the first place. There is still no pure “motoric” imagination that is not triggered by a prior imagination of the intended action outcomes. However, once a specific motor behaviour that implements these desired effects is activated, it can be supplemented or adjusted through associations to the effects this motor behaviour will *also* have, if carried out.

We believe that such effect-based accounts of motor imagery solve (or sidestep) most problems and inconsistencies in standard accounts. Note for example that, in standard accounts, imagery emerges from motor commands being fed to a forward model-like mechanism, which then plays through the sequence of perceptual consequences these actions will achieve. What is usually left open is where these motor commands come from in the first place. Why, if one already knows the sequence of movements one wants to imagine, is it necessary (a) to identify the precise motor command that brings about this imagination, and (b) how is the relevant chain of motor commands selected, so that it matches this (sometimes quite complex) imagination goal? This ambiguity is even more pronounced if one considers that the mechanisms (i.e., forward models, efference copies) that are proposed to serve as the basis for this imagination are usually assumed to have evolved to predict the outcomes of actions that the organism *does* indeed execute (i.e., for anticipatory error control or filtering out expected stimulation). How then is it possible (c) to fool these mechanisms into completing the same job for actions that one does decidedly *not* execute, because one only wants to imagine them? Effect-based accounts of motor imagery solve, or sidestep, all these problems by providing a straightforward associative account of how action imagery can affect the motor apparatus, and how, in turn, motoric activity can feed back into what people imagine.

1. **Accounting for the evidence**

We believe that a re-formulation of motor imagery as action planning decoupled from execution has various conceptual advantages over standard accounts. Here, we briefly review how the major findings, which are usually taken as evidence for the standard account, are explained by effect-based accounts of motor imagery.

*Motor imagery elicits muscular activity.* One central finding is that motor imagery often engenders measurable activation in the muscles involved in the action (e.g., Jacobson, 1931; 1932; Guillot et al., 2007). The traditional explanation is that this shows the working of an execution-related process, for example, the subliminal activation of motor routines that – via forward modelling – supply imagery with the relevant “material” to play through, such as the precise kinematics or timing of the action one imagines. In effect-based accounts, the explanation of this motor outflow is more straightforward. It simply reflects that the neuronal codes that make up one’s imagery of an action are the same ones that one would use – via the previously formed associations – to initiate this action. In such views, it is not surprising that some sub-liminal activation would persist and reach the relevant muscles, even if the motor threshold itself is not passed and no full action is released. In the words of William James “every mental representation of a movement awakens to some degree the actual movement which is its object” (James, 1890). Thus, while these motor activations may indicate that some form of action imagery takes place, they do not necessarily play a functional role in the imagery process.

Several findings suggest that this is exactly what is going on. Several studies show striking dissociations between imagery and motor processes. It has long been observed, for example, that muscular responses during mental practice are often only loosely related to the actions one tries to imagine (Heuer, 1985), and that motor imagery can elicit motor-like brain activity even if motor output is fully controlled (Zabicki et al., 2017). Moreover, motor imagery typically captures the (ideal) actions one would *want* to carry out but less so the actual motor behaviour that results, such as the errors one will make (e.g., Dahm & Rieger, 2019; Rieger, Martinez & Wenke, 2011) or the variability of one’s movements (Niziolek, Nagarajan & Houde, 2013). Such dissociations are problematic for accounts in which motor processes are the causal driver of action imagery, but they are of course fully in line with effect-based views where imagery is primarily perceptually driven, reflecting one’s prior knowledge of how these behaviours should look, sound and feel, and where motor involvement can enrich this imagery in some tasks but is by no means required (see above).

Other studies show that imagery of non-motor behaviour is often just as effective in causing overt behaviour as that of the motor behaviour itself, if this non-motor behaviour represents possible effects one can cause. One only needs to take a pendulum in one’s hand and image it is swinging: one’s hand will produce these swings, even though one does not imagine the hand itself moving, and is in fact often unaware that it does so (Chevreul, 1833; Easton & Shor, 1975, 1976, 1977; Wegner et al., 1998; Cantergi et al., 2021). Such findings can be replicated in lab-situations, where visualizing certain shapes is enough to induce a tendency to reproduce these shapes with one’s hands, even if these shapes belong to objects that one never has interacted with before (e.g., the circle shape of the moon), making any action imagery unlikely (Bach, Griffiths, Weigelt & Tipper, 2010). Both findings are much more in line with the idea that motor output during imagery is ultimately driven by bringing to mind the intended action *effects*, rather than the motor behaviours themselves.

A final example is tool use. For many tools, the effects they produce are clearly dissociable from the motor behaviours that cause them. For example, the Fulcrum effect (Kunde, Müsseler & Heuer, 2007) denotes the fact that in laparoscopic surgery, movements of the hand upwards produce downwards movements of the endoscopic tool, and vice versa (for a review of similar tool transformation findings, see Heuer & Sülzenbrück. 2013). Importantly, when people plan such actions, then bringing to mind the intended motions of the tool suffices to elicit the relevant motor behaviours, even if they are opposite to intended tool motion (Müsseler, Kunde, Gauspohl, & Heuer, 2008). For example, imaging a clockwise rotation of an object facilitates *counter-clockwise* manual rotations of a steering wheel, provided this produces an intended clockwise rotation of the task-relevant visual object (which applies to current displays in aviation, Janczyk, Pfister, Crognale, & Kunde, 2012). These findings therefore provide clear evidence that motor imagery – like action planning – occurs not in terms of motoric outflow, but in terms of the effects one wants to achieve with one’s actions (Hommel et al. 2001; Prinz, 1997), and there is evidence that also the neuronal coding of these actions seems to occur to a large extent in such an effect-based rather than motoric format (e.g., Ferrari, Rozzi & Fogassi, 2005).

*Action execution can help or hinder imagery*. A second class of important findings is that executing motor behaviours disrupts imagery of different actions, but helps imagery of similar ones (e.g., Wohlschläger, 1996; 2001; Callow et al., 2006; for a review, see Guillot, under review). In the standard view, these effects emerge because carrying out an action occupies one’s execution-related resources so that they cannot be deployed for motor imagery, or because a forward model that is fed by mismatching motor commands will also conjure up mismatching mental images. In principle, these explanations can also work in frameworks of effect-based action control, as they assume not only that effect imagery triggers motor behaviour, but also that motor behaviour, in turn, is associated to the relevant perceptual effects it will bring about (e.g., Hommel et al., 2001; Müsseler & Hommel, 1997). Executing movements that differ from imagination will therefore by necessity also conjure up inappropriate effect images, therefore interfering with the imagery process.

However, we suspect that there might be a simpler explanation for many of these findings. Executing a movement is not perception-free. Even if people are not looking at the body parts they move, they still receive (proprioceptive, tactile, etc.) feedback from their limbs and muscles. And of course, this feedback can also be congruent or incongruent with the perceptual effects one imagines, so that any interference can arise not only from *predicted* feedback, but also from the *actual feedback* that is inconsistent with what one currently imagines. A classic example is that people can mentally rotate an object more easily if they rotate their hands in the same direction, compared to the opposite direction (Wohlschläger, 1996; 2001; for a review of similar findings, see Guillot, under review). This does not necessarily reflect the anticipation of incongruent movement effects, but also the multisensory associations of clockwise proprioceptive feedback with clockwise visual motions (cf. de Lange, Helmich & Toni, 2006, Shenton, Schwoebel & Coslett, 2004).

Findings from deafferented patients – who receive no proprioceptive feedback – provide direct evidence for such multisensory associations. Usually, people find it difficult to draw when they only see the mirror image of their drawing hand, as there is a mismatch between visual and proprioceptive information: leftward hand movements look rightwards in the mirror and vice versa. To compensate, neurotypical agents often try to ignore such mismatching proprioceptive feedback, causing ‘functional haptic neglect’ (Liesner & Kunde, 2020; Heuer & Rapp, 2012)*.* Strikingly, however, the problems with mirror drawing are absent in deafferented patients, showing that the interference indeed largely emerges from a perceptual (proprioceptive to visual) locus (Lajoie et al., 1992).

*Temporal and kinematic similarities between imagery and action*. A central piece of evidence for motoric accounts is the tight temporal link between overt and imagined action (e.g., Decety, Jeannerod, Prablanc, 1989; Wohlschläger & Wohlschläger, 1998) and the finding that both are governed by similar regularities, such as Fitts law, or the two-thirds Power law between speed and curvature of biological movements (Decety & Jeannerod, 1995; Karklinsky & Flash, 2015; Papaxanthis et al, 2012). As these constraints are argued to emerge from the motor system, their re-occurrence in imagery is taken as evidence that imagery must reflect a “read out” of a motoric process.

Please note that such findings can be accommodated in effect-based accounts, if one assumes that not only activated action images can activate the efferent activities that bring them about (i.e., the inverse route), but also that activated motor actions can activate the perceptual effects they produce (i.e., the forward route), and in doing so can retrieve perceptual consequences that are not fully specified in the original action plan (see above). Nevertheless, we do believe that many of the findings that are usually provided as evidence for such influences should not be taken at face value. Typical research participants have ample perceptual experience with their own behaviour (and that of others) throughout their lifetime, and it would be no surprise that their imagination – and ability to plan their actions – has internalized these regularities and takes them into account. A strong case for motoric readout could therefore only be made in cases where imagery can be shown to be governed by a regularity that (a) reflects constraints from the motor apparatus, and which (b) participants had no prior perceptual experience with. We are not aware of any such demonstration (but see Rieger, Boe, Ingram, Bart, Dahm, under review, this volume, for a different view**)**. Instead, several studies have suggested that regularities originally attributed to the motor apparatus may in fact reflect perceptual or imaginistic constraints. For example, it is well-established that people find it almost impossible to perform particular irregular bimanual movements, for example when having to describe three circles with one hand while the other hand performs four (Kelso, 1995). These limitations have been put down to constraints of the motor apparatus, such as a synchronization of motor commands in the spinal cord. However, in an impressive series of demonstrations, Mechsner and colleagues (Mechsner, Kerzel, Knoblich & Prinz, 2001; Mechsner, 2003) have shown that these phenomena may instead reflect constraints of imaginistic action planning. The limitations were alleviated when participants were given help to perceptually coordinate – imagine – these actions, so that “impossible” movements suddenly became possible (for an earlier demonstration of similar effects, see Swinnen, Lee, Verschueren, Serrien & Bogaerds, 1997; for a review of related effects, see Shea, Buchanan & Kennedy, 2016).

In addition, many of the initial findings of tight temporal overlaps between overt and imagined actions have been recently challenged, with several showing either consistent over- or under-estimation of the timings of imagined actions, for different task and stimulus contexts (e.g., Reed, 2002; for a review, see Guillot & Collet, 2005). This temporal elasticity is problematic for approaches that assume that people’s mental motor imagery emerges from a subliminal engagement of the underlying motor routines, which should then closely match their actual time course. However, if imagery reflects a planning process that is decoupled from execution (see also Jeannerod, 1995; Glover & Baran, 2017), such differences are not surprising, and indeed expected, if one assumes that people flexibly decide to either faithfully follow their memories of action execution, or go through their major functional stages only, or even speed up or slow down specific steps to extract the information required for their particular imagery goal.

*Imagery and action activate similar brain regions*. A final important piece of evidence for motoric accounts is the tight overlap between brain structures activated by overt and imagined action (for reviews, see Lotze & Halsband, 2006; O’Shea & Moran, 2017). In our view, an interesting, but often ignored, aspect of these observations is that this overlap is primarily observed in structures associated with *action planning* – like premotor and parietal cortices – but it is much less robust in execution-related structures, like the primary motor cortices, basal ganglia and cerebellum (for review and meta-analysis, see O’Shea & Moran, 2017; Hétu et al., 2013; Munzert, Lorey & Zentgraf, 2009). For motoric accounts of action imagery, such a finding should be puzzling. Yet it is exactly what would be predicted by accounts of effect-based action control, where action imagery is conceptualized precisely as a shift towards planning away from execution-related resources (see also Jeannerod, 1995; Baran & Glover, 2017).

An elegant study confirmed this link between motor imagery and planning- instead of execution-related processes, while controlling for actual body movement (Raffin et al., 2012). The authors asked amputees, who still subjectively felt the presence of their phantom limbs, to either really move or imagine moving their absent limbs. No movement – and no proprioceptive feedback – was of course present in either case. Nevertheless, there were clear differences in brain activation. The execution instruction more strongly activated primary somatosensory and motor cortices and the anterior lobe of the cerebellum. In contrast, during imagination there was more activity in the parietal and occipital lobes, and the posterior cerebellum (see also Hanakawa, Dimyan & Hallett, 2008). Importantly, it was found that these regions matched those when the patients really moved, or only imagined moving, their still intact limbs on the other side. This shows both: that motor execution and imagery are indeed different, and that these differences map onto the proposed shift from motor to planning-related resources of effect-based control.

1. **Open questions and outlook**

Motoric views of motor imagery have left several puzzles open that have either been unresolved, or not been addressed. An effect-based view provides a new view on them, and – in many cases – clear avenues how they can be resolved, including testable predictions. Below we provide a first sketch for some of these open questions.

*When does imagery draw upon motor resources?* In effect-based accounts, motor imagery primarily reflects the planning stages of actions, that is, the bringing to mind of how the target actions would manifest perceptually, based on one’s prior experience with them. In contrast to conventional accounts, there is usually no need for imagery to draw upon knowledge encoded by the motor apparatus, for example, by accessing a forward model that is fed with the motor commands that would generate the behaviours one wants to imagine. As pointed out above, this does not mean that such an influence is not possible, however. The simple associative architecture of effect-based control predicts not only that the effects one intends will elicit the motor behaviours that will bring these effects about, but also that – via the same associations – the motor behaviours, once selected, activate the effects they will produce (e.g., Hommel et al., 2001; Müsseler & Hommel, 1997). These latter pathways therefore allow one to derive how one’s behaviours will play out, providing a secondary “motoric” path to imagery.

An important question is under what circumstances this secondary route is made use of. We suspect that this happens specifically when the effects usually used to plan a particular action do not correspond to the information required by the imagery task. For example, it is typically assumed that the effects agents use to control their behaviour are coded in a relatively sparse manner, capturing mainly the intended “distal” bodily or environmental effects (Hommel et al., 2001), the required transitions between current and intended perceptual states (Kunde, Schmidts, Wirth, & Herbort, 2017), or the major functional sub-steps of an action sequence (Schack & Mechsner, 2006). As the specific timing and kinematic of one’s skilled movements emerge, to a large extent, from biomechanical constraints within the muscular and skeletal system during execution, they are most likely not represented on this planning level, unless the skill specifically requires controlling them (e.g., in dance, or for unusual actions cf., Mechsner et al., 2001). If this is correct, then the motoric route might be necessary whenever the imagery task requires access to action features that are not captured by the perceptual goal states people use to control their actions.

Accordingly, for motor behaviour where timing and kinematics do not need accurate endogenous controlled, imagery should similarly only engender limited activation in brain structures involved in action execution, and outflow to muscles should be limited. However, as soon as one’s imagery task requires deriving accurate timing, precise kinematics, or access to internal bodily feedback, such motoric effects should emerge also for these movements. Several studies show that motoric indices of motor imagery indeed emerge for such instructions (e.g., Stinear et al., 2006), and this notion is also well supported by the implicit practice of motor imagery researchers. To ensure that motoric activations are found, researchers often instruct their participants to focus on the proprioception and kinesthesis, or the sensations of the actions, not just their visual representation, or tasks are made sufficiently complex to require such forward modelling.

*Why is mental practice effective?* One rarely addressed conflict in standard approaches is the evidence that mental practice is particularly effective in conditions in which it should not be. If mental practice relies on a subliminal play-through of motor routines, why is it more effective in sports with strong cognitive instead of motor components (like chess, golf, etc; see Ryan & Simons, 1981; 1983 for the original distinction), why are effect sizes usually smaller for purely motoric ones (for meta-analyses, see Driskell et al., 1994; Toth et al., 2020; Weinberg, 2008), and why is the myoelectric activity that accompanies it only loosely connected to the practiced movements (Heuer, 1985)? The findings are similar for studies that test how learners are best instructed in a motor skill. It is usually better to instruct people in terms of the consequences – the effects – they want to achieve their actions (e.g., focus on hitting the darts board in the bullseye) than reminding them of the proper motoric technique (pulling back the hand, flick of the finger, etc), which can even be counter-productive (for review, see Wulf, 2013).

If one assumes that mental practice is effective because it affords a subliminal playthrough of motor routines, stronger benefits in the *absence* of motoric instructions should be at least slightly puzzling. However, such results are of course precisely what would be expected under and effects-based imagery view. In such views, playing through one’s action would simply help in deriving better plans for one’s actions, without necessary input from efferent/motor activities. Repeated effect imagery alone can (a) make these effect images more accessible so that they are more easily brought to mind in the performance situation, it (b) can make these images sharper and more salient so that they become better drivers of muscular activities, and (c), especially for longer action sequences, it might help one to imagine different sub-steps and schedule them more efficiently.

Consistent with these views, recent work shows that athletes represent skilled actions (such as a golf putt) in a hierarchical manner (Schack & Mechsner, 2006). These perceptual-cognitive representations describe the interrelations between the action’s functional parts, each of them being linked to particular perceptual effects that need to be achieved by that component of the motor skill (Schack, 2004, 2020; Bläsing, Tenenbaum, & Schack, 2009). When novices physically practice these skills physically these mental representations become stabilized and structurally revised into functional groups (e.g., a preparation phase, a clubhead-ball impact phase, and an attenuation phase after the putt; Frank, Land, & Schack, 2013; Frank, 2016).

From an effect-based view, mental practice can similarly stabilize these acquired skill structures and cause similar revisions as physical practice (Frank, 2014; Schack, 2006; Schack & Frank, 2019). Indeed, when novices practice mentally, their perceptual-cognitive representation system shows such changes to a skill’s cognitive representation that match those after physical practice (Frank, Land, Popp, & Schack, 2014). Strikingly, however, while changes in novices’ representations also lead to overt behavioural changes after physical practice, they do not do so necessarily after mental practice, unless athletes had some prior physical experience in which the different parts of a skill’s structure were associated with the relevant motor activities that reliably produce these effects (Frank et al., 2014). In this way, the effect-based approach suggests some clear boundary conditions when mental practice should be effective, pointing specifically to the requirement of pre-existing ideomotor associations that can link the mentally practiced effects to the efferent activities that would achieve them (i.e., an inverse model).

An interesting avenue for future research are attempts to enrich actions by additional perceptual feedback (e.g., sonification for the case of auditory feedback, e.g., Effenberg, Fehse, Schmitz, Krueger, & Mechling, 2016). The idea here is to provide agents with auditory feedback of their body movements that is more discriminable than the perceptual (visual, interoceptive) feedback these movements naturally produce. This should not only enable agents to distinguish motor patterns that were not discriminable without such feedback, but also provide them with another means to imagine their movements. They can imagine how it *sounds* to move in specific ways, rather than just visually or proprioceptively imagine doing so, just as musicians plan their playing in terms of these musical effects instead of body movements (Brooks, 1995; Drost, Rieger, Brass, Gunter, & Prinz, 2005ab).

*Integration of motor with non-motor imagery?* Most everyday tasks require the integration of own action capabilities with the characteristics of the environment. For example, for a successful golf swing, the golfer needs to coordinate their (upper) body and limbs with respect to the grass surface, the club one uses, direction and speed of the wind, among other factors. Similarly, during everyday tool use, people need to consider motoric features (e.g., the flexion and extension of arms during hammering), relative to the mechanical properties of the tool and the object it is applied to (e.g., a nail vs. a pane of glass) (see Osiurak & Badets, 2016, for a review of mechanical problem solving in tool use).

An account that can describe how such actions are planned, or imagined, needs a means of integrating motoric with extra-motoric information form the physical and social environment (see Bach et al., 2005; 2010; 2014 for similar ideas in an action observation context). It is difficult to see how such integration would be implemented in conventional accounts of motor imagery that rely on a motor-to-perceptual route only, as it is unclear how non-motoric knowledge about the physical behaviour of objects could interact with motoric knowledge of one’s own action (but see Schubotz, 2007 for an interesting approach). However, if one assumes, as effect-based accounts do, that action planning generally relies on the same cognitive codes as imagery of physical events, such an integration is no mystery. One’s perceptual learning about the behaviour of physical system and about one’s own motor behaviour can be seamlessly integrated (see Prinz, 1992, for a similar argument in action control).

Some recent evidence from monkey single cell recordings studies shows that the perceptual systems indeed support such integration. In one study, monkeys first had to identify a relevant object, then trace its outline, and then direct an action towards its endpoint. Early visual cortex played a central role in each of these steps, initially highlighting the relevant object, then tracing its outline, and then its endpoint, which then served as the basis for eye movements towards it (Moro, Tolboom, Khayat, & Roelfsema, 2010, Roelfsema, Khayat, & Spekreijse, 2003; for a review, see Roelfsema & de Lange, 2016). These findings are therefore exactly in line with effect-based account, where relevant paths are first perceptually organized, and the exact same representations can then form as basis for action control. We are looking forward to future studies that show such interactions in human participants.

*Imagery (and control) of complex actions.* Imagery, especially for training purposes, often involves complex actions, such as the playing through of a martial arts kata, the complex motions of a platform dive, or even the different stages of a reach-to-grasp action (e.g., Driskell et al., 1994; Simonsmeier et al., 2020; Toth et al., 2020; Morris, Spittle & Watt, 2005). In contrast, most empirical research on ideomotor control focusses on much simpler actions, which involve mostly discrete stimuli and responses, and which occur for the most part ballistically (e.g., button presses). It may therefore seem challenging for effect-based approaches to provide an account for how more complex actions are controlled – and therefore imagined.

We believe that, fundamentally, the difference between the simple actions used in classical ideomotor experiments and those more complex actions is probably smaller than one may assume, as even the simple button presses involve a carefully orchestrated sequence and press and release of multiple muscles, and even very simple action effects (such as visual cue) need to be distinguished in often complex ways (in terms, of shape, timing, location, etc.) from other unrelated stimuli that can occur at the same time. Thus, many of the difficulties that characterize complex actions are already present for the simple actions that have been subject of most research in the lab.

A central assumption is that the representations used for effect-based control of both simple and complex actions are organized in a hierarchical fashion, with higher-level (distal) action goals at the top, and the required sequence of more (proximal) body-related effects at the bottom, each robustly linked to the efferent/motor activities that would bring these effects about (e.g., Hommel et al., 2001, Hommel, 2009; for a recent review, see Moeller & Pfister, 2022). During execution, imaginistic control can then access – and modify – each of these more proximal action components to fulfil task requirements, such as making a less forceful button press than usual (Cao, Kunde, & Haendel, 2020), varying the musical notes one is playing in a longer piece (e.g., Keller & Koch, 2008), or forming a more pronounced circular trajectory than usual during gesturing (e.g., Bach et al., 2010). Moreover, during learning more complex action hierarchies can be built form the ground up from simpler elements and assembled into larger chunks (e.g., Moeller & Pfister, 2019). Indeed, a large variety of work now reveals effect-based control in more complex actions (e.g., throwing, Land, 2018), music sequences (e.g., Keller & Koch, 2008), gesturing (Bach et al., 2010), motor sequence learning (e.g., Stöcker & Hoffmann, 2004; Brown, Friedgen & Koch, 2022) or in tool use (e.g., Massen & Prinz, 2009; Müsseler et al., 2008; for review and theoretical argument, see Badets & Osiurak, 2017), demonstrating that the described ideomotor principles do indeed apply such more complex behaviours.

When imagining such complex actions, imagery can then make use of the existing hierarchies, just as would be the case for imagery of more simple action. Similar to how they would plan the execution of such actions, agents can then simply step through the sequence on the desired level of the hierarchy, either in terms of the major action steps (e.g., the route when driving home from work), or by accessing the more fundamental proximal representation that reflect the body movements required to achieve these goals, explaining perhaps the variety in how the timing of imagined actions matches or does not match the timing of overt execution (for a review, see Guillot & Collet, 2005). Indeed, studies that probe the cognitive representation of more complex skills show that the same hierarchies underlie imagery and overt action, so that less effective cognitive representations translate into less effective action (e.g., Schack & Mechsner, 2006; for a review, see Land, Volchenkov, Bläsing & Schack2013), and that changes during imagery of a known motor skills induce analogous changes during overt action, and vice versa (Frank et al., 2014).

A related challenge is how imagery can model non-ballistic behaviours in which one’s required behaviour in one step depends on variations with the previous steps, for example when correcting errors in performance, or when compensating for unexpected contributions from external influences (e.g., a partner’s movements in dance, or a wobble of the balance board one stands on). To flexibly control – and imagine – such dynamic responses to feedback, it has been argued that proximal effect representations may not only code discrete perceptual end states, but perceptual *transitions* between a given (or imagined) perceptual state and an intended goal state (e.g., Kunde et al, 2017), or reduce the difference between both (i.e., prediction error minimization, Adams et al., 2013; Wolpert, 1997). Note however that feedback from one’s own actions – or from such external sources – is missing during imagery. Executive resources might therefore be necessary to supply such information (Glover & Baran, 2017; Martel & Glover, 2022), or imaginers might need to engage the “forward mode” of ideomotor learning to more directly retrieve which outcomes the particular motor behaviours they imagined will have on the next components of their actions. Imagery of such multistep actions, where one component depends on the previous ones, may therefore be a key example where motor imagery engages not only planning-related resources, but on more execution-related mechanisms to derive the specific outcomes one’s (imagined) motor behaviours will have.

*Evolution of motor imagery.* The ability to imagine one’s actions undoubtedly brings substantial advantages to human behaviour, allowing actors to play through different action alternatives in their mind before committing to one (e.g., Bennett, 2021), or to practice their skills offline before testing them “in the wild” (e.g., Simonsmeier et al, 2020; Toth et al., 2020). An interesting question is therefore when these imagery abilities, and the effect-based control of action on which they build, have evolved. Some theorists propose a co-evolution of ideomotor-like mechanisms together with abilities for action prediction, tool use and language (e.g., Badets & Osiurak, 2017). Others, in contrast, argue for an earlier emergence of effect-based control, and that even the behaviour of simple organisms is well-described not in terms of a simple stimulus-response learning, but as an attempt to resolve, through the action, the mismatch between perceptually defined (homeostatic) goal states, and the organism’s actual state (i.e., active inference, Adams et al, 2013; Friston, 2013).

Several pieces of evidence point, in our mind, towards such an earlier evolution. For example, the phenomenon of outcome devaluation (Adams & Dickinson, 1981) marks the finding that even strongly overlearned stimulus-response associations (i.e., habits) do not lead to action when the associated outcomes are undesirable in the given situation. This and related phenomena are present in several animals, certainly in rodents (for a review, de Wit & Dickinson, 2009). At least for these animals, therefore, action planning already occurs on the basis of desired action outcomes and suggests an emergence of effect-based control already in non-human mammals.

Other findings suggest that also the ability to imagine behaviours and their outcomes, without overt motor output, is present in animals. It is known since Tolman (1948) that rats sometimes stop at critical junctions in a maze, as if considering – playing through in their mind – which path leads to the better outcome. Indeed, a plethora of studies since then suggest that is exactly what happens (for a review, see Reddish, 2016). Rats that exhibit this “vicarious trial and error” behaviour are more successful in their navigation and arrive at the reward shortly after. Moreover, recording of hippocampal place cells confirm that during these instances the rat mentally travels along the paths within the maze, with associated firing of reward-related brain regions once the reward is (virtually) reached. This suggests again that rats, and potentially several species of birds (Sulikowski & Burke, 2015), have some abilities for motor imagery in the absence of action.

We are very much agnostic about this debate. For the account of motor imagery presented here it is only necessary that these abilities are fully formed in humans. We note however that the ideas of effect-controlled action provide an excellent framework to account for the findings in animals above. We therefore suspect that what sets the remarkable human action planning abilities apart from that of animals is not the ability for effect-based control and imagery itself, which may be more fundamental than usually assumed, but the ability to *withhold* action when imagining. The key evolutionary step would then be the development of the prefrontal cortex, and the associated cognitive resources for response inhibition or up-regulating one’s motor threshold. These abilities would enable humans to imagine their actions in peace, without engaging in overt action, while still being able to derive the consequences these actions will have, therefore opening up capacities for foresight, planning for (currently) counterfactual realities, and practicing skills in the absence of overt action.

1. **Conclusions**

We have briefly sketched how motor imagery can be conceptualized in effect-based views of action control. These views hold that actions are planned in term of their intended perceptual effects – how a successful action should look, feel, and sound –, which are then made reality by automatic motor processes. In such views, motor imagery reflects this imaginistic planning process, which is decoupled from execution. Imagery experience simply reflect the recall of one’s perceptual (visual, auditory, proprioceptive) experience with one’s actions, without requiring efferent (motoric) contributions. If these views are taken seriously, the term “motor imagery” becomes a misnomer; “effect imagery” might be a better description of how people imagine – and plan – their (motor) actions. The evidence reviewed here makes an explicit case for such an account.

**Compliance with Ethical Standards**

Conflict of interest: The authors declare that they have no conflict of interest.

Ethical approval: This article does not contain any studies with human participants or animals performed by any of the authors.

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**Footnotes**

1. Many of the muscles that control finger movements are located in the arms, not the fingers. The particular sound when snapping the fingers is produced when the finger hits the palm of one’s hand, not by the fingers rubbing together. One makes a bicycle go left or right by leaning in this direction, not by moving the handlebars.
2. Our terminology throughout the manuscript follows the classic distinction between action planning and control, which is central to both effect-based (e.g., Hommel et al., 2001; Wühr & Müsseler, 2001; Thomaschke et al. 2012) and conventional accounts of motor control (e.g., Woodworth, 1899; Glover, 2004). While both planning and execution are important components of successful action, they subserve different functions. Planning captures primarily cognitive and volitional contributions, that is, the setting of action goals and relevant parameters (e.g., the speed of the action, the intended kinematics, trajectory, etc.). Control, in contrast, reflects the downstream (efferent) processes, which ensure that these goals are achieved, by guiding motor output towards the specified goal states. Of course, in practice, the boundary between both components is fluid, and research often observes a gradual transition between planning-related contributions at the start of the action, towards more control-related contributions at the end (for a review, see Glover, 2004). Moreover, recent proposals argue that even those downstream execution related processes might be mediated by ideomotor-like mechanisms that move our effectors towards proprioceptively defined goal states (e.g., Adams et al., 2013).