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Functional imaging studies of emotion regulation: a synthetic review and evolving model of the cognitive control of emotion

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This paper reviews and synthesizes functional imaging research that over the past decade has begun to offer new insights into the brain mechanisms underlying emotion regulation. Toward that end, the first section of the paper outlines a model of the processes and neural systems involved in emotion generation and regulation. The second section surveys recent research supporting and elaborating the model, focusing primarily on studies of the most commonly investigated strategy, which is known as reappraisal. At its core, the model specifies how prefrontal and cingulate control systems modulate activity in perceptual, semantic, and affect systems as a function of one's regulatory goals, tactics, and the nature of the stimuli and emotions being regulated. This section also shows how the model can be generalized to understand the brain mechanisms underlying other emotion regulation strategies as well as a range of other allied phenomena. The third and last section considers directions for future research, including how basic models of emotion regulation can be translated to understand changes in emotion across the life span and in clinical disorders.

Keywords: amygdala; cognitive control; emotion; emotion regulation; prefrontal cortex

... Thy fate is the common fate of all,
Into each life some rain must fall. . .

Henry Wadsworth Longfellow
The Rainy Day (1842)

... 'Every cloud,' says the proverb, 'has a silver lining.'

P. T. Barnum
Struggles and Triumphs (1869)

It might be said that emotions are the weather of our lives. Some days, we experience the blue skies of happiness and the sunshine of joy. Other days, we are drenched by the rain clouds of sadness or buffeted by the hot winds of anger. How we respond adaptively to our emotional weather patterns—finding the silver lining in every dark cloud—has important consequences for our physical and mental well-being.^{1–7}

Although we cannot control the weather outside, we are capable of using myriad emotion regulation strategies to take control of our internal climates.⁸

Such strategies allow us to wholly or partially alter the nature, magnitude, and duration of our emotional responses, including initiating new ones. In recent years, great strides have been taken in using neuroscience techniques to understand the mechanisms underlying emotion regulation. In humans, this research has primarily used functional imaging to examine our ability to control affective responses using cognitive strategies. The overarching goals of this paper are to review the progress made by such research, synthesize from it conclusions that suggest expansion on and elaborations of a model of the cognitive control of emotion (MCCE), and show how the model can make sense of a wide range of emotion regulatory abilities and allied phenomena.

Toward these ends, the remainder of the paper is divided into three parts. In the first, we outline a basic MCCE whose core elements have been described previously.^{9,10} In the second section, we review current imaging research suggesting ways in which the model can evolve to integrate new findings on the

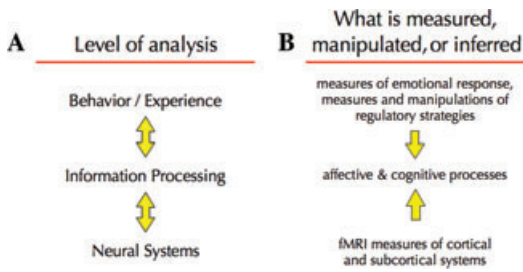


Figure 1. A multilevel approach to building a model of emotion regulation. (A) In cognitive, affective, and social neuroscience research, we seek to describe phenomena in terms of relationships among three levels of analysis: experience and behavior, psychological processes, and neural systems. The bidirectional arrows between levels indicate that the relationships among them are bidirectional. (B) Through measurement and/or experimental manipulation, neuroimaging research on emotion regulation can observe phenomena at the behavioral level and the neural level and use these observations to infer the nature of the intervening cognitive and/or affective processes. The direction of the arrows from the behavioral and neural levels toward the process level indicates the direction of causal inference (i.e., we can't observe the operation of these processes directly, but infer their operation based on behavioral and neural observations).

brain bases of emotion regulation as well as be applied to account for other related phenomena, such as affective learning, affect-based decision making, and affective expectancies. Throughout these first two sections we focus primarily on one strategy in particular—known as reappraisal—because it has received the bulk of empirical attention. In the third and last section, we summarize and consider directions for future basic and translational research.

A model of the cognitive control of emotion

Any model of emotion regulation (or any other phenomenon) is predicated on assumptions about how different levels of analysis fit together. Our assumptions follow those now commonplace in cognitive, affective, and social neuroscience in which researchers seek to describe phenomena in terms of the relationships among three levels of analysis: behavior/experience, process, and neural systems (Refs. 11–13; Fig. 1A). Neuroimaging research on emotion and its regulation can observe phenomena at the behavioral level (e.g., measures of emotional response and the specific regulatory strategies one might employ) and the neural level (e.g., fMRI measures of brain activity) and use these observations to infer the nature of the intervening cognitive and/or affective processes (Fig. 1B).

With this in mind, our review of current research will sometimes be organized in terms of phenomena described at the level of behavior, including regulatory goals, tactics, and target stimuli. In other cases it will be organized in terms of issues concerning the neural-level pathways on which the field has begun to make progress. Taken together, the data reviewed in each section constrains and influences our MCCE (see Fig. 2).

To understand how emotion regulation works, we must first have an idea of how emotions are generated. As such, our model has two main parts—descriptions of the mechanisms supporting emotion generation on the one hand and the mechanisms supporting emotion regulation on the other. For the sake of simplicity, we present the psychological and neural systems involved in the generation and regulation of emotion as being distinct, yet it should be noted that there is evidence to suggest that the underlying psychological¹⁴ and neural mechanisms^{15,16} are at least partially overlapping. Indeed, elsewhere we have noted that the distinction between emotion generation and regulation is blurry at best (e.g., Ref. 16), and which term one uses may reflect their usefulness for addressing a particular question more than hard and fast differences in their mechanisms. Here, we treat them separately to make points about the ways in which putative control and affect-triggering systems interact.

Mechanisms of emotion generation

Our account of how emotions are generated is multileveled¹² in its description of both the processes and the neural systems that give rise to emotional responses.

Processes involved in generating emotion

The black time line at the bottom of Figure 2A shows a simple model of four steps involved in generating emotional responses.¹⁷ In the first step, a stimulus is perceived in its current situational context. The stimulus could be an internal thought, feeling, or sensation, or any number of external cues, ranging from a facial expression or gesture to an action or event. At the second stage, one attends to some of these stimuli or their attributes. Whatever is in the focus of attention is passed along to subsequent emotion generative stages, whereas ignored or unattended stimuli may be either

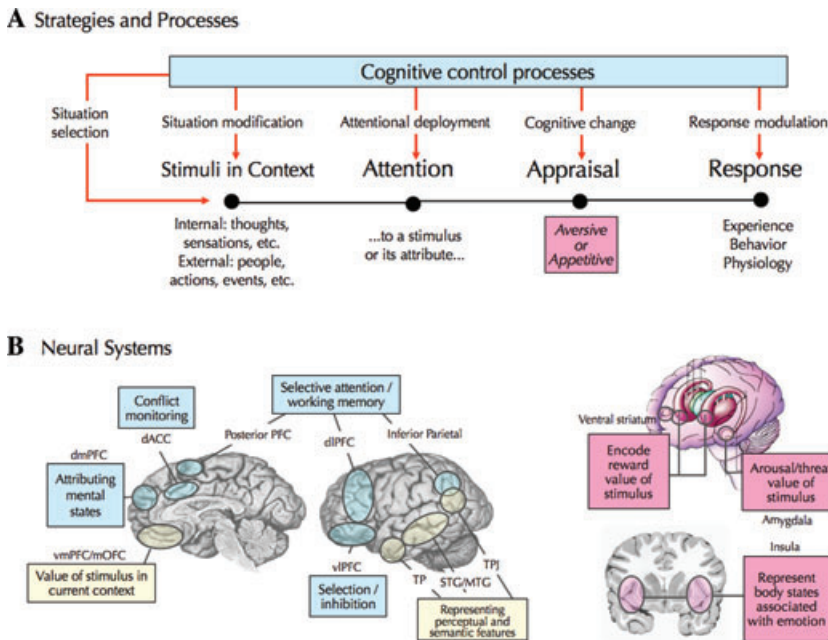


Figure 2. A model of the cognitive control of emotion (MCE). (A) Diagram of the processing steps involved in generating an emotion and the ways in which cognitive control processes (blue box) might be used to regulate them. As described in the text, the effects of different emotion regulation strategies (the red arrows descending from the cognitive control processes box) can be understood in terms of the stages of the emotion generation sequence that they influence. The pink box seen at the appraisal stage is meant to indicate that neural systems involved in generating emotion support this process. (B) Neural systems involved in using cognitive strategies, such as reappraisal, to regulate emotion (left, blue boxes), systems involved in generating those responses (left, pink boxes), and systems with an undefined or intermediary role in reappraisal (left, yellow boxes).

excluded from these stages or receive diminished subsequent processing. The third stage involves appraising the significance of stimuli in terms of their relevance to one’s current goals, wants, or needs. This is the stage focused on by appraisal theories of emotion, which describe the structure of different appraisals that lead to positive versus negative reactions in general and to specific types of emotional responses in particular.¹⁸ Because the current neuroscience literature suggests that there may not be specific neural systems for different discrete emotions,^{19,20} for present purposes, we simply distinguish between basic positive/appetitive versus negative/aversive appraisals that have been reliably associated with specific neural systems that are described below. Finally, the fourth stage involves translating these appraisals into changes in experience, emotion-expressive behavior, and autonomic physiology. Although these three indicators of emotional response do not always correlate with one another for reasons that are not perfectly understood,²¹ as noted below, emotion regulation

strategies can affect changes in some or all of them, depending on the strategy.

Neural systems involved in generating emotion

Reviews and meta-analyses of functional imaging studies^{19,20} indicate that a number of cortical and subcortical brain systems may play key roles in the appraisal and/or response stages of emotion generation. For present purposes, we focus on the four that have been most frequently discussed in studies of reappraisal in particular, and emotion regulation strategies more generally (see Fig. 2B; for examples of other emotion systems that may be modulated by emotion regulation, see Refs. 9 and 22).

The first is the amygdala, which is involved in the perception and encoding of stimuli relevant to current or chronic affective goals,^{23,24} ranging from rewards or punishments to facial expressions of emotion to aversive or pleasant images and films.^{25–27} Although the amygdala generally is sensitive to detecting and triggering responses to arousing

stimuli,²⁸ it exhibits a bias toward detecting cues signaling potential threats, like expressions of fear.^{29–31}

The second is the ventral striatum, which is involved in learning which cues (ranging from social signals, like smiling faces, to actions, to abstract objects) predict rewarding or reinforcing outcomes.^{32–34}

The third is the ventromedial prefrontal cortex (vmPFC), which integrates affective valuations of specific stimuli made by the amygdala and ventral striatum with inputs from other regions, including medial temporal lobe systems that provide historical information about prior encounters with the stimuli as well as inputs from brainstem motivational and prefrontal control centers that provide information about current behavioral goals.^{35–43} As such, vmPFC tracks the positive or negative valuation of stimuli in a context and goal-dependent manner.^{41,44–46} Examples of this include the finding that vmPFC activity to an image of a healthy but not tasty food depends on whether one has the goal to eat healthily,⁴⁷ and the findings that vmPFC lesions lead to context-inappropriate affective responses in both humans and animals.^{39,48,49}

The fourth brain system is the insula, which is thought to represent a viscerotopic map of ascending viscerosensory inputs from the body⁵⁰ and has been implicated in negative affective experience in general.^{51,52} There appears to be posterior–anterior functional gradient in the insula with posterior regions associated with primary representations of sensations from the body and anterior regions associated with interoceptive awareness of the body and in motivational and affective states, like disgust, that have a strong visceral component.^{51,53–56}

Mechanisms of emotion regulation

With an understanding of how emotions are generated in the first place, we can turn to an account of the processes and neural systems involved in regulating them.

Processes involved in emotion regulation

Although many behaviors can change our emotions, often these effects are unintended or incidental (e.g., your mood improves because you happen to have lunch with a friend) and as such are not considered to be examples of emotion regulation, *per se*. Instead, emotion regulation entails the modification of ongoing—or the initiation of new—emotional

responses through the active engagement of regulatory processes. That said, we can further distinguish between cases where emotion regulation is guided by regulatory goals that are implicit or outside awareness (e.g., Ref. 57) as compared with explicit and accessible to awareness. Although both are interesting and important, no neuroscience research has addressed the former case and a great deal has addressed the latter case. Therefore, we focus here on the deliberate deployment of an emotion regulation strategy in the service of explicit goals to change one's emotions. To understand how such explicit emotion regulation strategies work, it is useful to distinguish among five classes of strategies whose effects on emotion can be understood in terms of the stage of the emotion generation sequence on which they have an impact.⁵⁸

It is important to note that the distinctions made below originally were based on behavioral analyses of the aspects of emotional responses *targeted* by different strategies.⁵⁸ As such, this analysis was agnostic to the specific nature of the regulatory processes supporting each strategy, but tacitly assumed that all strategies drew upon some combination of cognitive control processes (designated by the blue box in Fig. 2A). In this regard, functional imaging has made a substantial contribution to our understanding of how emotion regulation works because it provides insight into the nature of the control processes supporting emotion regulation that is not obtainable from behavioral data alone.⁹

As illustrated by the top portion of Figure 2A, the first two strategies involve changing the nature of the stimulus inputs to the emotion generation cycle. In *situation selection*, you keep yourself away from stimuli that elicit unwanted emotions and put yourself in the presence of stimuli that elicit desired emotions. An example is staying away from a party where an old flame will be present if you don't want to feel pangs of sadness for having been dumped by her. *Situation modification* is when you find yourself in the presence of a stimulus that elicits an unwanted emotion and change something about the situation to alter its impact on you. In the old flame example, you might leave a party at which she is unexpectedly present or leave the room in which she is having a conversation. Although these two strategies are undoubtedly effective (e.g., Ref. 59), they can be difficult to study neurally and have received little

attention in imaging or using other neuroscience techniques (see below).

The remaining three strategies are all amenable to, and have been studied, using imaging, albeit to varying degrees. *Attentional deployment* controls what stimuli are gated into, or out of, the emotion generation process. The two most commonly studied exemplars¹⁰ are *selective attention*, which involves shifting the focus of attention toward or away from stimuli or their attributes, and *distraction*, which involves limiting attention to an external stimulus by focusing internally on information maintained in working memory. These types of strategies differ from situation selection in that they do not involve physically altering one's proximity or relationship to an emotional stimulus, but rather they manipulate attention so as to alter one's emotional response. *Cognitive change* involves changing the way one appraises the meaning of a stimulus. It is one of the most cognitively complex strategies insofar as it draws on any of a number of different higher cognitive processes to support changes in stimulus meaning, including language and memory, as well processes that also support other strategies, such as attention and response selection. The most commonly studied exemplar is *reappraisal*, which involves reinterpreting the meaning of a stimulus, including one's personal connection to it, to change one's emotional response. Finally, *response modulation* strategies target the systems for emotion-expressive behavior. The most commonly studied exemplar is expressive suppression,⁶⁰ which entails keeping one's face still so that observers would not know the emotion you are experiencing.

A great deal of behavioral and psychophysiological research has been devoted to comparing and contrasting the behavioral consequences of deploying each of these strategies. For example, it's known that attentional deployment and reappraisal can have downstream effects on various components of an emotional response because they target the early stages of the emotion generation sequence.^{60–65} By contrast, expressive suppression has an impact on only the behaviors it targets at the final response stage of emotion generation; when keeping your face still, emotional experience may subtly diminish,^{66,67} if at all, and your physiological arousal will increase from the effort.⁶⁰ There is also evidence that strategies differ in their long-term effects. For example, reappraisal, but not distraction, has been shown to

have long-lasting effects on one's tendency to have an emotional response to a stimulus,⁶⁵ presumably because only reappraisal involves an active change in how one represents the affective meaning of that stimulus.

Neural systems involved in emotion regulation

As foreshadowed previously, the use of functional imaging has provided insight into the nature of the control systems that support regulatory strategies as well as the affect systems that these strategies modulate to change an emotional response. This section discusses core conclusions that can be drawn from reappraisal studies and a model of emotion regulation that can be derived from it.

Reappraisal as a paradigm case. Reappraisal is an appropriate starting point for developing a MCCE for three reasons. First, because reappraisal is among the most cognitively complex strategies, a model of emotion regulation derived from reappraisal work may be generally applicable to other strategies and phenomena that typically will be cognitively simpler. Second, the majority of studies to date have focused on reappraisal because (i) it can be studied easily in an imaging environment and (ii) because it is *the* strategy referenced by countless aphorisms that advise us, “[to] look on the bright side. . .”; “[to] turn a sow's ear into a silk purse. . .”; “When life gives you lemons, make lemonade,” and, “[that] every dark cloud has a silver lining.” Third, in contrast to other areas of emotion regulation research (reviewed below) reappraisal studies tend to be more methodologically and conceptually similar to one another and therefore provide a stronger base for mechanistic inferences. With these considerations in mind, we now describe five key insights into the brain mechanisms supporting emotion regulation that have been derived from studies of reappraisal.⁹

Basic control system–affect system relationships.

When the first fMRI studies of reappraisal were published approximately 10 years ago, there were no imaging studies of any form of emotion regulation. To develop hypotheses about how reappraisal might work, an analogy was drawn between the use of cognition to control emotion and the use of cognition to control memory, attention, and other thought processes.⁴³ The simple idea was that prefrontal and cingulate systems would support control processes that modulate activity in posterior and subcortical

systems that generate emotional responses.¹⁰ A decade and over 50 imaging studies later, this initial hypothesis has been strongly supported.

Figure 2B schematically illustrates the brain systems shown by current research to be involved in the cognitive control of emotion via reappraisal. As such, Figure 2B diagrams the core elements of the MCCE. Three types of neural systems are primarily involved in generating and applying reappraisals.¹⁰ First, dorsolateral and posterior prefrontal cortex, along with inferior parietal regions generally implicated in selective attention and working memory, may be used to direct attention to reappraisal-relevant stimulus features and hold in mind reappraisal goals as well as the content of one's reappraisal.^{68–70} Second, dorsal regions of the anterior cingulate cortex implicated in performance monitoring may help track the extent to which one's current reappraisals are changing emotional responses in the intended way.⁷¹ Third, regions of ventrolateral prefrontal cortex implicated in selecting goal-appropriate (and inhibiting goal-inappropriate) responses and information from semantic memory may be used to deliberately select a new stimulus-appropriate reappraisal in favor of one's initial prepotent appraisal of that stimulus.^{72,73} Finally, to the extent that one's reappraisal involves focusing on and interpreting or reinterpreting one's own emotional states—or those of others—dorsomedial prefrontal regions implicated in attributing mental states also may be active.^{74,75}

With respect to the emotion-related regions that are modulated by reappraisal, the four regions described earlier in the section on neural systems for emotion generation all have been implicated—albeit to differing extents. Far and away, the most commonly modulated region is the amygdala, followed by the ventral striatum. The insula and the vmPFC are the least commonly modulated regions^{9,10} (although see the section on pathways below for a potential role of vmPFC in reappraisal as a modulator).

Although we will discuss the significance of the differential modulation of these regions in more detail later (see later sections on valence specificity and pathways), for now we can highlight the consistency with which they have been observed. Figure 3 plots peak activation foci for 43 studies (see Table 1) of reappraisal in healthy individuals as a function of reappraisal goals (panel A), reappraisal

tactics (panel B), and the valence of the emotion being regulated (panel C). Ignoring these distinctions for a moment, one can see that the control system–affect system relationships shown in Figure 2B and described previously have been observed reliably across numerous studies.

Moving beyond the basic model

With the consistency of the core control–affect system relationship as a foundation, we are now in a position to consider how this basic model—first proposed in 2002⁴³ and elaborated in 2005¹⁰—has evolved. Below we discuss first new conclusions that can be drawn about the model from recent studies of reappraisal. In this section, we pay special attention to two emerging features of the model: (i) the potential intermediary role of semantic/perceptual systems in reappraisal, and (ii) pathways linking control and affect systems. Next, we discuss the way in which the model can be applied to understanding regulatory strategies other than reappraisal as well as various allied phenomena involving control–affect system interactions.

Integrating new research on reappraisal

Recent research provides new insight into the distribution of emotion regulation–related activation foci as a function of reappraisal goals (i.e., the outcome one hopes to achieve by regulating, for example, increasing or decreasing an emotional experience), tactics (i.e., the specific subtype of reappraisal one implements), and the emotional valence of stimuli (i.e., whether the stimulus evokes a positive or negative emotional response). Here, we consider the implications of this work for the evolving MCCE.

Goal specificity

Arguably, the most common goal when using reappraisal is to decrease negative emotion, as when we attempt to make ourselves feel better about a disappointing paper rejection, an argument, and the like. It is not surprising then that this goal has been the focus of the majority of reappraisal studies (see Table 1). This is not the only goal that guides reappraisal, however. In some cases, as when we worry, ruminate, or make ourselves more anxious or fearful by elaborating on the meaning of unpleasant events, we are using reappraisal in service of the goal to increase emotion. A small, but growing, number of studies have examined this reappraisal goal as well.

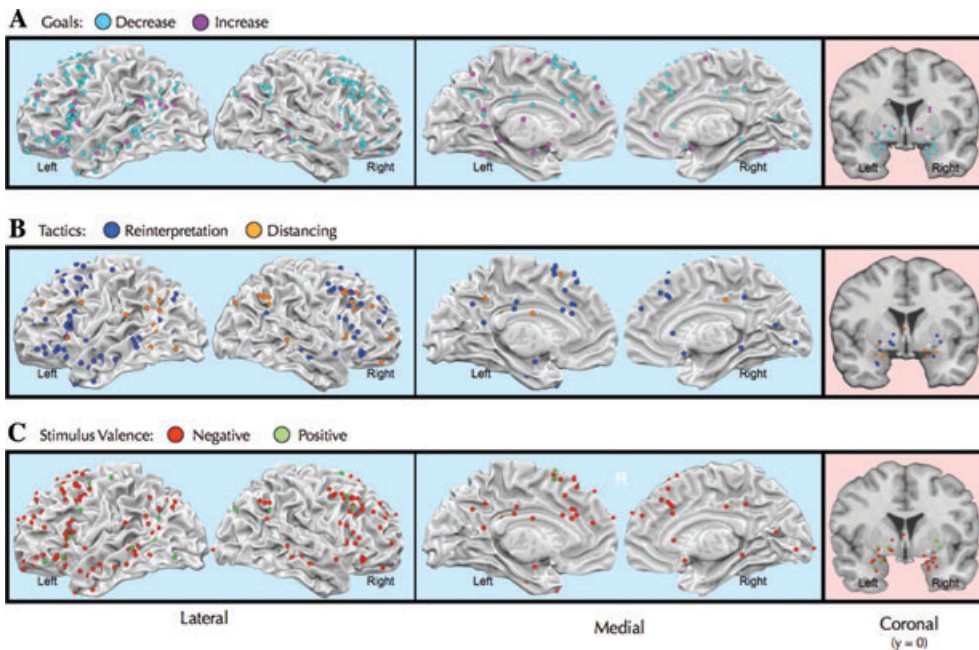


Figure 3. Plots of activation foci from the 43 studies of reappraisal described in the text and Table 1. (A) Plots of foci as a function of the goals to decrease or increase emotion. (B) Plots of foci as a function of the specific reappraisal tactics used—either reinterpreting the meaning of events depicted in stimuli or actively changing one’s psychological distance from them. (C) Plots of foci as a function of the valence of the stimuli eliciting the emotions that participants attempted to regulate. Blue boxes illustrate regions that are purported to support reappraisal (increase > look and decrease > look contrasts). Pink boxes illustrate regions that are purported to be modulated by reappraisal (look > decrease and increase > look contrasts; for clarity, only foci falling within the boundaries of the amygdala and striatum are shown).

Figure 3A plots peak activation foci for reappraisal studies of healthy individuals as a function of decrease versus increase goals. Perusal of this figure highlights three findings. First, whereas both increase and decrease goals recruit left prefrontal regions, decrease goals recruit right prefrontal regions to a much greater extent than do increase goals. There are two interpretations of this finding. First, it may be attributable to the fact that decreasing an emotional response is more difficult than increasing one, and therefore may require additional cognitive control resources.⁷⁶ Second, decreasing—but not increasing—an emotional response requires inhibiting or limiting the expression of a prepotent appraisal of a stimulus (e.g., as negative) in favor of selecting an alternative reappraisal (e.g., as neutral or even positive). Research shows that right dorsal—and especially ventrolateral—prefrontal cortex is involved in the selection and/or inhibition of various kinds of responses.^{77–80}

Second, there is some evidence that increase goals differentially involve anterior portions of dorsomedial prefrontal cortex (dmPFC). Of the 12 studies directly comparing increasing emotion with a control condition in which participants respond naturally, six show increases in anterior dmPFC.^{16,76,81–83} Of the six studies that did not, most showed activation in neighboring areas (such as anterior cingulate cortex).^{62,84–88} Given the role of dmPFC in making judgments about mental states^{75,89,90} and that the majority of reappraisal studies use photographs of people as stimuli (see Table 1), it is likely that these regions support attention to and elaboration of emotional states, intentions, and outcomes of the individuals depicted in these photos.

Third, whereas increase and decrease goals both seem to modulate the striatum (including both the caudate and putamen), they may differ in the way they modulate the amygdala. On the one hand,

Table 1. Neuroimaging studies of reappraisal in healthy individuals^a

Study	Participants	Design			Stimulus type	Timing of reapp cue	Amygdala
		Goal	Valence	Tactic			
Beauregard <i>et al.</i> ²⁰⁴	HYA	Dec	Pos	Dist	Videos	Early*	No
Domes <i>et al.</i> ⁸¹	HYA	Both	Neg	Both	Photos	Late	Yes
Eippert <i>et al.</i> ⁸⁴	HYA	Both	Neg	Both	Photos	Late	Yes
Erk <i>et al.</i> ¹⁶⁸	HC	Dec	Neg	Dist	Photos	Early	Yes
Goldin <i>et al.</i> ⁶⁶	HYA	Dec	Neg	Reint	Videos	Early	Yes
Harenski <i>et al.</i> ²⁰⁵	HYA	Dec	Neg	Both	Photos	Early	Yes
Hayes <i>et al.</i> ¹³⁹	HYA	Dec	Neg	Reint	Photos	Late	Yes
Herwig <i>et al.</i> ²⁰⁶	HYA	Dec	Both	Reint	Anticipate photos	Early	Yes
Hollmann <i>et al.</i> ²⁰⁷	HYA	Dec	Pos (food)	Reint	Photos	Early	No
Ichikawa <i>et al.</i> ⁸²	HYA	Both	Neg (errors)	Reint	Task errors	Early*	No
Kanske <i>et al.</i> ¹²⁷	HYA	Dec	Both	Both	Photos	Late	Yes
Kim <i>et al.</i> ⁹⁵	HYA	Dec	Both	Reint	Photos	Early	Inc pos only
Kober <i>et al.</i> ¹¹⁸	HYA smokers and non-smokers	Dec	Pos (food/cigs)	Reint	Photos	Early	Yes
Koenigsburg <i>et al.</i> ¹⁹⁹	HYA	Dec	Neg	Dist	Photos	Early	Yes
Krendl <i>et al.</i> ²⁰⁸	HYA	Dec	Neg	Unclear	Photos	Early	Yes
Kross <i>et al.</i> ¹¹²	HYA	Dec	Neg	Reint	Memories	Late	No
Lang <i>et al.</i> ⁸³	HC	Both	Neg	Dist	Scripts	Early	Inc only
Levesque <i>et al.</i> ²⁰⁹	HYA	Dec	Neg	Dist	Videos	Early*	No
Mak <i>et al.</i> ²¹⁰	HYA	Dec	Both	Unclear	Photos	Early*	No
McRae <i>et al.</i> ²¹¹	HYA	Dec	Neg	Reint	Photos	Early	Yes
McRae <i>et al.</i> ⁶¹	HYA	Dec	Neg	Reint	Photos	Early	Yes
McRae <i>et al.</i> ¹⁸⁰	Healthy aged 10–22	Dec	Neg	Reint	Photos	Early	No
McRae <i>et al.</i> ¹⁵	HYA	Dec	Neg	Reint	Photos of faces	Early	Yes
Modinos <i>et al.</i> ²¹²	HYA	Dec	Neg	Reint	Photos	Late	Yes
New <i>et al.</i> ⁸⁵	HC	Both	Neg	Reint	Photos	Late	Yes
Ochsner <i>et al.</i> ⁴³	HYA	Dec	Neg	Reint	Photos	Late	Yes
Ochsner <i>et al.</i> ⁷⁶	HYA	Both	Neg	Both	Photos	Early	Yes
Ochsner <i>et al.</i> ¹⁶	HYA	Inc	Neg	Both	Photos	Early	Yes
Ohira <i>et al.</i> ¹¹⁵	HYA	Dec	Both	Unclear	Photos	Early*	Yes
Opitz <i>et al.</i> ⁹⁶	HYA and HOA	Both	Neg	Reint	Photos	Late	No
Phan <i>et al.</i> ¹¹⁴	HYA	Dec	Neg	Reint	Photos	Early*	Yes
Pitskel <i>et al.</i> ⁸⁶	Healthy aged 7–17	Both	Neg	Reint	Photos	Early	Yes

Continued

Table 1. Continued

Study	Participants	Goal	Valence	Design			Amygdala
				Tactic	Stimulus type	Timing of reapp cue	
Schardt <i>et al.</i> ²¹³	HYA	Dec	Neg	Dist	Photos	Early	Yes
Schulze <i>et al.</i> ⁸⁷	HC	Both	Neg	Both	Photos	Late	No
Staudinger <i>et al.</i> ²¹⁴	HYA	Dec	Pos	Dist	Reward	Early*	No
Staudinger <i>et al.</i> ²¹⁵	HYA	Dec	Pos	Dist	Anticipate reward	Early*	No
Urry <i>et al.</i> ⁹⁴	HOA	Both	Neg	Reint	Photos	Late	Inc only
Urry <i>et al.</i> ²¹⁶	HOA	Both	Neg	Reint	Photos	Late	Yes
van Reekum <i>et al.</i> ⁸⁸	HOA	Both	Neg	Reint	Photos	Late	Dec only
Vrticka <i>et al.</i> ²¹⁷	HYA	Dec	Both	Reint	Photos	Early*	Yes
Wager <i>et al.</i> ¹¹⁷	HYA	Dec	Neg	Reint	Photos	Early	Yes
Walter <i>et al.</i> ²¹⁸	HYA	Dec	Neg	Dist	Photos	Early	Yes
Wincoff <i>et al.</i> ¹⁹⁵	HYA and HOA	Dec	Both	Dist	Photos	Late	Yes

*Studies are ordered first by year and second by alphabetical order.

Only studies that reported contrasts (i.e., not only functional connectivity or correlational analyses) for psychologically healthy individuals are included here. If a study included a patient sample but still reported results for its healthy adult controls separately, it was included.

HYA, healthy young adults, typically 18–30 yrs old; HOA, healthy older adult, typically aged 60 years or older; HC, healthy adult control participants matched to patients; For design, Goal, goal pursued by participants to increase or decrease emotional responses; Valence, Positive or negatively valenced emotional stimuli; Tactic, type of reappraisal used, distancing or reinterpreting; Stim Type, stimulus type; Timing of reapp cue, timing of instruction cue to reappraise relative to onset of stimulus, where early is just before stimulus onset and late is a few seconds after stimulus onset; Amygdala, whether modulation of amygdala was reported.

NOTE: All studies used event-related designs (different types of trials are presented in a randomized fashion so as to estimate responses on a trial-by-trial basis) except the nine studies designated by * in the “Timing of reapp cue” column, which indicates that they used a block design (trials are “blocked” by type, such that many of one type appear consecutively). Also, for the stimulus-type column, photo stimuli were drawn from the international affective picture system¹¹¹ unless otherwise specified. Goal: Dec, decrease; Inc, increase; Both, both increase and decrease conditions were used; Valence: Neg, negative; Pos, positive; Both, both positive and negative stimuli were used; Strategy: Both, both distancing and reinterpreting were used (this only applies to Ref. 76), or participants were given the choice of distancing or reappraising; Dist, become more or less psychologically distant; Reint, cognitively reinterpret; Unclear, unclear as to what tactic was instructed.

decrease goals reliably modulate the amygdala’s ventral (corresponding to the basal and lateral amygdala nuclei) and dorsal portions (corresponding to the central nucleus) as well as the sublentiform extended amygdala (SLEA^{30,91,92}) that lies between the amygdala and the striatum. On the other hand, increase goals may modulate only the dorsal amygdala/SLEA. One speculative interpretation of these data is that decrease goals influence perceptual and semantic inputs to the amygdala, which come through the basolateral complex, whereas in-

crease goals influence the outputs of the amygdala, which flow from the central nucleus.^{43,76} This hypothesis would fit with anatomical data showing that the basolateral complex has reciprocal connections with ventrolateral PFC as well as temporal and parietal regions implicated in visuospatial and semantic representation, whereas the central nucleus receives inputs from medial prefrontal regions and sends outputs to autonomic centers that implement various components of an emotional response.⁹³

The major caveat for all of these conclusions, however, is that very few studies have examined increase goals, and as a consequence, conclusions about the goal specificity of reappraisal-related activations must be considered tentative. That being said, the first study to directly compare increase and decrease goals within subjects obtained exactly the results described previously⁷⁶—both increasing and decreasing negative affect recruited left vLPFC and dlPFC and modulated the dorsal amygdala/SLEA (increasing affect increased amygdala activity, whereas decreasing negative affect decreased activity), yet it was also revealed that increasing negative affect recruited the dmPFC to a greater degree than did decreasing negative affect and decreasing negative affect recruited right vLPFC and modulated ventral amygdala to a greater extent than did increasing negative affect. At least two-thirds of subsequent studies comparing these goals have obtained results that are generally consistent with them^{76,84,87,94–96} (other findings also have been reported, including increase versus decrease differences only in the amygdala,⁸¹ striatal modulation,^{76,88,94} and greater right PFC activation for increasing than decreasing⁸⁴).

Tactic specificity

In the military, a distinction is commonly made between strategy and tactics. Strategy is the overall means by which a goal (e.g., win the war) is to be achieved (e.g., divide and conquer). Tactics are the specific ways in which strategies are implemented in a given circumstance (e.g., a quick infantry advance, an airstrike, etc.). In the same way, one can distinguish between reappraisal as a strategy that involves changing the meaning of a stimulus and the tactics used to implement that strategy.⁹⁷

Two different reappraisal tactics have been studied with imaging.^{9,76} The first can be called *reinterpretation*, which involves changing one's interpretation of the elements of the situation or stimulus that elicits emotion. For example, if one is presented with a photo of a sick man in the hospital that elicits feelings of sadness, one might reinterpret this image in a way that decreases emotion by thinking about the man's hearty constitution and that he will be healthy and well in the future. To increase emotion, one might instead think about how the man is in a great deal of pain and may, in fact, get worse and even perish. The second can be called

distancing, which involves changing one's personal connection to, or psychological distance from, the stimulus that elicits emotion. In the example of the photo of the sick man, one might decrease emotion by viewing the image from the detached perspective of an objective, third person observer and/or imagining that the pictured event took place a long time ago or in a faraway location. One might increase emotion by instead imagining that one is experiencing pictured events in the present moment, from a first-person perspective, which enables you to smell, hear, and directly observe what is taking place.

As Table 1 shows, about twice as many studies have examined reinterpretation as have examined distancing, with a few allowing participants to engage in either tactic, and only a single study directly comparing them.⁷⁶ Figure 3B plots peak activation foci for reappraisal studies of healthy individuals as a function of reinterpretation versus distancing tactics.

This figure illustrates three conclusions that can be drawn about reappraisal tactics. First, reinterpretation seems to differentially call upon ventral lateral prefrontal regions implicated in response selection and inhibition.^{73,98,99} Presumably, this reflects the fact that reinterpretation requires that one must look up and select alternative meanings for stimuli from semantic memory to a greater extent than does distancing. Second, distancing seems to recruit parietal regions implicated in spatial attention and representation to a greater extent, including perspective taking and the sense of agency.^{100–103} This may reflect the fact that distancing involves changing the conceptual and spatiotemporal perspective from which stimuli are experienced. Third, in general, the regions involved in reinterpretation appear to be more strongly left lateralized in prefrontal and temporal cortices, whereas regions involved in distancing appear to be more strongly right lateralized in prefrontal cortex. These patterns may reflect the differential dependence of reinterpretation and distancing on linguistic and semantic processes as opposed to spatial and attentional processes, which generally show a left versus right hemisphere pattern of relative specialization.^{76,104}

Here again, however, because comparatively fewer studies have examined distancing, firm conclusions concerning the tactic specificity of reappraisal-related activations await further

research that directly test the conclusions drawn previously.

Valence specificity

On average, the impact of negative emotional experiences seems to be greater than the impact of positive emotional experiences, both in the short and long term.¹⁰⁵ Indeed, problems with regulating negative emotion are more often a hallmark of clinical disorders than are problems with regulating positive emotion.¹⁰⁶ As such, it is not surprising that Table 1 shows that the number of reappraisal studies examining negative emotion outnumber those examining positive emotion more than three to one.

That said, two conclusions can be drawn from examining Figure 3C, which plots peak activation foci for reappraisal studies of healthy individuals as a function of the negative versus positive valence of stimuli (and the emotions they presumably elicit). First, whereas reappraisal of both negative and positive stimuli depends upon left-hemisphere regions, reappraising negative stimuli depends on right hemisphere regions as well. These findings might reflect the fact that, to date, the majority of studies of negative emotion involve decrease goals. As noted earlier, decrease goals may require more cognitive resources than increase goals, including placing greater demands on selection/inhibitory functions associated with right vLPFC.^{107–109} An alternative explanation is that positive and negative emotions generally involve approach versus avoidance motivations, which have been associated with the left versus right prefrontal cortex. This interpretation seems less likely, however, given that this motivation-related prefrontal asymmetry is commonly observed in EEG¹¹⁰ but not in fMRI studies.¹⁹

Second, it's apparent that reappraising negative stimuli typically modulates activity in the amygdala and less commonly activity in the striatum. By contrast, the handful of studies examining reappraisal of positive stimuli more commonly show modulation of the striatum, including the ventral portions associated with reward and reinforcement learning.^{33,34}

These conclusions are again tentative, however, because so few studies have examined reappraisal of positive stimuli and, in general, studies of reappraisal have focused overwhelmingly on decrease rather than increase goals. As a consequence, it is not yet clear whether the patterns noted previously

are attributable to the pursuit of decrease versus increase goals, the use of negative versus positive stimuli, or both.

Stimulus specificity

To date, 33 out of the 43 reappraisal studies shown in Table 1 have used photographic stimuli pulled from the International Affective Picture System (IAPS). These stimuli have been shown to reliably elicit experiential, physiological, and facial expressive components of an emotional response in a valence-specific manner.¹¹¹ As such, they provide a straightforward means of eliciting affective reactions in the scanner environment.

That said, the emotions elicited by such stimuli may or may not generalize to other contexts. For example, IAPS photos are selected so as to be normatively positive or negative.¹¹¹ Although this is suitable for many experimental agendas, other stimuli may be appropriate if one wants to examine the ability to reappraise specific emotions, the emotions elicited by idiosyncratically self-relevant autobiographical experiences,^{112,113} and so on.

With these considerations in mind, small numbers of studies have examined the ability to reappraise the specific emotions elicited by sad, sexual, or disgusting videos, scripts that elicit particular emotions, the recollection of autobiographical memories, anticipation of reward or shock, or the commission of an error (see Table 1). Because so few studies have used each of these stimuli, it is not useful at present to plot activation foci for them or to attempt to draw conclusions about how they might differ as a function of stimulus type. It remains for future research to directly address the question of how the nature of the stimulus *per se*, as opposed to the kind of emotion elicited, influences the neural systems involved in reappraisal.

Pathways linking control and affect systems

Studies of reappraisal—and more generally studies of any form of emotion regulation—implicitly or explicitly assume that prefrontal regions modulate emotional responses via their impact on affect systems like the amygdala and ventral striatum. Given the prevalence of this assumption, it is somewhat surprising that it has seldom been put to a direct test. To be sure, a number of studies have shown correlations between prefrontal and amygdala

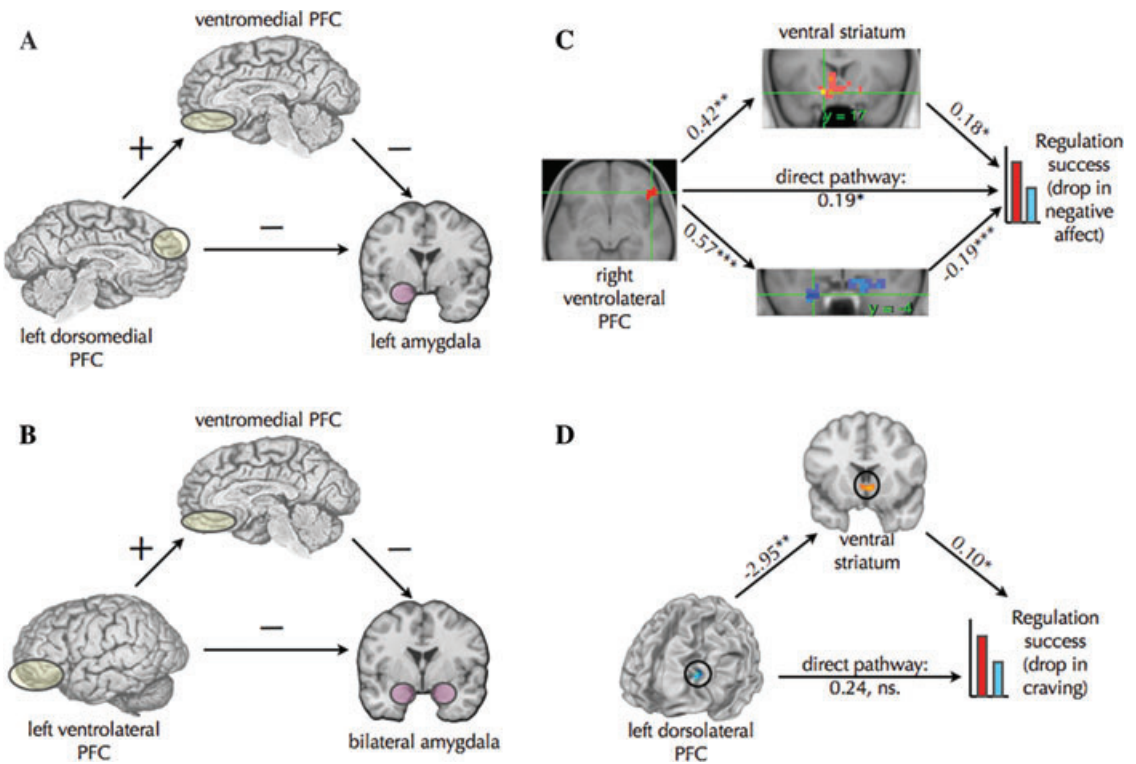


Figure 4. Two kinds of mediation pathways involved in reappraisal. (A) and (B) show pathways identified in two studies of the downregulation of negative emotion whereby dorsomedial or ventrolateral prefrontal regions diminish amygdala responses via their impact on ventromedial prefrontal cortex. These studies did not report weights for the mediation paths between regions or test for full versus partial mediation. (C) and (D) show pathways identified in two studies of the downregulation of negative or positive emotion whereby ventrolateral or dorsolateral prefrontal regions diminish self-reports of negative affect or craving via their impact on the amygdala or ventral striatum, respectively.

activity^{76,95,114} or correlations between some measure of emotional response (typically self-report) and either prefrontal^{76,114,115} or amygdala activity.^{76,94,116} Only four studies, however, have directly tested the mediation model implied by the hypothesis that control systems influence emotional response by influencing activity in affect systems (see Fig. 4).

The first two studies to use mediation examined the use of reappraisal to diminish responses to negative photos.^{94,116} Although both studies used amygdala reactivity as their measure of emotional response, neither reported a main effect of reappraisal on diminishing amygdala activity. Motivated by known connections between the amygdala and vmPFC, both studies looked for and found that individual differences in amygdala response were correlated inversely with responses of vmPFC. Mediation analyses showed that vmPFC mediated a

relationship between either left dmPFC⁹⁴ or left vlPFC¹¹⁶ and the amygdala (Fig. 4A and B) such that activity in these prefrontal regions was positively related with vmPFC activity, which, in turn, was negatively related to amygdala responses. There are, however, at least two qualifiers in interpreting these results. First, the study⁹⁴ identifying the left dmPFC region did so in an increase > attend (i.e., a no regulation baseline) > decrease contrast, meaning that it generally is *less* active when decreasing negative emotion than when responding naturally in a baseline “attend” condition. This suggests that to the extent one shows *less deactivation* when decreasing (relative to baseline), one will show greater activity in vmPFC, and, in turn, less amygdala response. It is not immediately clear how to interpret the reduced degree of dmPFC deactivation in this context. Second, the study¹¹⁶ identifying the vlPFC→vmPFC→amygdala pathway

collapsed across activity in both amygdalae that was extracted from structural ROIs. As such, it is not clear whether the prefrontal effects were stronger for one amygdala or the other. That said, when taken together, these two studies suggest that effective reappraisal involves PFC→vmPFC→amygdala pathways.

The second two studies used similar analytic approaches to study either the use of reappraisal to diminish responses to negative photos¹¹⁷ or for smokers, the use of reappraisal to diminish craving elicited by photographs of appetitive foods or cigarettes.¹¹⁸ The study of negative emotion¹¹⁷ showed that right vPFC activity predicted drops in self-reported negative emotion, and that this relationship was independently mediated by separate pathways through the amygdala and the ventral striatum (Fig. 4C). These two pathways were taken to reflect the use of reappraisal to minimize negative appraisals and enhance positive reappraisals, respectively (see also Table S1 from that paper, which shows left vPFC involvement as well). The study of craving¹¹⁸ showed that left dlPFC activity predicted drops in self-reported craving via modulation of activity in the ventral striatum (Fig. 4D). Together, these two studies suggest that effective reappraisal involves a pathway linking PFC→subcortex→emotion change, with the specific elements of the pathway depending on the nature of the stimulus and emotion involved.

Why are there differences between the results of these pairs of studies? On one hand, because different dependent measures of emotional response were used (amygdala response vs. self-reported emotion), it's possible that different reappraisal pathways will emerge depending on the type of response. On the other hand, it's also possible that differences in methodology may lead participants to reappraise differently, and, in turn, recruit different pathways for effective emotion regulation. Here, two differences between the pairs of studies may be relevant.

In the first pair of studies that identified the vmPFC-mediated pathway, both had participants that were up to 40 years older than the average participants in studies of reappraisal in young adults—aged 62–64 in one case⁹⁴ and 19–53 (average 33 years) years in the other.¹¹⁶ Participants in the second pair of studies were younger, as is the norm, averaging 22.3¹¹⁷ and 26.8¹¹⁸ years, respectively. Given findings that older adults may be impaired in some

kinds of reappraisal; the lateral PFC thins, whereas vmPFC thickens with age;¹¹⁹ and that even when not told to regulate, older adults can show greater connectivity between vmPFC and amygdala,¹²⁰ it is possible that the vmPFC plays a larger role in reappraisal for older compared with younger adults.

Second, the first pair of studies cued participants to reappraise approximately 4 seconds into an approximate 10-second presentation of an aversive photo (a *late cue*), whereas the second pair presented the cue to reappraise just before onset of aversive photos (an *early cue*). The early cue method is intended to provide participants with an opportunity to first have a naturalistic emotional response to an aversive photo before they begin to regulate and has been used in 14 studies (see Table 1). The late cue method models real-world situations where the goal to reappraise comes online just as one encounters an emotionally evocative stimulus and has been used in 29 studies (see Table 1).

Although the early cue method is analogous to real-world situations where the goal to regulate comes online only after one already is having an emotional response, there is a potential problem with trying to model this in the lab. During the initial free viewing of an aversive photo, participants may try out a few reappraisals just in case they are asked to subsequently reappraise on that trial. If this were the case, then we might expect one or both of two kinds of results.

One possibility is that the ability to detect an effect of reappraisal on amygdala responses would be diminished for late cue studies, either because the amygdala responded early and then habituated, or because once participants were asked to reappraise, the amygdala's response could have already decreased because the participants had begun generating/practicing potential reappraisals before the explicit instruction cue appeared. Although neither of the mediation studies in question showed whole-brain amygdala effects, and only one showed effects using ROIs, weak effects of reappraisal on the amygdala are probably related to other factors (like age—see above) given that roughly the same ratio (roughly 2/3 to 3/4) of studies using the late and early cue methods show reappraisal-related amygdala modulation, especially for studies using photos (see Table 1).

A second possibility is that the late versus early timing of reappraisal cues changes the nature of

one's reappraisals, even if, on average, they have similar effects on amygdala responding. For example, in late cues studies, if participants have had a chance to view stimuli for a few seconds and to think about potential reappraisals before being explicitly told to reappraise, then vmPFC recruitment could reflect decision processes about which of a set of prepared reappraisals they prefer and can best use for the stimulus at hand (see also section on decision making below).

To date, no imaging studies have compared late and early cues. But one behavioral/psychophysiological study comparing these cues found that the effects of increase goals on some physiological measures are greater for late than early cues, but that the effects of decrease goals were similar for each cue type. Future work could fruitfully illuminate these issues.

All this said, it is of course possible that both kinds of pathways are important and that a multistep vIPFC→vmPFC→amygdala/striatum→emotion response pathway may be observed in future studies. To date, however, no published studies have expressly tested for the existence of this complex pathway underlying reappraisal success.

The role of perceptual and semantic systems

A related issue is whether and how reappraisal involves modulation not just of systems involved in affective appraisal and response, but of systems involved in representing the perceptual and semantic properties of stimuli as well. As shown in Figure 3, activation of a number of these systems is often seen during reappraisal, including regions along the middle and superior temporal sulci involved in representing the visual properties of stimuli, including nonverbal social cues to emotion like movements of lips and eyes;^{121–123} temporal polar regions implicated in representing episodic and semantic emotion knowledge;¹²⁴ and regions near the temporal–parietal junction involved in representations of beliefs, including “false” beliefs of the sort one generates when considering alternative reappraisals of stimuli.^{125,126}

These data raise at least three questions. First, there is the question of when activation of these regions will be seen. Certainly, cognitive change strategies like reappraisal may involve these regions, given that it involves an active reworking of the

meaning of a stimulus. Other strategies that do not focus on meaning may not involve these regions, however. Consistent with this, two studies directly comparing reappraisal and distraction found that reappraisal differentially recruited all three of the temporal regions listed above.^{61,127} Along these lines, it is also likely that these regions will be more involved in regulating responses to visual stimuli given the role of the temporal lobe in the “ventral visual stream” for representing information about object identity^{128–130} (although this remains to be tested directly). As noted previously, there is not yet enough work using different kinds of stimuli to say whether reappraisal of stimuli in nonvisual modalities (e.g., somatosensory or auditory) may involve modulation of corresponding modality-specific regions (e.g., somatosensory or auditory cortices).

Second, if these regions are more active during reappraisal, there is the question as to why this is the case. Does greater activity here reflect increased attention to perceptual and semantic aspects of stimuli? Access to/retrieval of alternative “views” of reappraised stimuli? The process of actively restructuring one's (visual) mental image of a stimulus? All three interpretations are possible and could be tested in future work.

Third, there is the question of whether these temporal regions play a part in the regulation pathways described earlier—playing an intermediary role, for example—between prefrontal control systems and affective appraisal systems. This possibility was raised in early reappraisal studies (e.g., Ref. 43) where it was suggested that even though dorsolateral PFC regions do not have direct connections to subcortical regions like the amygdala, they may nevertheless modulate them via their impact on perceptual/semantic systems. In line with this view, PFC could change one's mental representation of a stimulus's meaning from the top down, and that representation of the reappraised stimulus would feed forward to the amygdala (and other structures that trigger affective responses). Because the amygdala now “sees” the reappraised stimulus, its response changes. Although plausible, this hypothesis has yet to be directly tested.

Summary

Extant data from functional imaging studies of reappraisal strongly support the MCCE depicted in

Figure 2B. Although many questions remain to be addressed about how specific control systems modulate specific affect systems as a function of reappraisal goals and tactics or various aspects of stimuli and the emotions they elicit, a core control–affect system dynamic is now well established.

Generalizability to other forms of regulation

Given the robustness of the MCCE (Fig. 2) in accounting for reappraisal, the question naturally arises as to whether this model can be generalized to account for other types of emotion regulation strategies.

As noted previously, the majority of functional imaging studies of emotion regulation have focused on reappraisal. That said, the other four main classes of emotion regulation strategies diagrammed in Figure 2A have been targeted by imaging studies to varying degrees. Here, we consider each in turn.

Situation selection and modification

The two situation-focused strategies, situation selection and situation modification, have received little attention thus far in human imaging research. As noted earlier, this is at least partially attributable to the difficulty of devising appropriate lab paradigms for studying them. The lone human imaging study of situation selection builds on the rodent literature on avoidance conditioning. In a typical task, a rat learns to perform an action that allows it to avoid presentation of an aversive stimulus (e.g., Refs. 131 and 132). In a human analogue of this procedure, Delgado *et al.* found that avoidance conditioning activates vIPFC and dlPFC control systems and modulates the amygdala.¹³³ These findings provide an initial suggestion that situation selection may call systems that maintain regulatory goals and select context-appropriate avoidance responses.

Attentional deployment

By contrast, studies of attentional deployment have been relatively common, second in number only to studies of reappraisal. One set of these studies has examined the use of selective attention to shift visual spatial attention away from an affectively valenced stimulus or stimulus attribute and toward a neutral one. Another set of these studies has examined the use of distraction to shift the focus of attention inward onto some internally maintained mental representation (e.g., a relevant working memory load, self-generated stimulus-irrelevant thoughts, a pleas-

ant mental image, and so on). As has been reviewed in detail elsewhere,^{10,134} interpreting the findings of both of these kinds of studies is clouded by three issues. First, almost all of the studies of selective attention, and many studies of distraction, use stimuli that do not elicit strong emotional responses, such as facial expressions of emotion. As such, these studies are concerned with the regulation of evaluative judgment or perception rather than affective responding, *per se*. Second, when highly arousing and affect-inducing stimuli are employed, they most often are stimuli that cause physical pain. Although responses to painful stimuli have a strong negatively valenced affective component, this component may itself have a distinct neural signature because of its recruitment of dedicated pain-specific neural pathways.^{135,136} As such, it is an empirical question whether the regulation of pain is similar to or different from the regulation of negative affective responses more generally. Third, these studies are highly heterogeneous, often employing very different stimuli and methods of controlling the focus and level of attention, without a clear metric for assessing the extent to which attention has or has not been paid to a given affective stimulus. Given these limitations, we refer the reader to other reviews of this literature,^{137,138} although noting that they are generally consistent with the model depicted in Figure 2B, insofar as activation of prefrontal systems and modulation of affect systems (like the amygdala) is often (but not always) reported.

Response modulation. Finally, both behavioral and imaging studies of *response modulation* have focused on expressive suppression, which is the ability to hide behavioral manifestations of emotion.⁶⁰ The two imaging studies of expressive suppression asked participants to suppress facial expressions of disgust elicited by a film clip.^{66,139} Both found that expressive suppression not only activated dorsolateral and ventrolateral PFC regions associated with maintaining goals, response selection, and inhibition,^{72,73,77} but also it increased activation of the insula, which is involved in triggering affective responses. Amygdala findings were more mixed, however, with one study reporting increases⁶⁶ and one decreases¹³⁹ in activity during suppression. Increases in insula and amygdala fit with psychophysiological studies, demonstrating that expressive suppression

boosts the autonomic component of emotional responding.⁶⁰

In total, the available literature on emotion regulation strategies other than reappraisal is in some cases limited and in other cases somewhat confusing, but in general supports the idea that all emotion regulation strategies involve interactions between cognitive control and affect regions. Future neuroimaging research must apply the same rigorous and thorough approach to these other strategies that has already been applied to reappraisal.

Generalizability to other related phenomena

Given the robustness of the MCCE (Fig. 2B) in accounting for multiple forms of regulation, a natural next question is whether it can be generally applied to other allied phenomena, such as affective/emotional learning, decision making, and expectancies. These phenomena are typically considered in separate literatures, but seem to involve related cognitive–affective dynamics. Although space limitations prohibit an in-depth discussion, here, we briefly examine the broad applicability of the model in each of these three cases.

Affective/emotional learning. At the outset of this paper we made a distinction between goal-directed forms of emotion regulation, which are the focus of this review, and other behaviors that may have regulatory effects on emotion despite lacking a specific goal to do so. There are a number of forms of affective or emotional learning that fit the latter description. One of the most common examples is extinction of a conditioned fear response. In the traditional fear conditioning paradigm,¹⁴⁰ an animal learns that an ostensibly neutral stimulus, such as a light (known as the conditioned stimulus or CS), predicts the occurrence of an intrinsically aversive stimulus, such as electric shock (known as the unconditioned stimulus or UCS). Over time, the repeated pairing of the light and shock lead the animal to respond to the light itself with an anticipatory fear response. Elegant animal studies have shown that fear conditioning depends on communication between input and output nuclei of the amygdala.^{140,141} Fear extinction involves the repeated presentation of the CS in the absence of the UCS.¹⁴² Over time, the organism learns that the CS no longer predicts shock, ceases to have its

anticipatory fear response, and fear is said to be extinguished. Importantly, extinction is known to involve the laying down of a new context-dependent memory.¹⁴³ In the current temporal context, the CS does not predict shock, whereas in the past temporal context it did. Rodent lesion studies have shown that whereas the initial acquisition of extinction requires only the amygdala, the ability to retain and express memory for extinction depends on vmPFC.¹⁴² In keeping with this finding, studies in humans have shown that both the magnitude of vmPFC activation and vmPFC thickness predict the speed of extinction.^{144–146}

In the present model, phenomena like extinction (or stimulus–reward reversal learning, which also depends upon vmPFC^{147,148}), are somewhat hybrid phenomena. On the one hand, they can be viewed as an example of emotion generation, insofar as one is learning to express a new emotional response to a given stimulus. On the other hand, they can be viewed as an implicit form of emotion regulation where one does not have an explicit goal to regulate, but the behaviors in which one engages directly, alter the nature of one's emotional response.

Beyond this, there are a number of ways in which prefrontal control systems may have a regulatory impact on affective learning. For example, as noted earlier, in some cases reappraisal may involve interactions between PFC, vmPFC, and the amygdala, when reappraisal paradigms give participants a chance to respond emotionally and potentially plan reappraisals before deciding whether to implement them. Interactions of this sort also have been observed in studies that use distraction to regulate a conditioned response.^{149,150} In these studies, one is initially conditioned to expect either a painful shock or reward UCS following a visual CS (e.g., a yellow triangle). Later, one regulates the conditioned response to the CS by thinking about a calm and neutral scene unrelated to either the CS or the UCS. In both cases, effective regulation involves activation of left dlPFC and modulation of both the amygdala and/or ventral striatum and the vmPFC.

Affective decision making. Affective decision making involves choosing among several stimuli that one may purchase, consume, or own. In some cases, these choices are a simple matter of selecting the option that has the greatest value. Imaging research suggests that activation in systems thought to

represent affect and value like the ventral striatum, insula, and vmPFC is sufficient to support and even predict such choices.^{151,152} But in other cases, the choice options may be of similar value, or the reasons for valuing them may conflict with one another. In the model, such cases may draw on the control systems shown in Figure 2B to modulate the values associated with choice options, essentially guiding a top-down revaluation of them to facilitate choice.

Perhaps the simplest example of this is where the act of choice itself arouses conflict as one decides which features of choice options they can't live without and which features of choice options they must forgo. Classically, this decision conflict is thought to arouse cognitive dissonance, which the act of choosing reduces by placing a higher value on chosen and a lesser value on unchosen stimuli.¹⁵³ Imaging studies show that these choice-induced changes in value involve control systems like the anterior cingulate cortex, which may signal the presence of choice conflict and motivate value change, and systems like the ventral striatum, which may represent the revalued stimuli.^{153–156}

Another type of choice that commonly requires the use of control occurs when an individual must decide between options that fit short-term versus long-term goals. This is the dilemma faced by a dieter who must decide whether to eat a cupcake or an apple. Consuming the cupcake satisfies the short-term goal of hedonic pleasure, whereas eating the apple satisfies the long-term goal of living a healthy lifestyle. A recent imaging study⁴⁷ of this choice dilemma showed that food choices reflecting a greater valuation of long-term health over short-term tastiness involve the modulation of vmPFC by dlPFC. This is consistent with the idea that the cognitive control of choice involves interactions between systems for maintaining choice goals (e.g., dlPFC) and systems representing the value of choices with respect to those goals (e.g., vmPFC).

This same logic applies to studies of intertemporal choice and delay of gratification,¹⁵⁷ where imaging^{158,159} and transcranial magnetic stimulation (TMS)¹⁶⁰ studies suggest that lateral PFC control systems can be used to effortfully represent the value of a larger delayed reward and guide selection of it over a smaller but immediately available reward.

More generally, the model can be applied to other choices where control is needed to modulate the af-

fective valuations placed on choice options, ranging from risky decision making¹⁶¹ to interpersonal contexts in which one must decide whether to be fair toward or punish others.^{162,163}

Affective expectancies. In parallel to the growth and development of imaging research on emotion regulation, there has been a tremendous surge of interest in the brain mechanisms underlying the influence of expectancies on behavior.¹⁶⁴ In imaging, expectancies have been studied either by cueing participants that an upcoming stimulus will have particular properties (e.g., that it will or will not be painful, will be a neutral or aversive image, and so on) or by inducing beliefs about the effects of a placebo drug on their experience (e.g., that an analgesic cream will reduce pain).

In the model, these phenomena all involve the use of prefrontal control systems to set and maintain an expectation, which, in turn, influences the responses of affect generating systems. For example, imaging studies show that expectancies and placebo beliefs about pain activate lateral prefrontal/parietal control systems and/or medial prefrontal systems^{164–167} that may maintain expectations about upcoming events. In turn, these systems may influence the way one attends to and appraises the meaning of expected stimuli, thereby increasing or decreasing activity in affect systems to be consistent with the nature of one's expectations.

Summary and future directions for basic and translational research

The overarching goal of this paper has been to review and synthesize current functional imaging research on emotion regulation. Toward that end, we outlined a basic model of the processes and neural systems involved in emotion generation and emotion regulation and surveyed various domains of research that support it. At its core, the MCCE specifies how prefrontal and cingulate control systems modulate activity in affect systems as a function of one's regulatory goal, tactics, and the nature of the stimuli and emotions being regulated. Although the model was built primarily from studies of one type of cognitive change strategy known as reappraisal, it is generally applicable to understanding the brain mechanisms underlying the other emotion regulation strategies depicted in Figure 2A as well as a range of other allied phenomena.

That said, there is much work yet to be done. At various points during the review we've highlighted the limitations of current knowledge and the shortcomings of current methodologies. Future work is needed to clarify the mechanisms underlying all of the emotion regulation strategies discussed here as well as the roles the brain systems supporting emotion regulation (Fig. 2B) play in affective learning, affective decision making, and affective expectancies. Progress is essential to refine our understanding of the distinctions made here, but also to address new questions about how emotion regulation mechanisms operate. For example, although it is certainly important that regulation strategies have immediate effects on emotional responses, it is also important that their effects be long lasting. Indeed, whether regulatory effects last is critical both in everyday and clinical contexts in which one could repeatedly re-encounter an emotionally evocative stimulus (e.g., the risk of running into a girlfriend who dumped you because you work for the same company). To date, this issue has been addressed only twice—once in an fMRI study¹⁶⁸ reporting that the effects of reappraisal on diminishing amygdala responses may endure for up to 40 minutes in healthy adults, but not those with major depression, and once in an ERP study¹⁶⁹ showing that the effects of reappraisal on arousal-related responses endure for up to 30 minutes. Clearly, more work is needed here.

In so doing, it will be important for this work to increasingly make use of techniques other than functional imaging (e.g., ERP,^{169–177} TMS,^{160,178} and lesion methodologies¹⁷⁹), as well as to integrate insights gained from human studies with the large body of literature on affective and regulatory phenomena in nonhuman primates and rodents.^{39,142,144} Progress on all of these fronts is absolutely critical if we are to develop a model of interactions between control and affect systems that can make sense not just of emotion regulatory phenomena, but of all the other types of phenomena that recruit these systems as well.

Another important direction for future research is the translation of basic research of emotion regulation to understanding the full range of normal to abnormal differences in emotional responding and regulatory ability. This is critical both for understanding the mechanisms underlying this variability and for testing the boundaries of basic models of emotion regulatory mechanisms.

One domain in which this will prove important is understanding how and why our emotional lives evolve as we grow from childhood through adolescence into adulthood and old age. On one hand, there is growing evidence that childhood and adolescence are critical times for the development of the emotion regulatory abilities needed to adaptively regulate affective impulses and the deleterious health behaviors (e.g., obesity, substance use) they can promote. A small but growing number of studies have begun to address this issue by asking how the neural mechanisms of reappraisal and emotional reactivity develop from adolescence into young adulthood. Some early results suggest that reappraisal ability increases linearly with age, whereas emotional reactivity remains relatively constant^{180,181} (but see Ref. 182). On the other hand, although physical health and cognitive abilities tend to decline with age,^{183–185} older adults report more emotional stability and a greater ratio of positive to negative experiences in their daily life, with the extent of positive emotion predicting longevity.^{186–188} Although many have hypothesized that this “rosy glow” of old age is due in part to more effective emotion regulation, to date there is little evidence directly testing this idea.^{187,189,190} One conundrum to resolve here will be the apparent dependence of emotion regulation on the same kinds of prefrontal control systems that decline with age. This raises the question of how regulatory abilities improve as the underlying neural machinery declines.^{191,192} Early results suggest that it may depend on the strategies older adults deploy, with spared or greater regulatory ability shown for strategies and tactics that fit with long-term goals and have become habitual.^{9,96,190,193–196}

A second important goal for translational research will be to understand how potential dysfunction in the mechanisms of emotion generation and regulation may underlie various forms of psychiatric and substance use disorders (for a more in depth discussion, please see Ref. 197). This translational direction is being pursued in studies of reappraisal across various disorders, ranging from depression^{116,168,198} to borderline personality disorder,^{83,87,199} social anxiety disorder,^{200,201} phobia,²⁰² posttraumatic stress disorder,^{83,85} cocaine users,²⁰³ and smokers.¹¹⁸ These studies can be useful in two ways. First, they may show disorder-specific patterns of altered function in control and affect systems. For example, current data suggest that depressed

individuals may show impaired recruitment of vIPFC during reappraisal,¹¹⁶ suggestive of an impairment of top-down control, whereas borderline individuals may show heightened amygdala responses coupled with diminished cingulate responses, suggestive of a failure to monitor paradoxical increases in affective responding when attempting to decrease emotion.¹⁹⁹ Second, imaging methods for studying emotion regulation may be used before and after treatment regimes as predictors of and markers of improvement. Although such studies are only beginning to emerge, they hold great promise for understanding why some individuals improve and whether different treatments (e.g., drugs versus cognitive behavioral therapy) have different mechanisms of action.

In the long run, the hope is that integrating basic and translational perspectives will help specify which individuals are at greatest risk for maladaptive health behaviors and emotional outcomes, at what ages this risk is greatest, and which regulatory mechanisms could be targeted in future interventions during particular points in the life course. Although realization of this dream is still a long way away, current research provides a strong foundation for getting there.

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Conflicts of interest

The authors declare no conflicts of interest.

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