Title

Neural precursors of decisions that matter—an ERP study of deliberate and arbitrary choice

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Abstract

The onset of the readiness potential (RP)—a key neural correlate of upcoming action—was repeatedly found to precede subjects' reports of having made an internal decision. This has been taken by some as evidence against a causal role for consciousness in human decision-making and thus as a denial of free-will. Yet those studies focused on purposeless, unreasoned, arbitrary decisions, bereft of consequences. It remains unknown to what degree these specific neural precursors of action generalize to deliberate decisions, which are more ecological and relevant to real life. We therefore directly compared the neural correlates of deliberate and arbitrary decision-making during a $1000-donation task to non-profit organizations. While we found the expected RPs for arbitrary decisions, they were strikingly absent for deliberate ones. Our results are congruent with the RP representing the accumulation of noisy, random fluctuations, which drive arbitrary—but not deliberate—decisions. The absence of RPs in deliberate decisions further points to different neural mechanisms underlying deliberate and arbitrary decisions and thus challenges the generalizability of studies that argue for no causal role for consciousness in decision making from arbitrary to deliberate, real-life decisions.

Significance Statement

The extent of human free will is among the most hotly contended, long-standing debates in the history of human thought. Previous studies demonstrated that neural precursors of upcoming actions—especially the EEG readiness potential component—precede subjects' reports of having decided to move. Some took this as evidence against free-will. However, these experiments focused on arbitrary decisions—e.g., randomly raising the left or right hand. We directly compared neural precursors of deliberate decisions (actual $1000 donations to NPOs) and arbitrary ones, and found readiness potentials before arbitrary decisions, but—critically—not before deliberate decisions. This supports the interpretation of readiness potentials as byproducts of accumulation of random fluctuations in arbitrary but not deliberate decisions. It also points to different neural mechanisms underlying deliberate and arbitrary decisions and thus challenges the generalizability of previous neuroscientific studies of arbitrary decisions to deliberate ones.

MAIN TEXT

Introduction

Humans typically experience freely selecting between alternative courses of action, say, when ordering a particular item off a restaurant menu. Yet a series of human studies using EEG (1-3), fMRI (4-7), intracranial (8), and single-cell recordings (9) challenged the validity of this common experience, finding neural correlates of decision processes hundreds of milliseconds and even seconds prior to the moment that subjects reported having consciously decided. Some have claimed, following these and other findings, that the subjective human experience of freely deciding is but an illusion, because human actions are unconsciously initiated before the conscious decision to act (1, 10, 11). This debate has been captivating scholars from many disciplines in and outside of academia (10, 12-16).

Critically, in the above studies, subjects were either told to move their right hand or flex their right wrist; or they were instructed to move either the right or left hand (14, 17, 18). Thus, their decisions were always unreasoned, purposeless, and bereft of any real consequence. This stands in sharp contrast to real-life decisions that are reasoned, purposeful, and bear consequences (19)—which clothes to wear, what route to take to work, as well as more formative decisions about life partners, career choices, and so on. Such deliberate decisions are also at the center of the philosophical debate on free will (17, 20). They typically involve more conscious and lengthy deliberation, and might thus be more tightly bound to conscious processes.

Deliberate decisions have been widely studied in the field of neuroeconomics (21, 22) and in perceptual tasks (23). Yet, interestingly, little has been done in that field to assess the relation between decision-related activity, subjects' conscious experience of deciding, and the neural activity instantaneously contributing to this experience. Though some studies compared, for example, internally driven and externally cued decisions (24, 25), or stimulus-based and intention-based actions (26), these were typically arbitrary decisions and actions that had no real implications.

Here, we compared neural precursors of deliberate and arbitrary decisions—and in particular the readiness potentials (RP), as they were the focus of so many previous studies of voluntary action—on the same subjects, in an EEG experiment. Our experiment utilized a donation-preference paradigm, in which a pair of non-profit organizations (NPOs) were presented in each trial. In deliberate-decision trials, subjects chose to which NPO they would like to donate $1000. In arbitrary-decision trials, both NPOs received an equal donation of $500, irrespective of subjects' key presses (Fig. 1). In both conditions, subjects were instructed to report their decisions as soon as they made them, and their hands were placed on the response keys, to make sure they could do so as quickly as possible. Notably, while the visual inputs and motor outputs were identical between deliberate and arbitrary decisions, the decisions' meaning for the subjects was radically different: in deliberate blocks, the decisions were meaningful and consequential—reminiscent of important, real-life decisions—while in arbitrary blocks, the decisions were meaningless and bereft of consequences—mimicking previous studies of volition. Demonstrating differences in RP between arbitrary and deliberate decisions would first challenge the generalizability of the RP (from arbitrary to deliberate decisions) as an index for internal decision-making. Second, it would more generally suggest that different neural mechanisms might be at play between deliberate and arbitrary decisions. This, in turn, would question the generalizability of studies focused on arbitrary decisions to everyday, ecological, deliberate decisions—regardless of whether these studies relied on the RP or not.

Results

Behavioral Results

Figure 1: Experimental paradigm. The experiment included deliberate (red, left panel) and arbitrary (blue, right panel) blocks, each containing nine trials. In each trial, two NPO names were presented, and subjects were asked to either choose to which NPO they would like to donate (deliberate), or to simply press either right or left, as both NPOs would receive an equal donation (arbitrary). They were specifically instructed to respond as soon as they reached a decision, in both conditions. Within each block, some of the trials were easy (lighter colors) decisions, where the subject's preferences for the two NPOs substantially differed (based on a previous rating session), and some were hard decisions (darker colors), where the preferences were more similar; easy and hard trials were intermixed within each block. To make sure subjects were paying attention to the NPO names, even in arbitrary trials, and to better equate the cognitive load between deliberate and arbitrary trials, memory tests (in light grey) were randomly introduced, where subjects were asked to determine which of four NPO names appeared in the immediately previous trial.

To validate the experimental paradigm, we manipulated decision difficulty as well as decision type (Fig. 1). We reasoned that this should affect only deliberate decisions, and not arbitrary ones. This was borne out in our subjects' behavioral patterns. Subjects' reaction times (RTs) were substantially slower for deliberate than for arbitrary decisions (Fig. 2, left; $F(1,17)=126.11$, $p<0.0001$ for the main effect of decision type, as revealed by a 2-way ANOVA with decision type (arbitrary/deliberate) and difficulty (easy/hard). The ANOVA also showed a main effect of decision difficulty, $F(1,17)=18.76$, $p=0.0004$; all analyses were performed on log-transformed data, because the raw RTs violated the normality assumption ($W=0.94$, $p=0.001$)). Moreover, subjects were slower for hard vs. easy deliberate decisions ($F(1,17)=20.12$, $p=0.0003$ for the interaction between decision type and decision difficulty; hard vs. easy deliberate decisions: $t(17)=4.78$, $p=0.0002$, yet not for arbitrary decisions: $t(17)=1.01$, $p=0.33$). This further demonstrates that, in deliberate decisions, subjects were making meaningful decisions, affected by the difference in the values of the two NPOs, while in arbitrary decisions they were not.

The consistency between subjects' choices throughout the main experiment and the NPO ratings they gave prior to the main experimental session was also analyzed (see Methods). As expected, subjects were highly consistent with their own, previous ratings when making deliberate decisions, but not when making arbitrary ones (Fig. 2, right; 2-way ANOVA: $F_{(1,17)}=946.55$, $p<0.0001$) for the main effect of decision type. A main effect of decision difficulty was also found ($F_{(1,17)}=57.39$, $p<0.0001$). Again, decision type and decision difficulty interacted ($F_{(1,17)}=25.96$, $p<0.0001$: subjects were much more consistent with their choices in easy vs. hard deliberate decisions ($t_{(17)}=11.15$, $p<0.0001$), than they were in easy vs. hard arbitrary decisions ($t_{(17)}=2.50$, $p=0.028$). Nevertheless, though subjects were around chance (i.e., 0.5) in their consistency in arbitrary decisions (ranging between 0.39 and 0.64), it seems that some subjects were slightly influenced by their preferences in easy-arbitrary decisions trials, resulting in the significant difference between hard-arbitrary and easy-arbitrary decisions above. Finally, no differences were found between subjects' tendency to press the right vs. left key in the different conditions (both main effects and interaction: $F<1$).

Figure 2: Behavioral results. Response Times (RTs; left) and Consistency Grades (CG; right) in arbitrary (blue) and deliberate (red) decisions. Each dot represents the average RT/CG for easy and hard decisions for an individual subject (hard decisions: x-coordinate; easy decisions: y-coordinate). Group means and SEs are represented in dark red and dark blue crosses. The histograms at the bottom-left corner of each plot sum the number of dots with respect to the solid diagonal line. The dashed diagonal line represents equal RT/CG for easy and hard decisions; data points below that diagonal indicate longer RTs or higher CGs for hard decisions. In both measures, arbitrary decisions are more centered around the diagonal than deliberate decisions, showing no or substantially reduced differences between easy and hard decisions.

Figure 3: The readiness potentials for deliberate and arbitrary decisions. (A) Mean and SE of the Readiness Potential (RP) in deliberate (red shades) and arbitrary (blue shades) easy and hard decisions in electrode Cz, as well as scalp distributions. Zero refers to time of right/left movement, or response, made by the subject. Notably, the RP significantly differs from zero and displays a typical scalp distribution for arbitrary decisions only. The scalp distribution was calculated over the averaged activity during the last 500 ms before response, across subjects. The inset shows the mean amplitude of the RP, with 95% confidence intervals over the same time window. Response-locked potentials with an expanded timecourse, and stimulus-locked potentials are given in Fig. S3B and S3A, respectively. The same (response-locked) potentials as here, but with a *movement-locked baseline* of -1 to -0.5 s (same as in our Bayesian analysis below), are given in Fig. S3C. **(B)** Individual subjects' Cz activity in the four conditions (n=18). The linear-regression line for voltage against time over the last 1000 ms before response onset is designated by a dashed, black line. Note that the waveforms converge to an RP only in arbitrary decisions.

The RP is generally held to index unconscious readiness for upcoming movement (1, 14, 27, 28); although more recently alternative interpretations of the RP have been suggested (29-33). It has nevertheless been the standard component studied in EEG versions of the Libet paradigm (1, 2, 14, 28-31, 34-36). Here, the RP was measured over electrode Cz in the different conditions by averaging the activity across trials in the 2 s prior to subjects' movement. Focusing on the last 500 ms before movement onset for our statistical tests, we found a clear RP in arbitrary decisions, yet RP amplitude was not significantly different from 0 in deliberate decisions (Fig. 3A; ANOVA $F(1,17)=11.86$, $p=0.003$ for the main effect of decision type; in t-tests against 0, corrected for multiple comparisons, an effect was only found for arbitrary decisions (hard: $t(17)=5.75$, $p<0.0001$; easy: $t(17)=5.09$, $p=0.0004$) and not for deliberate ones (hard: $t(17)=1.24$, $p>0.5$; easy: $t(17)=1.84$, $p=0.34$). In a similar manner, regressing voltage against time for the last 1000 ms before response onset, the downward trend was significant for arbitrary decisions (Fig. 3B; $p<0.0001$, for both easy and hard conditions) but not for deliberate decisions (hard: $p>0.5$, easy: $p=0.35$; all Bonferroni corrected for multiple comparisons). Notably, this pattern of results was

also manifested for single-subject analysis (Fig. S1; 14 of the 18 subjects had significant downward slopes for arbitrary decisions—i.e., p<0.05, Bonferroni corrected for multiple comparisons—when regressing voltage against time for every trial over the last 1000 ms before response onset; but only 5 of the 18 subjects had significant downward slopes for the same regression analysis for deliberate decisions; see Methods. In addition, the average slopes for deliberate and arbitrary decisions were -0.43±0.31 and -2.30±0.44 (mean±SE), respectively, a significant difference: t(17)=3.51, p=0.001).

To test whether the null result for RP amplitude in deliberate decisions stems from a genuine absence of effect or from insufficient or underpowered data, we used Bayesian statistics. Specifically, the Bayes factor allowed us to compare the probability of observing the data given H_0 (i.e., no RP in deliberate decisions) against the probability of observing the data given H_1 (i.e., RP exists in deliberate decisions). We followed the convention that a BF < 0.33 implies substantial evidence for lack of an effect (that is, the data is three times more likely to be observed given H_0 than given H_1), 0.33 < BF < 3 suggests insensitivity of the data, and BF > 3 denotes substantial evidence for the presence of an effect (H_1) (37). We found strong evidence that arbitrary trials (pooled across difficulty for more statistical power) are different from 0 (BF=$2.82 \cdot 10^6$), but inconclusive evidence for deliberate trials (BF=1.46, again for pooled trials). Notably, however, this result might reflect the general, slow negative trend of the deliberate trials data (Fig. 3A) rather than a typical RP, which is locked to movement-onset and shows a sharp decline towards that. To remove the effect of this slow negative drift, we used a baseline period of -1000 ms to -500 ms relative to *movement* onset (i.e., a baseline that immediately preceded our time of interest window) instead of our usual baseline that preceded stimulus onset (see Methods). Under this analysis, we found evidence that deliberate decisions are not different from 0 (BF=0.332), supporting the claim that the RP during the last 500 ms before response onset was completely absent (BF for arbitrary decisions was $5.07 \cdot 10^4$).

We further tested whether differences in reaction time between the conditions, eye movements, the hand with which the subjects executed the movement, and subjects' consistency scores might explain our effect. We also tested whether the RPs might reflect some stimulus-locked potentials or be due to baseline considerations. We found that none of these factors drive the effect (see Supplementary Results).

EEG Results: Lateralized Readiness Potential (LRP)

The LRP, which reflects activation processes within the motor cortex for action preparation after action selection (38, 39), was measured by subtracting the difference potentials (C3-C4) in right-hand response trials from this difference in left-hand responses trials and averaging the activity over the same time window (2, 38). In this purely motor component, no difference was found between the two decision types (Fig. S6; ANOVA all Fs<1). It should be noted that the LRP we receive is relatively large (compared to (2)). Our analysis of EOG channels suggests that some of that LRP might be driven by eye movements (we repeated the LRP computation on the EOG channels instead of C3 and C4). However, the shape of the eye-movement-induced LRP is very different from the LRP we calculated from C3 and C4. Also, the differences that we found between conditions in the EOG LRP are not reflected in the C3/C4 LRP. So, while our LRP might be boosted by eye movements, it is not strictly driven by these eye movements.

Drift Diffusion Model (DDM)

The main finding of this study—the absence of RP in deliberate decisions, suggesting different neural underpinnings of arbitrary and deliberate decisions—is in line with recent work that used a drift-diffusion model (DDM) to claim that the RP might arise from the nonlinearities associated with threshold crossing together with time-locking neural activity to response onset, rather than from specific preparation for or ballistic-like initiation of movement (29). DDMs of decision-making typically feature a process that rises toward a threshold. The crossing of that threshold reflects the onset of the decision in the model, typically leading to action. Schurger and colleagues (29) modelled arbitrary decisions, and suggested that there the threshold crossing leading to response onset is largely determined by spontaneous subthreshold fluctuations of the neural activity. This challenged the common view of the RP as a neural correlate of unconscious preparation for upcoming action (28). Instead, Schurger and colleagues claimed, time-locking to response onset ensures that these spontaneous fluctuations appear, when averaged over many trials, as a gradual increase in neural activity.

To further assess this interpretation of the RP, we expanded the model developed by Schurger and colleagues (29) to a DDM that was composed of a *value-assessment* component and a *noise-generation* component. Under this assumption, Cz-electrode activity mainly reflects the noise-generation component. (Note that we suggest that noise generation might be a key function of the (pre-)SMA and other brain regions underneath the Cz electrode in this specific task. When subjects make arbitrary decisions, these might be based on some symmetry-breaking mechanism, which is driven by random fluctuations that are here simulated as noise, following Schurger et al. Thus, we do not claim or think that noise generation is the main purpose or function of these brain regions in general.) In addition to the value generation and noise generation components, each trial was modeled as a race to threshold between two alternatives: one where the subject chose the NPO that was rated higher in the earlier rating session (the *congruent* option) and the other where the subject preferred the lower-rated NPO (the *incongruent* option). Each race was represented as a leaky stochastic accumulator (the Methods and Supplementary Methods provide more details and the model parameters).

We fit our DDM to our average empirical reaction-times, which were 2.13, 2.52, 0.98 and 1.00 s for the different conditions (henceforth, magnitudes are given for deliberate easy, deliberate hard, arbitrary easy, and arbitrary hard, respectively, in this order). The model's corresponding mean RTs were 2.04, 2.46, 0.94, and 0.96 s for these conditions (Fig. 4A, B). The model was further fit to the empirical consistency ratios (the proportions of congruent decisions), which were 0.99, 0.83, 0.54 and 0.49. The model's corresponding consistency ratios were 1.00, 0.84, 0.53 and 0.53. The model then predicted the shape of the ERP in its noise component (assumed to be reflected by Cz-electrode activity) for each decision type: a continuing, RP-like increase in activity (with a negative sign) for arbitrary decisions, but only a very slight increase in activity for deliberate decisions (Fig. 4C; see also Fig. S7). This was in line with our empirical results (Fig. 3A). Note that that the Schurger model aims to account for neural activity leading up to the decision to move, but no further (29). Similarly, we expect our DDM to fit Cz neural data only up to around -0.1 s (100 ms before response onset). We also make no claims that ours is the only, or even optimal, model that explains our results. Rather, by extending the Schurger model, we show how that interpretation of the RP could also be applied to our more-complex paradigm.

Figure 4: Empirical and model RTs and model prediction for Cz activity. (A) The empirical distributions of subjects' RTs across the four decision types. **(B)** The equivalent distributions of RTs for the model. **(C)** The model's prediction for the ERP activity in electrode Cz across all four decision types.

Discussion

Since the publication of Libet's seminal work claiming that neural precursors of action, in the form of the RP, precede subjects' reports of having consciously decided to act (1), a vigorous discussion has been raging among neuroscientists, philosophers, and other scholars about the meaning of these findings for the debate on free will (recent collections include (40-42)). Some claim that these results have removed conscious will from the causal chain leading to action (10, 14, 43). Others are unconvinced that these results are decisive for, or even applicable to, the free-will debate (17, 20, 44, 45). At the heart of much of this debate lies the RP, thought to represent unconscious decision/planning mechanisms that initiate subjects' decisions prior to their conscious experience of deciding (1, 27).

Notably, the RP and similar findings showing neural activations preceding the conscious decision to act have typically been based on arbitrary decisions of different types (1, 2, 4, 5, 15, 35, 46). This, among other reasons, rested on the notion that for an action to be completely free, it should

not be determined in any way by external factors (35)—which is the case for arbitrary, but not deliberate, decisions (where each decision alternative is associated with a value, and the values of alternatives typically guide one's decision). But this notion of freedom faces several obstacles. First, most discussions of free will focus on deliberate decisions, asking when and whether these are free (47-49). This might be because everyday decisions to which we associate freedom of will—like choosing a more expensive but more environmentally friendly car, helping a friend instead of studying more for a test, donating to charity, and so on—are generally deliberate, in the sense of being reasoned, purposeful, and bearing consequences (although see (50)). In particular, the free will debate is often considered in the context of moral responsibility (e.g., was the decision to harm another person free or not) (14, 51-55), and free will is even sometimes defined as the capacity that allows one to be morally responsible (16, 44). In contrast, it seems meaningless to assign blame or praise to arbitrary decisions. Thus, though the scientific operationalization of free will has typically focused on arbitrary decisions, the common interpretations of these studies—in neuroscience and across the free will debate—have often alluded to deliberate ones. Here, we show that inference from arbitrary to deliberate decisions may not be justified, as the neural precursors of arbitrary decisions, and in particular the RP, do not generalize to meaningful ones (17, 20). This therefore challenges previous studies relying on arbitrary decisions, regardless of whether they were based on the RP or not.

Interestingly, while the RP was present in arbitrary decisions but absent in deliberate ones, the LRP—a long-standing, more-motor ERP component—was indistinguishable between the different decision types. This provides evidence that, at the motor level, the neural representation of the deliberate and arbitrary decisions that our subjects made may have been indistinguishable, as was our intention when designing the task.

Our finding and the model thus suggest that two different mechanisms may be involved in arbitrary and deliberate decisions. Earlier literature demonstrated that deliberate, reasoned decision-making—which was mostly studied in the field of neuroeconomics (21) or using perceptual decisions (23)—elicited activity in the prefrontal cortex (PFC; mainly the dorsolateral (DLPFC) part (56, 57) and ventromedial (VMPFC) part/orbitofrontal cortex (OFC) (58, 59) and the anterior cingulate cortex (ACC) (60, 61). Arbitrary, meaningless decisions, in contrast, were mainly probed using variants of the Libet paradigm, showing activations in the Supplementary Motor Area (SMA), alongside other frontal areas like the medial frontal cortex (62, 63) or the frontopolar cortex, as well as the posterior cingulate cortex (4, 9) (though see (64), which suggests that a common mechanism may underlie both decision types). Possibly then, arbitrary and deliberate decisions may differ not only with respect to the RP, but may be subserved by different underlying neural circuits, which makes generalization from one class of decisions to the other more difficult. Deliberate decisions are associated with more lateralized and central neural activity while arbitrary ones are associated with more medial and frontal ones. This appears to align with the different brain regions associated with the two decision types above, as also evidenced by the differences we found between the scalp distributions of arbitrary and deliberate decisions (Fig. 3A). Future studies are needed to further explore this potential divergence in the neural regions between the two decision types.

To be clear, and following the above, we do not claim that the RP captures all unconscious processes that precede conscious awareness. However, some have suggested that the RP represents unconscious motor-preparatory activity before any kind of decision (e.g., (35)). But our results provide evidence against that claim, as we do not find an RP before deliberate decisions, which also entail motor preparation. What is more, in deliberate decisions in particular, it is likely that there are neural precursors of upcoming actions—possibly involving the above neural circuits

as well as circuits that represents values—which are unrelated to the RP. Note also that we did not attempt to separately measure the timing of subjects' conscious decision to move. Rather, we instructed them to hold their hands above the relevant keyboard keys and press their selected key as soon as they made up their minds. This was both to keep the decisions in this task more ecological and because we think that the key method of measuring decision onset (using some type of clock to measure Libet's W-time) is highly problematic (see Methods).

Aside from highlighting the neural differences between arbitrary and deliberate decisions, this study also challenges a common interpretation of the function of the RP. If the RP is not present before deliberate action, it does not seem to be a necessary link in the general causal chain leading to action. Schurger and colleagues (29) suggested that the RP reflects the accumulation of stochastic fluctuations in neural activity that lead to action, following a threshold crossing, when humans arbitrarily decide to move. The shape of the RP therefore results from the manner in which it is computed: averaged over trials that are locked to response onset (that directly follows the threshold crossing). Our results and our model are in line with that interpretation and expand upon it, suggesting that the RP represents the accumulation of noisy, random fluctuations that drive arbitrary decisions, while deliberate decisions are mainly driven by the values associated with the decision alternatives (65). (For discussion of other, alternative interpretations of our results see Supplementary Discussion.)

Our drift-diffusion model was based on the assumption that every decision is driven by a component based on the values of the decision alternatives (the subject's support for the two NPOs in our case) and by another component representing noise—random fluctuations in neural activity. The value component plays little to no role in arbitrary decisions, so action selection and timing depend on when the accumulation of noise crosses the decision threshold for the congruent and incongruent decision alternatives. In deliberate decisions, in contrast, the value component drives the decisions, while the noise has a smaller effect. Thus, in arbitrary decisions, action onset closely tracks threshold crossings of the noise accumulation. But, in deliberate decisions, the noise component is at more random levels at the moment of response onset. Hence, locking the ERP to response onset and averaging over trials to obtain the RP, leads to a relatively flat signal for deliberate decisions but to the expected RP shape in arbitrary decisions. This provides strong evidence that the RP does not reflect subconscious movement preparation. Rather, it is induced by threshold crossing of random fluctuations in arbitrary decisions, which do not drive deliberate decisions; accordingly, the RP is not found there. Further studies of the causal role of consciousness in deliberate versus arbitrary decisions are required to test this claim.

Our study, which suggests that RPs do not precede deliberate decisions, further motivates future investigations into other precursors of action besides the RP using EEG, fMRI or other techniques. It would be of interest to find the neural activity that precedes deliberate decisions in particular. And it would also be of interest to find neural activity, which is not motor activity, that is common to both deliberate and arbitrary decisions.

Materials and Methods

Study Design

The objective of this study was to compare ERPs elicited by arbitrary and deliberate decision-making, and in particular the RP. We further manipulated decision difficulty to validate our manipulation of decisions type: we introduced hard and easy decisions which corresponded to small and large differences between subjects' preferences for the pairs of presented NPOs,

respectively. We reasoned that if the manipulation of decision type (arbitrary vs. deliberate) was effective, there would be behavioral differences between easy and hard decisions for deliberate choices but not for arbitrary choices (because differences in preferences should not influence subjects' arbitrary decisions). Our 2 x 2 design was therefore decision type (arbitrary vs. deliberate) by decision difficulty (easy vs. hard). Each condition included 90 trials, separated into 10 blocks of 9 trials each, resulting in a total of 360 trials and 40 blocks. Blocks of different decision types were randomly intermixed. Decision difficulty was randomly counterbalanced across trials within each block.

Experimental Procedure

In the first part of the experiment, subjects were presented with each of the 50 NPOs and the causes with which the NPOs were associated separately (see Supplementary Table S2). They were instructed to rate how much they would like to support that NPO with a $1000 donation on a scale of 1 ("I would not like to support this NPO at all) to 7 ("I would very much like to support this NPO"). No time pressure was put on the subjects, and they were given access to the website of each NPO to give them the opportunity to learn more about the NPO and the cause it supports.

After the subjects finished rating all NPOs, the main experiment began. In each block of the experiment, subjects made either deliberate or arbitrary decisions. The instructions for carrying out their decisions were nevertheless identical for both decision types: Subjects were instructed to report their decisions as soon as they made them in both conditions. They were further asked to place their fingers on the response keys, so they could respond as quickly as possible. Two succinct cause descriptions, representing two actual NPOs, were presented in each trial (Fig. 1). In deliberate blocks, subjects were instructed to choose the NPO to which they would like to donate $1000 as soon as they decided. Subjects were informed that one of their selections would be randomly picked and the selected NPO would receive $20. In addition, one of the selected NPOs from one subject would be randomly picked to receive a $1000 donation. Arbitrary trials were similar to deliberate trials except that both NPOs that appeared in the randomly selected decision would receive $10 each, and both NPOs from the randomly picked subject would receive $500 each. Thus, subjects' decisions in the deliberate blocks were meaningful and consequential, while their decisions in arbitrary blocks had no impact on the final donations that were made (for more details, see Supplementary Methods).

ERP recording methods

The EEG was recorded using an Active 2 system (BioSemi, the Netherlands) from 64 electrodes distributed based on the extended 10–20 system and connected to a cap, and seven external electrodes. Four of the external electrodes recorded the EOG: two located at the outer canthi of the right and left eyes and two above and below the center of the right eye. Two external electrodes were located on the mastoids, and one electrode was placed on the tip of the nose. All electrodes were referenced during recording to a common-mode signal (CMS) electrode between POz and PO3. The EEG was continuously sampled at 512 Hz and stored for offline analysis.

ERP analysis

ERP analysis was conducted using the "Brain Vision Analyzer" software (Brain Products, Germany) and in-house Mathworks Matlab scripts. Data from all channels were referenced offline to the average of all channels. The data were then digitally high-pass filtered at 0.1 Hz using a Finite Impulse Response (FIR) filter to remove slow drifts. A notch filter at 59-61 Hz was applied

to the data to remove 60-Hz electrical noise. The signal was then cleaned of blink and saccade artifacts using Independent Component Analysis (ICA) (66). Signal artifacts were detected as amplitudes exceeding ±100 µV, differences beyond 100 µV within a 200 ms interval, or activity below 0.5 µV for over 100 ms (the last condition was never found). Sections of EEG data that included such artifacts in any channel were removed (150 ms before and after the artifact), leaving an average number of 70.38 trials with a range of 36-86 out of 90 trials per condition. Channels that consistently had artifacts were replaced using interpolation (4.2 channels per subject, on average).

The EEG was segmented by locking the waveforms to subjects' movement onset, starting 2s prior to the movement and ending 0.2s afterwards, with the segments averaged separately for each decision type (Deliberate/Arbitrary x Easy/Hard) and decision content (right/left). The baseline period was defined as the time window between -1000 ms and -500 ms prior to *stimulus* onset, that is, the onset of the causes screen, rather than prior to movement onset. In addition to the main baseline, we tested another baseline—from -1000 ms to -500 ms relative to *movement* onset—to investigate whether the baseline period influenced our main results (Figs S3B, C). Furthermore, we segmented the EEG based on *stimulus* onset, using the same baseline, for stimulus-locked analysis (Fig S3A).

To assess eye movements during the experiment, we defined the radial eye signal as the average over all 4 EOG channels, when band-pass filtered to between 30 and 100 Hz. We then defined a saccade as any signal that was more than 2.5 standardized IQRs away from the median of the radial signal for more than 2 ms. Two consecutive saccades had to be at least 50 ms apart. The saccade count (SC) was the number of saccades during the last 500 ms before response onset (67) (see also (68-70)).

Additional details about the study design, subjects, stimuli and apparatus, experimental procedure, statistical analysis, and model and simulations are provided in the Supplementary Materials and Methods section.

Supplementary Materials

References and Notes

1. Libet B, Gleason C, Wright E, & Pearl D (1983) Time of conscious intention to act in relation to onset of cerebral activity (readiness-potential): The unconscious initiation of a freely voluntary act. *Brain* 106(3):623.
2. Haggard P & Eimer M (1999) On the relation between brain potentials and the awareness of voluntary movements. *Experimental Brain Research* 126(1):128-133.
3. Salvaris M & Haggard P (2014) Decoding Intention at Sensorimotor Timescales. *PLoS One* 9(2):e85100.
4. Soon C, Brass M, Heinze H, & Haynes J (2008) Unconscious determinants of free decisions in the human brain. *Nature Neuroscience* 11(5):543-545.
5. Soon C, He A, Bode S, & Haynes J (2013) Predicting free choices for abstract intentions. *Proceedings of the National Academy of Sciences* 110(15):6217-6222.
6. Bode S, *et al.* (2011) Tracking the unconscious generation of free decisions using uitra-high field fMRI. *PloS one* 6(6):e21612.
7. Bode S & Haynes JD (2009) Decoding sequential stages of task preparation in the human brain. *Neuroimage* 45(2):606-613.
8. Perez O, *et al.* (2015) Preconscious prediction of a driver's decision using intracranial recordings. *Journal of cognitive neuroscience*.
9. Fried I, Mukamel R, & Kreiman G (2011) Internally generated preactivation of single neurons in human medial frontal cortex predicts volition. *Neuron* 69:548-562.
10. Wegner D (2002) *The illusion of conscious will* (MIT Press).
11. Harris S (2012) *Free will* (Simon & Schuster, Inc., New York, NY).
12. Frith C, Blakemore S, & Wolpert D (2000) Abnormalities in the awareness and control of action. *Philosophical Transactions of the Royal Society B: Biological Sciences* 355(1404):1771.
13. Jeannerod M (2006) *Motor cognition: What actions tell the self* (Oxford University Press).
14. Haggard P (2008) Human volition: towards a neuroscience of will. *Nature Reviews Neuroscience* 9(12):934-946.
15. Lau H, Rogers R, Haggard P, & Passingham R (2004) Attention to intention. *Science* 303(5661):1208.
16. Mele A (2006) *Free will and luck* (Oxford University Press).

17.   Roskies A (2010) How Does Neuroscience Affect Our Conception of Volition? *Annual Review of Neuroscience* 33:109-130.
18.   Hallett M (2016) Physiology of free will. *Annals of neurology* 80(1):5-12.
19.   Ullmann-Margalit E & Morgenbesser S (1977) Picking and choosing. *Social Research* 44(4):757-783.
20.   Breitmeyer BG (1985) Problems with the psychophysics of intention. *Behavioral and brain sciences* 8(04):539-540.
21.   Kable JW & Glimcher PW (2009) The neurobiology of decision: consensus and controversy. *Neuron* 63(6):733-745.
22.   Sanfey A, Loewenstein G, McClure S, & Cohen J (2006) Neuroeconomics: cross-currents in research on decision-making. *Trends in Cognitive Sciences* 10(3):108-116.
23.   Gold J & Shadlen M (2007) The neural basis of decision making.
24.   Wisniewski D, Goschke T, & Haynes J-D (2016) Similar coding of freely chosen and externally cued intentions in a fronto-parietal network. *NeuroImage* 134:450-458.
25.   Thut G*, et al.* (2000) Internally driven vs. externally cued movement selection: a study on the timing of brain activity. *Cognitive Brain Research* 9(3):261-269.
26.   Waszak F*, et al.* (2005) Intention-based and stimulus-based mechanisms in action selection. *Experimental brain research* 162(3):346-356.
27.   Kornhuber H & Deecke L (1990) Readiness for movement—the Bereitschafts potential-story. *Current Contents Life Sciences* 33(4):14.
28.   Shibasaki H & Hallett M (2006) What is the Bereitschaftspotential? *Clinical Neurophysiology* 117(11):2341-2356.
29.   Schurger A, Sitt JD, & Dehaene S (2012) An accumulator model for spontaneous neural activity prior to self-initiated movement. *Proceedings of the National Academy of Sciences* 109(42):E2904-E2913.
30.   Trevena J & Miller J (2010) Brain preparation before a voluntary action: Evidence against unconscious movement initiation. *Consciousness and cognition* 19(1):447-456.
31.   Miller J, Shepherdson P, & Trevena J (2011) Effects of Clock Monitoring on Electroencephalographic Activity. *Psychological Science* 22(1):103-109.
32.   Schmidt S, Jo H-G, Wittmann M, & Hinterberger T (2016) 'Catching the waves'–slow cortical potentials as moderator of voluntary action. *Neuroscience & Biobehavioral Reviews* 68:639-650.
33.   Verleger R, Haake M, Baur A, & Śmigasiewicz K (2016) Time to Move Again: Does the Bereitschaftspotential Covary with Demands on Internal Timing? *Frontiers in human neuroscience* 10.
34.   Hallett M (2007) Volitional control of movement: The physiology of free will. *Clinical Neurophysiology* 118(6):1179-1192.
35.   Libet B (1985) Unconscious cerebral initiative and the role of conscious will in voluntary action. *Behavioral and brain sciences* 8:529-539.
36.   Libet B, Wright E, & Gleason C (1982) Readiness-potentials preceding unrestricted 'spontaneous' vs. pre-planned voluntary acts. *Electroencephalography and Clinical Neurophysiology* 54(3):322-335.
37.   Jeffreys H (1998) *The theory of probability* (OUP Oxford).
38.   Eimer M (1998) The lateralized readiness potential as an on-line measure of central response activation processes. *Behavior Research Methods, Instruments, & Computers* 30(1):146-156.
39.   Masaki H, Wild-wall N, Sangals J, & Sommer W (2004) The functional locus of the lateralized readiness potential. *Psychophysiology* 41(2):220-230.
40.   Mele A (2015) *Surrounding free will: philosophy, psychology, neuroscience* (Oxford University Press).

41. Sinnott-Armstrong W & Nadel L eds (2011) *Conscious will and responsibility: A tribute to Benjamin Libet* (Oxford University Press).
42. Pockett S, Banks WP, & Gallagher S (2009) *Does consciousness cause behavior?* (Mit Press).
43. Haggard P (2005) Conscious intention and motor cognition. *Trends in Cognitive Sciences* 9(6):290-295.
44. Mele A (2009) *Effective intentions: the power of conscious will* (Oxford University Press, USA).
45. Nahmias E, Shepard J, & Reuter S (2014) It's OK if 'my brain made me do it': People's intuitions about free will and neuroscientific prediction. *Cognition* 133(2):502-516.
46. Sirigu A*, et al.* (2004) Altered awareness of voluntary action after damage to the parietal cortex. *Nature Neuroscience* 7(1):80-84.
47. Frankfurt HG (1971) Freedom of the Will and the Concept of a Person. *The journal of philosophy* 68(1):5-20.
48. Hobbes T (1994) *Leviathan: with selected variants from the Latin edition of 1668* (Hackett Publishing Company).
49. Wolf S (1990) *Freedom within reason* (Oxford University Press).
50. Deutschländer R, Pauen M, & Haynes J-D (2017) Probing folk-psychology: Do Libet-style experiments reflect folk intuitions about free action? *Consciousness and Cognition* 48:232-245.
51. Strawson G (1994) The impossibility of moral responsibility. *Philosophical Studies* 75(1):5-24.
52. Fischer JM (1999) Recent work on moral responsibility. *Ethics* 110(1):93-139.
53. Sinnott-Armstrong W (2014) *Moral Psychology: Free Will and Moral Responsibility* (Mit Press).
54. Roskies AL (2012) How does the neuroscience of decision making bear on our understanding of moral responsibility and free will? *Current opinion in neurobiology* 22(6):1022-1026.
55. Maoz U & Yaffe G (2016) What does recent neuroscience tell us about criminal responsibility? *Journal of Law and the Biosciences* 3(1):120-139.
56. Sanfey AG, Rilling JK, Aronson JA, Nystrom LE, & Cohen JD (2003) The neural basis of economic decision-making in the ultimatum game. *Science* 300(5626):1755-1758.
57. Wallis JD & Miller EK (2003) Neuronal activity in primate dorsolateral and orbital prefrontal cortex during performance of a reward preference task. *Eur J Neurosci* 18(7):2069-2081.
58. Ramnani N & Owen AM (2004) Anterior prefrontal cortex: insights into function from anatomy and neuroimaging. *Nature Reviews Neuroscience* 5(3):184-194.
59. Wallis JD (2007) Orbitofrontal cortex and its contribution to decision-making. *Annu. Rev. Neurosci.* 30:31-56.
60. Bush G, Luu P, & Posner MI (2000) Cognitive and emotional influences in anterior cingulate cortex. *Trends in cognitive sciences* 4(6):215-222.
61. Carter CS*, et al.* (1998) Anterior cingulate cortex, error detection, and the online monitoring of performance. *Science* 280(5364):747-749.
62. Krieghoff V, Waszak F, Prinz W, & Brass M (2011) Neural and behavioral correlates of intentional actions. *Neuropsychologia* 49(5):767-776.
63. Brass M & Haggard P (2008) The what, when, whether model of intentional action. *The Neuroscientist* 14(4):319.
64. Hughes G, Schütz-Bosbach S, & Waszak F (2011) One action system or two? Evidence for common central preparatory mechanisms in voluntary and stimulus-driven actions. *Journal of Neuroscience* 31(46):16692-16699.

65. Maoz U*, et al.* (2013) Predeliberation activity in prefrontal cortex and striatum and the prediction of subsequent value judgment. *Front Neurosci* 7:225.
66. Junghofer M, Elbert T, Tucker D, & Rockstroh B (2000) Statistical control of artifacts in dense array EEG/MEG studies. *Psychophysiology* 37(04):523-532.
67. Keren AS, Yuval-Greenberg S, & Deouell LY (2010) Saccadic spike potentials in gamma-band EEG: characterization, detection and suppression. *Neuroimage* 49(3):2248-2263.
68. Croft RJ & Barry RJ (2000) Removal of ocular artifact from the EEG: a review. *Neurophysiologie Clinique/Clinical Neurophysiology* 30(1):5-19.
69. Elbert T, Lutzenberger W, Rockstroh B, & Birbaumer N (1985) Removal of ocular artifacts from the EEG—a biophysical approach to the EOG. *Electroencephalography and clinical neurophysiology* 60(5):455-463.
70. Shan Y, Moster ML, & Roemer RA (1995) The effects of time point alignment on the amplitude of averaged orbital presaccadic spike potential (SP). *Electroencephalography and clinical neurophysiology* 95(6):475-477.

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Neural precursors of decisions that matter—an ERP study of deliberate and arbitrary choice

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<u>Supplementary material</u>

Supplementary Results

In this section, we show that differences in reaction time between the conditions, eye movements, the hand with which the subjects executed the movement, and subjects' consistency scores all do not drive our effect. We further show that the RPs in arbitrary decisions are not a reflection of a stimulus-locked potentials and are not due to baseline considerations as part of the analysis of the difference in reaction times.

<u>Differences in reaction times (RT) between conditions, including stimulus-locked potentials and baselines, do not drive the effect</u>

RTs in deliberate decisions were typically more than twice as long as RTs in arbitrary decisions. We therefore wanted to rule out the possibility that the absence of RP in deliberate decisions stemmed from the difference in RTs between the conditions. We carried out four analyses for this purpose. First, we ran a median split analysis—dividing the subjects into two groups based on their RTs: lower and higher than the median, for deliberate and arbitrary trials, respectively. We then ran the same analysis using only the faster subjects in the deliberate condition (M=1.91 s, SD=0.25) and the slower subjects in the arbitrary condition (M=1.25 s, SD=0.23). If RT length affects RP amplitudes, we would expect the RP amplitudes to be more similar between these two groups. However, though there were only half the data points, a similar pattern of results to those over the whole dataset was observed (Fig. S2A). Deliberate and arbitrary decisions were still reliably different ($F_{(1,17)}=5.22$, p=0.03), with significant RPs found in arbitrary (easy: $t(8)=4.57$, p=0.0018; hard: $t(8)=4.09$, p=0.0035), but not deliberate (easy: $t(8)=1.92$, p=0.09; hard: $t(8)=0.63$, p=0.54) decisions. In addition, the RPs for arbitrary decisions were not significantly different between the subjects with above-median RTs and the entire population for the easy or hard conditions (easy: $t(25)=0.14$, p>0.5; hard: $t(25)=0.56$, p>0.5). Similarly, the RPs for deliberate decisions were not significantly different between the subjects with below-median RTs and the entire population for the easy or hard conditions (easy: $t(25)=-0.34$, p>0.5; hard: $t(25)=0.17$, p>0.5). This suggest that RTs do not reliably affect Cz activation for deliberate or arbitrary decisions in our results.

Second, we regressed the difference between RPs in deliberate and arbitrary decisions (averaged over the last 500 ms before response onset) against the difference between the RTs in these two conditions for each subject (Fig. S2B). Again, if RT length affects RP amplitudes, we would expect differences between RTs in deliberate and arbitrary conditions to correlate with differences between RPs in the two conditions. But no correlation was found between the two measures (r=0.22, $t(17)=0.86$, p=0.4). We further tried regressing the RP differences on RT differences. The regression did not produce any reliable relation between RT and RP differences (regression line: y = 0.54 [CI -0.8, 1.89] x - 0.95 [CI -2.75, 0.85]; the R^2 was very low, at 0.05

(as expected from the r value above), and, as the confidence intervals suggest, the slope was not significantly different from 0, $F(1,17)=0.74$, $p=0.4$).

A third concern that could relate to the RT differences among the conditions is that the RP in arbitrary blocks might actually be some potential evoked by the stimuli (i.e., the presentations of the two causes), specifically in arbitrary blocks, where the RTs are shorter (and thus stimuli-evoked effects could still affect the decision). In particular, a stimulus-evoked potential might just happen to bear some similarity to the RP when plotted locked to response onset. To test this explanation, we plotted the potentials in all conditions, locked to the onset of the stimulus (Fig. S3A). We also plotted the response-locked potentials across an expanded timecourse for comparison (Fig. S3B). If the RP-like shape we see in Figs. 3A and S3B is due to a stimulus-locked potential, we would expect to see the following before the 4 mean response onset times (indicated by vertical lines at 0.98 and 1.00, 2.13, and 2.52 s for arbitrary easy, arbitrary hard, deliberate easy, and deliberate hard, respectively) in the stimulus-locked plot (Fig. S3A): Consistent potentials, which precede the mean response times, that would further be of a similar shape and magnitude to the RPs found in the decision-locked analysis in the arbitrary condition (though potentially more smeared for stimulus locking). We thus calculated a stimulus-locked version of our ERPs, using the same baseline (Fig. S3A). As the comparison between Fig. S3A and S3B clearly show, no such consistent potentials are found before the 4 response times, nor are these potentials similar to the RP in either shape or magnitude (their magnitudes are at the most around 1μV, while the RP magnitudes we found are around 2.5 μV; Figs. 3A, S3B). This analysis thus suggests that it is unlikely that a stimulus-locked potential drives the RP we found.

A fourth concern is that the differences in RTs may have an additional effect on the results: Because the main baseline period we used thus far was 1 to 0.5 s before stimulus onset, the duration from the baseline to the decision also varied widely between the conditions. To make sure this difference in temporal distance between the baseline period and the response to which the ERPs were locked does not drive our results, we recalculated the potentials for all conditions with a *response-locked* baseline of -1 to -0.5 s (Fig. S3C)—the same baseline we used for the Bayesian analysis. The rationale behind this choice of baseline was to have the time that elapsed from baseline to response onset be the same across all conditions. As is evident in Fig. S3C, the results for this new baseline were very similar to those for the stimulus-locked baseline we used before. Focusing again on the -0.5 to 0 s range before response onset for our statistical tests, we found a clear RP in arbitrary decisions, yet RP amplitude was not significantly different from 0 in deliberate decisions (Fig. S3C; ANOVA $F(1,17)=12.09$, $p=0.003$ for the main effect of decision type; in t-tests against 0, corrected for multiple comparisons, an effect was only found for arbitrary decisions (hard: $t(17)=4.13$, $p=0.0007$; easy: $t(17)=4.72$, $p=0.0002$) and not for deliberate ones (hard: $t(17)=0.38$, $p>0.5$; easy: $t(17)=1.13$, $p=0.27$). This supports the notion that the choice of baseline does not strongly affect our main results. Taken together, these results provide strong evidence against the claim that the differences in RPs stem from or are affected by the differences in RTs between the conditions.

Eye movements do not affect the results

Though ICA was used to remove blink artifacts and saccades (see Methods), we wanted to make sure our results do not stem from differential eye movement patterns between the conditions. We therefore computed a saccade-count metric (SC; see Methods) for each trial for all subjects. Focusing again on the last 500 ms before response onset, we computed mean (± s.e.m.) SC values of 1.65±0.07 and 1.67±0.06 saccades for easy and hard deliberate decisions, respectively, and 1.69±0.07 and 1.73±0.07 saccades for easy and hard arbitrary decisions, respectively. We found no reliable differences between the number of saccades during deliberate and arbitrary trials (ANOVA $F_{(1,17)}=2.56$, $p=0.13$ for the main effect of decision type).

We further investigated potential effects of saccades by running a median-split analysis— dividing the trials for each subject into two groups based on their SC score: lower and higher than the median, for deliberate and arbitrary trials, respectively. We then ran the same analysis using only the trials with more saccades in the deliberate condition (SC was 2.02±0.07 and 2.04±0.07 for easy and hard, respectively) and those with less saccades for the arbitrary condition (SC was 1.33±0.07 and 1.31±0.08 for easy and hard, respectively). If the number of saccades affects RP amplitudes, we would expect that the differences in RPs between arbitrary and deliberate trials will diminish, or even reverse (as now we had more saccades in the deliberate condition). However, though there were only half the data points for each subject in each condition, a similar pattern of results to those over the whole dataset was observed (Fig. S4): Deliberate and arbitrary decisions were still reliably different within the median-split RPs ($F_{(1,17)}=16.70$, $p<0.001$), with significant RPs found in arbitrary (easy: $t_{(17)}=4.79$, $p=0.002$; hard: $t_{(17)}=5.77$, $p<0.001$), but not deliberate (easy: $t_{(17)}=0.90$, $p=0.38$; hard: $t_{(17)}=0.30$, $p>0.5$) decisions. In addition, we compared the RP data across all the trials with the median-split RP data above. No significant differences were found for arbitrary decisions (easy: $t_{(17)}=1.02$, $p=0.32$; hard: $t_{(17)}=0.75$, $p=0.46$) or for deliberate decisions (easy: $t_{(17)}=1.63$, $p=0.12$; hard: $t_{(17)}=1.47$, $p=0.16$). Taken together, the analyses above provide strong evidence against the involvement of eye movements in our results.

Testing other factors that could potentially account for the results

We further tested whether our results depend on the hand with which subjects responded. Again, the same pattern of results persisted when separating left-hand button presses from right-hand ones (Fig. S5). Second, we took a closer look at subjects' behavior in the easy arbitrary condition, where some subjects had a consistency score that was further above 0.5 (chance) than others. It seems like those subjects had a greater difficulty ignoring their preferences, despite the instructions to do so. We therefore wanted to test to what extent the RP of those subjects was similar to the RPs of the other subjects. Focusing on the 8 subjects that had a consistency score above 0.55 (M=0.59, SD=0.03) and comparing their RPs to those of the 10 other subjects (consistency M=0.50, SD=0.06) in easy arbitrary trials, we found no reliable differences ($t_{(16)}=0.94$, $p=0.36$). This is not surprising, as the mean consistency score of these subjects – though indeed higher than chance – was still far below their consistency score for easy deliberate decisions (M=0.99, SD=0.02).

Supplementary Discussion

In this supplementary discussion, we introduce two alternative explanations of our results, and argue that they are less plausible than the interpretation we suggested in the main text. First, one could claim that—in the deliberate condition only—the NPO names act as a cue, thereby turning what we term internal, deliberate decisions into no more than simple responses to external stimuli. Under this account, if the preferred NPO is on the right, it is immediately interpreted as "Press right". It would therefore follow that the absence of decisions in deliberate trials results in the absence of the RP in those trials. However, the reaction time and consistency values that we observed provide evidence against this interpretation. We found longer reaction times for hard-deliberate decisions than for easy-deliberate ones (2.52 versus 2.13 s, on average, respectively; Fig. 2) and higher consistencies with the initial ratings for easy-deliberate decisions than for hard-deliberate decisions (0.99 versus 0.83, on average, respectively; Fig. 2). If the NPO names acted as mere cues, we would have expected no differences between reaction times or consistencies for easy- and hard-deliberate decisions. In addition, there were 50 different causes in the first part of the experiment. So, it is highly unlikely that subjects could memorize all 1225 pairwise preferences among these causes and simply transform any decision between a pair of causes into a stimulus instructing to press left or right.

Another, alternative interpretation of our results might be that subjects engage in (unconscious) deliberation also during arbitrary decisions (1). If so, the RP in the arbitrary decisions might reflect the extra effort required to break the symmetry under this condition. However, this interpretation entails a longer reaction time for arbitrary than for deliberate decisions, because of the heavier cognitive load, which is the opposite of what we found (Fig. 2A). We would also expect the simpler deliberation in arbitrary-easy trials to result in a shorter reaction-time than that of arbitrary-hard. But this is not what we find (Fig. 2A).

Supplementary Materials and Methods

Sample size and exclusion criteria

In a preliminary study that we ran, we found that a sample size of at least 16 subjects results in a clear, averaged RP. So, we decided beforehand to collect 20 subjects for this study, taking into account that some could be excluded as they would not meet the following predefined inclusion criteria: at least 30 trials per experimental condition remaining after artifact rejection; and averaged RTs (across conditions) which deviated by 3 or less standard deviations from the group mean. Subjects were informed about the overall number of subjects that would participate in the experiment when the NPO lottery was explained to them (see below). So, we had to finalize the overall number of subjects who would participate in the study—but not necessarily the overall number of subjects whose data would be part of the analysis—before the experiment began. After completing data collection, we ran only the EEG preprocessing and behavioral-data analysis to test each subject against the exclusion criteria. This was done before we looked at the data with respect to our hypothesis or research question. Two subjects did not meet the inclusion criteria: the data of one subject (#18) suffered from poor signal quality which resulted in less

than 30 trials after artifact rejection; another subject (#12) had RTs longer than 3 standard deviations from the mean. All analyses were thus run on the 18 remaining subjects.

We further excluded single trials in which subjects pressed the wrong button as well as trials where subjects' RTs were less than 200 ms, more than 10s, or more than 3 standard deviations away from that subject's mean in that condition (mean number of excluded trials =7.17, SD=2.46, which are 1.99% of the trials).

Subjects

Twenty healthy subjects participated in the study. They were California Institute of Technology (Caltech) students as well as members of the Pasadena community. All subjects had reported normal or corrected-to-normal sight and no psychiatric or neurological history. They volunteered to participate in the study for payment ($20 per hour). Subjects were prescreened to include only participants who were socially involved and active in the community (based on the strength of their support of social causes, past volunteer work, past donations to social causes, and tendency to vote). The data from 18 subjects was analyzed; two subjects were excluded from our analysis (see Study Design above). The experiment was approved by Caltech's Institutional Review Board, and informed consent was obtained from all participants after the experimental procedures were explained to them.

Stimuli and apparatus

Subjects sat in a dimly lit room. The stimuli were presented on a 21" Viewsonic G225f (20" viewable) CRT monitor with a 60-Hz refresh rate and a 1024×768 resolution using Psychtoolbox version 3 and Mathworks Matlab 2014b (2, 3). They appeared with a gray background (RGB values: [128, 128,128]). The screen was located 60 cm away from subjects' eyes. Stimuli included names of 50 real, non-profit organizations (NPOs). Twenty organizations were consensual (e.g., the Cancer Research Institute, or the Hunger project), and thirty were more controversial: we chose 15 causes that were widely debated (e.g., pro/anti guns, pro/anti abortions), and selected one NPO that supported each of the two sides of the debate. This was done to achieve variability in subjects' willingness to donate to the different NPOs. In the main part of the experiment, succinct descriptions of the causes (e.g., pro-marijuana legalization, pro-child protection; for a full list of NPOs and causes see Table S2) were presented in black Comic Sans MS.

Extended Experimental Procedure

In the first part of the experiment, subjects were presented with each NPO separately. They were instructed to rate how much they would like to support that NPO with a $1000 donation on a scale of 1 ("I would not like to support this NPO at all) to 7 ("I would very much like to support this NPO"). No time pressure was put on the subjects, and they were given access to the website of each NPO to give them the opportunity to learn more about the NPO and the cause it supports.

After the subjects finished rating all NPOs, the main experiment began. In each block of the experiment, subjects made either deliberate or arbitrary decisions. Subjects were clearly

instructed to report their decisions as soon as they made them for both arbitrary and deliberate decisions, and they were asked to place their fingers on the corresponding keys to make sure they could respond as quickly as possible. Two succinct cause descriptions, representing two actual NPOs, were presented in each trial (Fig. 1). In deliberate blocks, subjects were instructed to choose the NPO to which they would like to donate $1000 by pressing the <Q> or <P> key on the keyboard, for the NPO on the left or right, respectively, as soon as they decided. Subjects were informed that at the end of each block one of the NPOs they chose would be randomly selected to advance to a lottery. Then, at the end of the experiment, the lottery will take place and the winning NPO will receive a $20 donation. In addition, that NPO will advance to the final, inter-subject lottery, where one subject's NPO will be picked randomly and will be given a $1000 donation. It was stressed that the donations were real and that no deception was used in the experiment. To persuade the subjects that the donations are real, we presented a signed commitment to donate the money, and promised to send them the donation receipts after the experiment. Thus, subjects knew that in deliberate trials, every choice they made was not hypothetical, and could potentially lead to an actual $1020 donation to their chosen NPO.

Arbitrary trials were identical to deliberate trials except for the following crucial differences. Subjects were told that, at the end of each block, the pair of NPOs in one randomly selected trial would advance to the lottery together. And, if that pair wins the lottery, both NPOs would receive $10 (each). Further, the NPO pair that would win the inter-subject lottery would receive a $500 donation each. Hence it was stressed to the subjects that there was no reason for them to prefer one NPO over the other in arbitrary blocks, as both NPOs would receive the same donation regardless of their button press. Subjects were told to therefore simply press either <Q> or <P> as soon as they decided to do so.

Thus, while subjects' decisions in the deliberate blocks were meaningful and consequential, their decisions in the arbitrary blocks had no impact on the final donations that were made. In these trials, subjects were further urged not to let their preferred NPO dictate their response. Note that we did not ask subjects to report their "W-time" (time of consciously reaching a decision), because this measure was shown to rely on neural processes occurring after response onset (4) and to potentially be backward inferred from movement time (5). Even more importantly, clock monitoring was demonstrated to have an effect on RP size (6), so it could potentially confound our results (7).

In addition, we manipulated decision difficulty (Easy/Hard) throughout the experiment, randomly intermixed within each block. Decision difficulty was determined based on the rating difference between the two presented NPOs. NPO pairs with 1 or at least 4 rating-point difference were designated hard or easy, respectively. Based on each subject's ratings, we created a list of NPO pairs, half of each were easy choices and the other half hard choices.

Each block started with an instruction written either in dark orange (Deliberate: "In this block choose the cause to which you want to donate $1000") or in blue (Arbitrary: "In this block both causes may each get a $500 donation regardless of the choice"). The background color of the screen was light gray here and throughout the experiment. Short-hand instructions appeared at the top of the screen throughout the block in the same colors as that block's initial instructions; Deliberate: "Choose for $1000" or Arbitrary: "Press for $500 each" (Fig. 1).

Each trial started with a screen that was blank except for a centered, black fixation cross. The fixation screen was on for a duration drawn from a uniform distribution between 1 and 1.5 s. Then, the two cause-descriptions appeared on the left and right side of the fixation cross (left/right assignments were randomly counterbalanced), and remained on the screen until the subjects reported their decisions with a key press—<Q> or <P> on the keyboard for the cause on the left or right, respectively. The cause corresponding to the pressed button then turned white for 1 s, and a new trial started immediately. If subjects did not respond within 20 s, they received an error message and were informed that if this trial would be selected for the lottery, no NPO would receive a donation. However, this did not happen for any subject on any trial.

To assess the consistency of subjects' decisions during the main experiment with their ratings in the first part of the experiment, subjects' choices were coded in the following way: each binary choice in the main experiment was given a consistency grade of 1, if subjects chose the NPO that was rated higher in the rating session, and 0 if not. Then a consistency grade was calculated as the mean over all the choices. A consistency grade of 1 indicates perfect consistency with one's ratings, 0 – perfect inconsistency, and 0.5 – chance performance.

We wanted to make sure subjects were carefully reading and remembering the causes also during the arbitrary trials to better equate memory load, attention, and other cognitive aspects between deliberate and arbitrary decisions—except those aspects directly associated with the decision type, which was the focus of our investigation. We therefore randomly interspersed 36 memory catch-trials throughout the experiment (thus more than one catch trial could occur per block). On such trials, four succinct descriptions of causes were presented and subjects had to select the one that appeared in the previous trial. A correct or incorrect response added or subtracted 50 cents from their total, respectively. (Subjects were informed that if they reached a negative balance, no money will be deducted off their payment for participation in the experiment.) Thus, subjects could earn $18 more for the experiment, if they answered all memory test questions correctly. Subjects typically did well on these memory questions, on average erring in 2.5 out of 36 memory catch trials (7% error) and gaining an additional $16.75 (SD=3.19).

Statistical Analysis

Differences greater than expected by chance were assessed using two-way ANOVAs with decision type (deliberate, arbitrary) and decision difficulty (easy, hard), using IBM SPSS statistics, version 24. For both RP and LRP signals, the mean amplitude from 500 ms before to button-press onset were used for the ANOVAs. Greenhouse–Geisser correction was never required as sphericity was never violated (8).

Trend analysis on all subjects' data (Fig. 3B) was carried out by regressing the voltage for every subject against time for the last 1000 ms before response onset using first-order polynomial linear regression. We used every 10th time sample for the regression (i.e., the 1st, 11th, 21st, 31st samples, and so on) to conform with the individual-subject analysis (see below). For the individual-subject analysis, the voltage on all trials was regressed against time in the same manner (i.e., for the last 1000 ms before response onset and using first-order polynomial linear regression). As individual-trial data is much noisier than the mean over all trials in each subject, we opted for standard robust-regression using iteratively reweighted least squares (implemented

using the *robustfit()* function in Mathworks Matlab). The iterative robust-regression procedure is time consuming. So, we used every 10th time sample instead of every sample to make the procedure's run time manageable.

Model and Simulations

All simulations were performed using Mathworks Matlab 2014b. The model was devised off the one proposed by Schurger and colleagues (9). They built a drift-diffusion model (10, 11) for arbitrary decisions only, which included a leaky stochastic accumulator (with a threshold on its output) and a time-locking/epoching procedure. Their model amounted to iterative numerical integration of the differential equation

$$\delta x_i = (I - kx_i)\Delta t + c\xi_i\sqrt{\Delta t}$$

where I is the drift rate, k is the leak (exponential decay in x), ξ is Gaussian noise, and c is a noise-scaling factor (we used $c = 0.05$). Δt is the discrete time step used in the simulation (we used $\Delta t = 0.001$, similar to our EEG sampling rate). Here I represents a constant urgency to respond that is inherent in the demand characteristics of the task, evidenced by the fact that no subject took more than 20 s to decide on any trial. The leak term, k, ensures that the model would not be too linear; i.e., it prevented the urgency from setting up a linear trajectory for the accumulator toward the threshold. Hence, due to the leak term, doubling the magnitude of the threshold would make the accumulator rarely reach the threshold, instead of reaching it in roughly twice the amount of time (up to the noise term) had we not introduced a leak term.

Our model accounted for both arbitrary and deliberate decisions and was built to fit our empirical results, based on two Schurger-like components. The first one accumulated activity that drove arbitrary decisions (i.e., random fluctuations (9)). The second component drove deliberate decisions based on subjects' values associated with the decision alternatives. Henceforth we term these *Noise* and *Value* components for ease of description. Our model used its Noise component for arbitrary decisions and its Value one for deliberate decisions.

Schurger and colleagues modeled only the decision when to move. But our subjects decided both when and which hand to move. So, we had to extend the Schurger model in that respect as well. We did this using a race-to-threshold mechanism between the two decision alternatives. In our paradigm, the difference in rating was either 1 (hard decisions) or 4-6 (easy decisions; see "Procedure & Experimental Design" in Methods), so there was always an alternative that was ranked higher than the other. Choosing the higher or lower alternative was termed a congruent or incongruent choice with the initial ratings, respectively.

Using a parameter sweep, we found the values of the thresholds, urgency, and leak that best fit our average empirical reaction times for (easy, hard) x (deliberate, arbitrary) decisions as well as our empirical consistency ratios for those 4 decision types. The model's reaction time was defined as the overall time (where each step took $\Delta t = 0.001$ s) that it took until the first threshold crossing in the race-to-threshold pair. We used the same threshold value of 0.15 and leak value of $k=0.5$ for all model types. The only parameter that was modulated across (deliberate,

arbitrary) x (easy, hard) decisions x (congruent, incongruent) decision alternatives was the urgency, I (Table S1). These parameters were then fixed when fitting the simulated Cz activity across all conditions.

Each simulation consisted of either 120 runs of the model, the same as the number of empirical trials per condition (Fig. 4C), or 10000 runs of the model for a smoother reaction-time distribution for the model (Fig. 4B). For each run of the model, we identified the first threshold crossing point and extracted the last second (1000 steps) before the crossing in each run. If the first crossing was earlier than sample no. 1,000 by $n > 0$ samples, we padded the beginning of the epoch with n null values (NaN or "not-a-number" in Matlab). These values did not contribute to the average across simulated trials, so the simulated average RP became noisier at earlier time points in the epoch. Our model was therefore similarly limited to the Schurger model in its inability to account for activity earlier than the beginning of the trial. (Fig. 3C).

Figure S1: Individual-subjects RPs. Six examples of for individual subjects' RPs (ERPs in the Cz electrode aligned to response onset) for deliberate decisions (in red) and arbitrary ones (in blue).

Figure S2: Relations between RTs and RPs. (A) The subjects with above-median RTs for arbitrary decisions (in blue) and below-median RTs for deliberate decisions (in red), show the same pattern that was found in the main analysis (compare Fig. 3A in main text). **(B)** A regression of the difference between the RPs versus the difference between the RTs for deliberate and arbitrary decisions for each subject. The equation of the regression line is $y = 0.54$ [CI -0.8, 1.89] $x - 0.95$ [CI -2.75, 0.85]. The R^2 is 0.05. One subject, #7, had an RT difference between deliberate and arbitrary decisions that was more than 6 interquartile ranges (IQRs) away from the median difference across all subjects. That same subject's RT difference was also more than 5 IQRs higher than the 75th percentile across all subjects. That subject was therefore designated an outlier and removed from the regression analysis.

Figure S3: Stimulus- and response-locked Cz-electrode ERPs with different baselines and overlaid events. (A) Stimulus-locked waveforms including the trial onset range, baseline period, and mean reaction times for all four experimental conditions. **(B)** Response-locked waveforms with mean stimulus onsets for all four conditions as well as the offset of the highlighting of the selected cause and the start of the next trial. **(C)** Same potentials and timeline as Fig. 3A, but with a *response-locked* baseline of -1 to -0.5 s—the same baseline used for our Bayesian analysis.

Figure S4: Relations between eye movements and RPs. Trials with below-median number of saccades for arbitrary decisions (in blue) and above-median number of saccades for deliberate decisions (in red), show the same pattern that was found in the main analysis (compare Fig. 3 in main text).

Figure S5: RPs for left-hand and right-hand trials. The RP for trials where subjects moved their left hand (left panel) and their right hand (right panel) separately. Dividing the trials into left-hand and right-hand ones results in only half the number of trials per subject, compared with Fig. 3. So, easy and hard decisions were pooled together to get back to approximately the same number of trials. Notably, the main pattern of results persists for each type of trials (right/left responses), where an RP is found for arbitrary decisions only.

Figure S6: Lateralized readiness potential. The lateralized readiness potential (LRP) for deliberate and arbitrary, easy and hard decisions. No difference was found between the conditions (ANOVA all Fs<1).

Figure S7: Model runs. The first 20 model runs for all four decision conditions. The integration threshold, at 0.15, is designated by a dashed line in all decision conditions. Here t = 0 s designates the beginning of the model's run.

Table S1: Values of the urgency parameter across decision types. Values of the urgency parameter, *I*, in our model across (deliberate, arbitrary) x (easy, hard) decisions x (congruent, incongruent) decision alternatives.

Urgency (*I*) values	Congruent		Incongruent	
	Easy	Hard	Easy	Hard
Deliberate	0.0400	0.1010	0.0228	0.0000
Arbitrary	0.1650	0.1648	0.1650	0.1566

Table S2: NPO names and causes acronyms

NPO	Cause	NPO website
Consensual NPOs		
American Society on Aging	Pro Quality of Life for the Elderly	http://asaging.org/
Conservation Fund	Pro Environment protection'	http://www.conservationfund.org/
Bill & Melinda Gates Foundation	Pro Education	http://www.gatesfoundation.org/
Global Fund for Women	Pro Women's Rights	https://www.globalfundforwomen.org/
The Hunger Project	Pro Hunger Relief	https://www.thp.org/
Oxfam International	Pro Poverty & Disaster Relief	http://www.oxfam.org/
World Wild Life Fund (WWF)	Pro Species Conservation	http://worldwildlife.org/
Cancer Research Institute	Pro Cancer Research	http://www.cancerresearch.org/
Habitat for Humanity	Pro Housing for All	http://www.habitat.org/
Reading is Fundamental	Pro Advancement of Literacy	http://www.rif.org/
International Institute for Conservation of Historic and Artistic Works	Pro Culture & Arts Preservation	https://www.iiconservation.org/
Big Brothers and Big Sisters of America	Pro Youth Development	http://www.bbbs.org/site/c.9iILI3NGKhK6F/ b.5962335/k.BE16/Home.htm

United Nations Children's Fund (UNICEF)	Pro Child Protection	http://www.unicef.org/
Doctors without Borders (Medecins sans frontieres)	Pro Disaster Medical Care	http://www.msf.org/
Soldiers' Angels	Pro Veterans & Military	http://www.soldiersangels.org/heroes/index.php
Disability Rights International	Pro Disabilities Rights	http://www.disabilityrightsintl.org/
National Crime Prevention Council (NCPC)	Pro Crime Prevention	http://www.ncpc.org/
Amnesty International	Pro Human Rights	https://www.amnesty.org/
Peace Corps	Pro Peace & Development	http://www.peacecorps.gov/
World Health Organization	Pro World Health	http://www.who.int/en/
Controversial NPOs		
Planned Parenthood	Pro Abortion & Family Planning	http://www.plannedparenthood.org/
Pro-Life Alliance	Anti Abortion & Family Planning	http://www.prolifealliance.com/
Human Rights Campaign	Pro LBGTQ Rights	http://www.hrc.org/
National Organization for Marriage	Anti LBGTQ Rights	https://www.nationformarriage.org/
Stem for Life Foundation	Pro Stem Cell Research	http://www.stemforlife.org/
Christian Dental & Medical Association	Anti Stem Cell Research	http://www.cmda.org/
Greenpeace	Pro Action Against Climate Change	http://www.greenpeace.org/international/en/
Global Climate Scam	Anti Action Against Climate Change	http://www.globalclimatescam.com/
National Association for Gun Rights	Pro Gun Rights	http://www.nationalgunrights.org/
Coalition to Stop Gun Violence	Pro Gun Control	http://csgv.org/

American Gas Association	Pro Fracking for Natural Gas	http://www.aga.org/Pages/default.aspx
Americans Against Fracking	Anti Fracking for Natural Gas	http://www.americansagainstfracking.org/
StandWithUs (Israel)	Pro Israel	http://www.standwithus.com/
Palestinian Centre for Human Rights	Pro Palestine	http://www.pchrgaza.org/portal/en/
National Organization for the Reform of Marijuana Laws	Pro Marijuana Legalization	http://norml.org/
Citizens Against Legalizing Marijuana	Anti Marijuana Legalization	http://www.calmca.org/
Understanding Animal Research	Pro Scientific Experiments on Animals	http://www.understandinganimalresearch.org.uk/
International Association Against Painful Experiments on Animals	Anti Scientific Experiments on Animals	http://www.iaapea.com/
Federation for American Immigration Reform	Pro Immigration Reform	http://www.fairus.org/
American Immigration Control	Anti Immigration Reform	http://www.immigrationcontrol.com/
Human Cloning Foundation	Pro Human Cloning	http://www.humancloning.org/
Americans to Ban Cloning	Anti Human Cloning	http://www.cloninginformation.org/
Americans United for Separation of Church and State	Pro Separation of Church & State	https://www.au.org/
Christian Coalition of America	Anti Separation of Church & State	http://www.cc.org

Death with Dignity National Center	Pro Euthanasia (Assisted Suicide)	http://www.deathwithdignity.org/
Euthanasia Prevention Coalition	Anti Euthanasia (Assisted Suicide)	http://www.epcc.ca/
The Alliance for Better Foods	Pro Genetically Modified Foods	http://www.betterfoods.org/
Non-GMO Project	Anti Genetically Modified Foods	http://www.nongmoproject.org/
Answers in Genesis	Pro Creationism Teaching	https://answersingenesis.org
National Center for Science Education	Pro Evolution Teaching	http://ncse.com/

References

1. Tusche A, Bode S, & Haynes J-D (2010) Neural responses to unattended products predict later consumer choices. *Journal of neuroscience* 30(23):8024-8031.
2. Brainard DH (1997) The psychophysics toolbox. *Spatial vision* 10:433-436.
3. Pelli DG (1997) The VideoToolbox software for visual psychophysics: Transforming numbers into movies. *Spatial vision* 10(4):437-442.
4. Lau H, Rogers R, & Passingham R (2007) Manipulating the experienced onset of intention after action execution. *Journal of cognitive neuroscience* 19(1):81-90.
5. Banks WP & Isham EA (2009) We infer rather than perceive the moment we decided to act. *Psychological Science* 20(1):17.
6. Miller J, Shepherdson P, & Trevena J (2011) Effects of Clock Monitoring on Electroencephalographic Activity. *Psychological Science* 22(1):103-109.
7. Maoz U*, et al.* (2015) On reporting the onset of the intention to move. *Surrounding Free Will: Philosophy: Psychology, Neuroscience*, ed Mele AR (Oxford University Press).
8. Picton T*, et al.* (2000) Guidelines for using human event-related potentials to study cognition: recording standards and publication criteria. *Psychophysiology* 37(02):127-152.
9. Schurger A, Sitt JD, & Dehaene S (2012) An accumulator model for spontaneous neural activity prior to self-initiated movement. *Proceedings of the National Academy of Sciences* 109(42):E2904-E2913.
10. Ratcliff R (1978) A theory of memory retrieval. *Psychological Review* 85(2):59-108.
11. Usher M & McClelland JL (2001) The time course of perceptual choice: the leaky, competing accumulator model. *Psychological review* 108(3):550.