Transcranial ultrasound selectively biases decision-making in primates

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Keywords: noninvasive, neuromodulation, macaque, choice behavior

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Abstract

Transcranial focused ultrasound has the promise to evolve into a transformative noninvasive way to modulate activity of neuronal circuits deep in the brain. The approach may enable systematic and causal mapping of how individual brain circuits are involved in specific behaviors and behavioral disorders. Previous studies demonstrated neuromodulatory potential, but the effect polarity, size, and spatial specificity have been difficult to assess. Here, we engaged non-human primates (*macaca mulatta*) in an established task that provides a well defined framework to characterize the neuromodulatory effects. In this task, subjects decide whether to look at a right or a left target, guided by one the targets appearing first. Previous studies showed that excitation/inhibition of oculomotor circuits leads to contralateral/ipsilateral biases in this choice behavior. We found that brief, low-intensity ultrasound stimuli (300 ms, 0.6 MPa, 270 kHz) delivered to the animals' left/right frontal eye fields bias the animals' decisions to the right/left visual hemifield. The effect was modest, about on the order of that produced when injecting moderate amounts of potent neuromodulatory drugs into the same regions in this task. The polarity of the effects suggested a neuronal excitation within the stimulated regions. No effects were observed when we applied the same stimuli to control brain regions not involved in oculomotor target selection. Together, using an established paradigm, we found that transcranial ultrasound is capable of modulating neurons to the extent of biasing choice behavior of non-human primates. A demonstration of tangible, double-dissociated effects on behavior of primates constitutes a critical step toward applying this noninvasive neuromodulation method in investigations of how specific neural circuits are involved in specific behaviors or disease signs.

Introduction

Noninvasive and spatially specific approaches to modulating neuronal activity have the potential to revolutionize the diagnoses and treatments of a variety of brain disorders. One of such approaches, ultrasound, can be applied through the intact skull and skin and focused into tight regions deep in the human brain (Ghanouni et al., 2015). At the focus, ultrasound has been shown to modulate neural activity (Naor et al., 2016; Fini and Tyler, 2017; Kubanek, 2018; Tyler et al., 2018; Fomenko et al., 2018). By virtue of its noninvasiveness and spatial focus, the approach has a unique promise in modulating the activity of specific circuits in a systematic fashion. This could enable us, for the first time, to characterize the *causal* contribution of specific brain circuits to specific behaviors or behavioral disorders *in humans*. In order to bring the approach into clinics, we need to be able to establish three critical aspects of the neuromodulatory effects. First, it is crucial to characterize the polarity of the neuromodulatory effects—whether neurons are excited or inhibited by ultrasound. Second, we need to determine how strong the effects are. In order for the method to be useful for mapping brain function in a causal manner, the effects on neurons must be strong enough to manifest in behavior. For example, if clinicians are to determine which brain nuclei underlie a patient's essential tremor, the neuromodulatory effects on a particular nucleus in question must be strong enough to yield measurable changes in the tremor amplitude. And third, it is critical to validate that the neuromodulatory effects are confined to the focal region of the ultrasound.

This information has been difficult to infer from the approaches and metrics used in previous studies. The bulk of work on ultrasonic neuromodulation has been performed in lightly anesthetized mice (Tufail et al., 2011; King et al., 2013; Kim et al., 2014; Mehić et al., 2014; Ye et al., 2016; Kamimura et al., 2016; Li et al., 2016; Sato et al., 2018). In these studies, applications of low-intensity stimuli to peri-motor regions often lead to visible movements of the limbs or other body parts. There have been concerns that the small size of the rodent brain, relative to the dimensions of the focal spot, results in reflections, standing waves, and consequently, artifactual effects (Guo et al., 2018; Sato et al., 2018). Within this debate, it has been argued that the neuromodulatory effects of ultrasound might be local (King et al., 2014; Mehić et al., 2014; Kamimura et al., 2016), but it has also been argued that these effects might be merely due to auditory or vestibular artifacts (Guo et al., 2018; Sato et al., 2018). Either way, the probability of eliciting movements on a given stimulation trial ("success rate") has been difficult to reproduce consistently and is strongly dependent on the kind and level of anesthesia (Naor et al., 2016). On the other hand, studies using larger mammals including sheep, macaques, and humans have shown effects on less direct metrics including EEG activity, MRI BOLD, or reaction time (Deffieux et al., 2013; Hameroff et al., 2013; Legon et al., 2014; Lee et al., 2015; Lee et al., 2016; Lee et al., 2016; Wattiez et al.,

2017; Legon et al., 2018). It has been difficult to judge from these studies how strong the effects are and in what direction they point because there has been no particular prediction framework within which to interpret these effects. In addition, it has not been clear how local the effects are (Lee et al., 2016).

Here, we characterize the size and polarity of the neuromodulatory effects of ultrasound using a well-established task (Oppenheim, 1885; Rorden et al., 1997; Ro et al., 2001; Schiller and Tehovnik, 2003; Kubanek et al., 2015) in awake behaving non-human primates (NHPs). In this task, a subject decides whether to look at a right or a left target, guided by one of the targets appearing slightly earlier than the other target. Previous studies showed that specific, neuroinhibitory or neuroexcitatory perturbations of specific visuomotor regions produce specific shifts in subjects' decisions regarding which target to choose. Thus, this prior research provides predictions of what behavior to expect if a neuromodulatory approach such as ultrasound is excitatory or inhibitory. Moreover, these studies enable us to gauge the size of the neuromodulatory effects from the magnitude of the behavior shift. Finally, the large brain of NHPs and hemispheric symmetry allows us to assess whether the neuromodulatory effects are specific to the stimulated regions.

Results

We engaged two macaque monkeys in a task that is often used in neurology to diagnose the impact of brain lesions such as those induced by stroke (Oppenheim, 1885; Rorden et al., 1997; Ro et al., 2001). In this paradigm (Fig. 1A), one visual target is shown in the left and one in the right visual hemifield, with a short, controlled delay between the onsets. Typically, in this task, healthy, normal subjects tend to look at the target that appeared first. Stroke or lesions of specific nodes of the oculomotor network, such as the frontal eye fields (FEF) or the lateral intraparietal area (LIP), strongly affect this behavior (Rorden et al., 1997; Ro et al., 2001; Schiller and Tehovnik, 2003; Wardak et al., 2004; Kubanek et al., 2015). These oculomotor circuits preferentially represent targets in their contralateral visual hemifield (Fig. 1B). As a consequence, when neurons in these circuits are affected by a stroke, the contralateral visual hemifield is underrepresented, and subjects preferentially decide to look at the ipsilesional target (Rorden et al., 1997; Ro et al., 2001). These effects are also observed for other kinds of neural perturbations, such as when neuromodulatory agents are injected into these regions. For example, when muscimol—a potent neuroinhibitory drug—is injected into left FEF of macaques (Schiller and Tehovnik, 2003), animals performing this task exhibit a strong ipsilateral—leftward bias (Fig. 1C, red). In contrast, injecting a drug with opposite, disinhibitory effects such as bicuculine into the same region (Schiller and Tehovnik, 2003) leads to an opposite, contralateral bias (Fig. 1C, blue). Analogous results

are obtained for neuromodulatory interventions into area LIP (Hanks et al., 2006; Schiller and Tehovnik, 2003; Wardak et al., 2004; Kubanek et al., 2015). This task therefore provides a well established framework that enables us to interpret the neuromodulatory effects of ultrasound—or any other intervention, for that matter. Excitatory interventions or stimuli bias subjects' decisions in the contralateral direction, whereas inhibitory interventions in the opposite direction.

We used this framework to evaluate the polarity and size of the effects of ultrasound on neurons (Fig. 1A). In a given session, ultrasound was applied to the animals' left or right FEF. Ultrasound was applied in blocks of 3-6 trials and was strictly interleaved with trials in which ultrasound was not applied to exclude potential effects of session time. The stimulus was applied while an animal was making his decision: 100 ms prior to the onset of the first target, and the stimulation lasted for 300 ms. Given that animals responded on average within 171 and 263 ms (monkey A and B, respectively), this perturbation influences a substantial portion of the decision-making process.

The ultrasound stimulus (Fig. 1D) had neuromodulatory parameters (0.6 MPa, 270 kHz, 300 ms duration, 500 Hz pulse repetition frequency, 50% duty cycle) that fell into commonly used ranges. We chose a relatively low carrier frequency of 270 kHz so that we could be confident that the stimulus would effectively penetrate the animals' skull (Deffieux et al., 2013). As a consequence of this choice, the stimulus was relatively broad as indicated by our measurements of free field pressure (Fig. 1D). Full width at half maximum pressure was 10.5 mm in the lateral dimension and 21.8 mm in the axial direction below the skull.

Animals showed typical choice behavior in this task, being sensitive to the difference in target onset times (Fig. 2A, black curves). As in previous studies (Rorden et al., 1997; Ro et al., 2001; Schiller and Tehovnik, 2003; Wardak et al., 2004; Kubanek et al., 2015), the earlier a target appeared before the consecutive target, the more likely the animal was to choose that target. Critically, transcranial ultrasound had a strong influence on this choice behavior (Fig. 2A, blue curves). In the trials in which ultrasound was applied to left FEF (left column), animals were more likely to choose the rightward target. The effect reversed polarity when ultrasound was applied to right FEF (right column): in this case, animals were more likely to choose the rightward target. Thus, these single session examples suggest that ultrasound biases the animals' decisions in the contralateral direction. Indeed, this effect was consistent across the individual sessions and monkeys (Fig. 2B). For each session, at the point of equal preference when no ultrasound was applied (see Materials and Methods), ultrasound targeting left FEF increased the animal's choices proportion of rightward choices (Fig. 2B, left). The effect was especially strong in monkey A, who chose the rightward target, when stimulated, in 65.9% of cases at the point of otherwise equal preference (50%). That effect was significantly different from 50% across the 8 sessions in this

animal ($p = 0.0015$, $t_7 = 5.0$). The effect is substantial—at a nearly 66% bias, the monkey chose the rightward target nearly twice as often as the leftward target. The effect pointed in the same direction for monkey B, and was highly significant across the two animals ($p = 0.0002$, $t_{15} = 4.9$). As in the single session examples (Fig. 2A, right column), the effect reversed polarity on the trials in which right FEF was stimulated (Fig. 2B, right). The monkeys chose the rightward target at the point of equal preference only in 42.2% of cases, and this differed significantly from the 50% equal preference ($p = 0.0009$, $t_{15} = -4.12$). These effects are graphically summarized at the bottom of Fig. 2B. Stimulation of left FEF significantly increased the proportion of rightward choices, whereas stimulation of left FEF significantly increased the proportion of leftward choices. The contralateral nature of these shifts suggests that ultrasound led to a heightened target representation by neurons within the stimulated regions (Fig. 2).

It is worth noting that the point of equal preference for monkey A is generally substantially distinct from 0 (Fig. 2A, top row, black curves). This means that the animal shows an inherent preference for one of the targets. Such an inherent bias is very common in this free choice task, and can vary substantially from session to session (Noudoost and Moore, 2011). To ensure that this variability did not influence our results, monkey B performed the same task with the exception that he was primed to choose the target that appeared first (see Materials and Methods). This step greatly reduces the mean and variance of inherent bias (Kubanek et al., 2015). Fig. 2B shows that from retrospect, this additional control was not necessary—the effects point in the same direction in both animals, and are on average of comparable size.

We next investigated the effects on the animals' decision-making in more detail. In particular, we quantified how ultrasound affected sigmoid fits, parameterized by 4 variables, to the choice data (see Materials and Methods). We found that ultrasound specifically influenced the shift of the curves along the horizontal axis. This amount of the horizontal shift is shown for each session as a histogram in Fig. 3. Left FEF stimulation shifted the decision curves on average by -7.8 ms (Fig. 3, left), and this effect was highly significant across the sessions ($p = 0.003$, $t_{15} = -3.5$; two-sided t-test). The effect reversed polarity during right FEF stimulation, shifting the decision curves by 4.4 ms on average. Also this effect was highly significant ($p = 0.0007$, $t_{15} = 4.2$). No other parameter of the fit was affected. These shifts corroborate the finding of Fig. 2 that ultrasound stimulation biases the animals' choices in the contralateral direction.

Stimulation of the FEF with moderate electric currents is known to elicit saccades into the contralateral hemifield (Bruce et al., 1985; Tehovnik et al., 2000). The magnitude of the stimulating current influences the saccadic endpoint—the larger the current, the farther away from the central fixation point a saccade lands (Bruce et al., 1985; Tehovnik et al., 2000). We therefore tested whether ultrasonic

stimulation of FEF produces a similar phenomenon. We did find a small but significant effect on the saccadic endpoints (Fig. 4). The effects point in the expected direction based on previous electrical microstimulation studies. For left FEF stimulation, the animals' saccade endpoints attained an additional $0.20\,^{\circ}$ for contralateral saccades (Fig. 4, left). This effect is small given that the eccentricity of the targets was $6\,^{\circ}$, but it was highly significant ($p = 0.001$, $t_{14} = -4.0$; two-sided t-test; one session lacked saccade trace data). The effect was not significant for ipsilateral choices ($p = 0.95$). Reversely, stimulation of right FEF brought the endpoints farther in the opposite, leftward direction (Fig. 4, right). It did so by $-0.23\,^{\circ}$ (contralateral choices; $p = 0.039$, $t_{15} = -2.3$) and $-0.30\,^{\circ}$ (ipsilateral choices; $p = 0.040$, $t_{15} = -2.3$). As in a previous study that subjected the FEF to ultrasound of similar parameters (Deffieux et al., 2013), we did not observe ultrasound directly eliciting saccades. This suggests that the effects of sonic stimulation are distinct to those of electrical stimulation. We did not observe a significant effect of ultrasound stimulation on the animals' reaction time, either for contralateral or for ipsilateral choices.

We investigated how rapidly the effect emerges and whether it is cumulative or, in contrast, whether there is an adaptation. Our blocked paradigm allows us to assess the effect dynamics. In particular, we gathered choice data separately for each trial number within a block. To increase the power of this analysis, we pooled choice data across left and right FEF stimulation trials. For this to be meaningful, we inverted the choices made on right FEF trials such that the effects have the same sign as on the left FEF trials, before pooling the data together. These data are presented in Fig. 5. The figure reveals that the biasing effect of ultrasound emerges immediately, on the first stimulated trial within a stimulation block. Interestingly, the effect diminishes in amplitude and becomes insignificant ($p = 0.082$, two-sided t-test) in trial 4 of a block. This suggests that the perceptual system of the animals adapts to the stimulation, either at the level of FEF neurons or within a broader perceptual system.

The differential effects of right and left FEF stimulation (Fig. 2, Fig. 3, Fig. 4) demonstrate that the effects are specific to the stimulated regions. To further validate this observation, we applied the same stimuli to control regions 10 mm more posterior to FEF, i.e., the motor cortex. Motor cortex is not involved in oculomotor choices and so in this case, there should be no effects on animals' choices or saccade endpoints. We collected data in 11 sessions of left motor cortex stimulation and 11 sessions of right motor cortex stimulation. Besides the change in the stimulation location, data were collected in the same way as with the FEF stimulation. In contrast to FEF stimulation, motor cortex stimulation did not elicit significant biases in choice behavior (Fig. 6). There were no significant shifts from equal preference (left motor cortex: $p = 0.43$, $t_{10} = -0.82$; right motor cortex: $p = 0.36$, $t_{10} = 0.95$) and no significant horizontal shifts (left motor cortex: $p = 0.5$, $t_{10} = -0.70$; right motor cortex: $p = 0.31$, $t_{10} = -1.1$). Neither were there significant shifts in saccadic endpoints ($p > 0.13$ for all four combinations of left and

right motor cortex stimulation and contralateral and ipsilateral choices).

Discussion

We report that transcranial ultrasound can modulate neurons to the extent that it can influence spatial choices of non-human primates. We used an established task that allowed us to assess the polarity, size, and spatial specificity of the effects.

We found that stimulation of left/right FEF biased animals' choices rightward/leftward (Fig. 2). Based on previous studies, this finding suggests that our pulsed ultrasound stimulus enhanced the representation of the contralateral target within the FEF. In comparison, neuronal inhibition—be it temporary using injected drugs or permanent following lesions—produces shifts of opposite polarity (Rorden et al., 1997; Ro et al., 2001; Schiller and Tehovnik, 2003; Wardak et al., 2004; Kubanek et al., 2015). We therefore conclude that our stimulus, at least in part, excited neurons within the FEF. This finding is supported by a study that recorded neuronal responses within supplementary eye fields in response to ultrasonic FEF stimulation in an anti-saccade task (Wattiez et al., 2017).

The effect was moderate. It was much smaller than effects attained using large injections of potent neuromodulatory drugs (Fig. 1C), but of comparable size to injections of smaller drug volumes (Kubanek et al., 2015) or electrical microstimulation (Hanks et al., 2006). It is possible that the magnitude of our effects is diminished by our block design, which frequently interleaved stimulated and non-stimulated blocks within a session. In such a block design, carry-over effects and adaptation may reduce the total effect. Nonetheless, the finding that noninvasive ultrasound can produce effects of similar magnitudes as those induced by drugs injected through craniotomies has strong implications for future research of basic brain function. Ultrasound, by virtue of its noninvasiveness and spatial flexibility, may for the first time enable us to screen the contribution of specific brain regions to a given behavior or disease sign systematically, one by one, and in a personalized fashion (Kubanek, 2018).

The ultrasonic effect was specific to the stimulated region, based on two lines of evidence. First, reversing the stimulation hemisphere reversed the effect polarity (Fig. 2, Fig. 3). This constitutes a double dissociation of the effect through different brain regions. In addition, there were no effects when stimulating control regions that are not involved in oculomotor choice, i.e., right and left motor cortex (Fig. 6). Being able to demonstrate a double dissociation regarding the stimulation site is critical to control for generic artifacts that can be associated with propagating ultrasound (Guo et al., 2018; Sato et al., 2018).

We discovered that the biasing effect manifests mainly as a horizontal shift of the decision curves (Fig. 3). This finding suggests that our stimulus enhanced the representation of the contralateral target within the FEF. A horizontal shift suggests that the stimulation biased animals' perception of the timing of the targets. The effect therefore seems, at least in part, of a perceptual nature. (A simple motor bias would manifest as an upward or downward shift of the decision curves.) Notably, the slope of the decision curves remained unchanged. This indicates that ultrasound did not fundamentally impair the animal's ability to perceive the stimuli. On the contrary, it apparently enhanced the representation of the contralateral target such that it was more likely to be chosen on a given stimulation trial.

There was a small but significant effect on the amplitude of saccades directed into the hemifield contralateral to the stimulated FEF (Fig. 4). Affecting a saccade metric such as saccade endpoint may appear as evidence of ultrasound acting, in part, on motor aspects of saccade planning within FEF. An equally likely possibility, however, is that ultrasound enhanced or shifted the perception of the contralateral target, thus resulting in a slight increase in saccade amplitude. Recording neural activity from FEF and from other, non-stimulated nodes of the oculomotor network such as the parietal area LIP, might help to distinguish between these possibilities.

Our blocked design allowed us to assess the effect progression as a function of the number of consecutive stimuli (Fig. 5). A neuromodulation effect can be constant in time, cumulative—increasing with each additional intervention, or diminishing—decreasing with each additional intervention. We found evidence for the latter kind. The effect emerged immediately, within the first stimulated trial. It then gradually decreased in size until becoming insignificant at about 4-5th consecutive stimulation trial. This indicates that the system adapted to the repetitive stimulation. The adaptation may occur at the molecular level, whereby the molecular machinery gradually loses sensitivity to repetitive excitation. At the molecular level, there is an emerging view that the effects of ultrasound on neurons are of a mechanical kind. In particular, the mechanical forces associated with propagating ultrasound displace membranes and this way open mechanosensitive ion channels (Tyler, 2011; Kubanek et al., 2018; Prieto et al., 2018). It has been demonstrated that such mechanosensing systems can indeed adapt to repetitive stimulation at the molecular level (Geffeney and Goodman, 2012). Nonetheless, the adaptation could also occur at a high perceptual level. For instance, the animal may learn to gradually ignore an ultrasound-mediated enhancement, since it brings no particular relevance to the task—and in fact—may be detrimental to task performance. Future recordings of neural activity may help distinguish between these two possibilities.

We used a relatively low stimulation frequency (Deffieux et al., 2013) to diminish the role of the skull in neuromodulatory outcomes (Lee et al., 2016). As a consequence, the pressure field associated with our stimulus was relatively broad (Fig. 1D). Although this may appear as a drawback from the perspective of

future applications, this in fact provided two benefits in regard to the basic aims of this study. First, the stimulus provided a certain level of tolerance in our FEF targeting. Second, the oblong depth geometry enabled us to stimulate an entire depth of the anterior bank of the arcuate sulcus, a region associated with the FEF. However, it is worth noting that the relatively broad stimulus in part likely influenced other neighboring regions, such as the DLPFC. Future studies can use much more circumscribed stimuli to realize the focusing strength of ultrasound (e.g, about 3 mm half-width when applied through the human skull using large, helmet-like arrays (Ghanouni et al., 2015) and less than 1 mm half-width when applied through a mouse skull (Li et al., 2016)).

The choice paradigm (Fig. 1) used in this study offers several benefits to future studies. First, it can be applied to characterize the effect polarity, size, and spatial specificity of any insonation—and, for that matter—of any neuromodulation protocol. This includes non-invasive (transcranial magnetic/electrical stimulation) and invasive (optogenetics, electrical microstimulation, pharmacological injections) neuromodulation approaches. Second, the block paradigm can be modified (e.g., increased in duration), so that also plastic, long-term effects associated with long-term stimulation (Dallapiazza et al., 2017) can be assessed. Third, the paradigm enables to quantify the effect polarity, size, and local specificity noninvasively, from a subject's choice behavior. And finally, the task is easy to learn and master for animals, which is of tremendous benefit in NHP studies.

In summary, we used a task commonly employed in neurology and NHP research to quantify the polarity, size, and spatial specificity of the effects of transcranial focused ultrasound on neurons in NHPs. We demonstrate that ultrasound can noninvasively modulate neurons in oculomotor circuits and so substantially influence animals' spatial decisions. We show that the effect points in the contralateral direction, that its size is comparable to moderate injections of neuromodulatory drugs into oculomotor regions, and that the effect is localized since stimulating the opposite hemisphere reverses the effect's polarity. A major contribution of the study is the demonstration that the effects of ultrasound on neurons in NHPs are of sufficient magnitude to modulate behavior. This is critical because flexible, systematic neuromodulation of neural circuits will enable rapid and causal screening of the candidate circuits involved in specific disorders. This way, ultrasonic neuromodulation could realize its potential in noninvasive and personalized diagnoses of a variety of brain conditions.

Materials and Methods

Subjects

Two adult male rhesus monkeys (macaca mulatta, monkey A: 13 kg, monkey B: 7 kg) participated in this study. The animals sat head-fixed in a custom designed monkey chair in a completely dark room. Visual stimuli were displayed on a LCD monitor positioned 25 cm in front of the animals' eyes. Eye position was monitored using a camera (EyeLink). All procedures conformed to the Guide for the Care and Use of Laboratory Animals and were approved Stanford University Institutional Animal Care and Use Committee.

Task

Monkeys were trained in a visual discrimination task that has been used to measure neural deficits or enhancements in previous studies (Rorden et al., 1997; Schiller and Chou, 1998; Ro et al., 2001; Wardak et al., 2002; Scherberger et al., 2003; Balan and Gottlieb, 2009). Briefly, in this task, monkeys first acquired a fixation target. After a short delay, a first target (gray square of 0.5° by 0.5°) appeared in the left (right) part of the screen, 6° away from the center of fixation. After a random delay ($[0, 130]$ ms, adjusted to the performance of each monkey), a second target, of identical parameters, appeared in the right (left) part of the screen. Once presented, both targets remained present until a choice was made. To receive a liquid reward, the animals had to make a saccade to one of the targets after the appearance of the first target within a 2° acceptance window, and remain in the window for at least 100 ms. In monkey A, choice of either target was rewarded. This free choice task is commonly associated with a substantial bias preference for one of the targets, and this bias varies considerably across days (Noudoost and Moore, 2011). To test whether our results are independent of this bias, monkey B was only rewarded for choosing the target that appeared first. This effectively mitigates a bias (Kubanek et al., 2015). The effects of ultrasonic stimulation had the same polarity and were comparable in magnitude in both monkeys.

Ultrasonic stimulation

We stimulated the macaque frontal eye fields (FEF) using the same approach and transducer as described previously (Deffieux et al., 2013). The main difference is that our transducer operated at 270 kHz instead of 320 kHz, and we used a longer stimulus, 300 ms instead of 100 ms. The single element transducer (H-115, diameter 64 mm, Sonic Concepts), geometrically focused to 63 mm, was used with a coupling cone filled with agar. The height of the cone was chosen such that the geometric focus was located 5 mm below the skull, to ensure that ultrasound stimulated neurons within the entire depth of the arcuate sulcus. Pulsed stimulus (300 ms duration, 500 Hz pulse repetition frequency, 50% duty cycle; Fig. 1D) was generated using a commercial function generator (33520B, Keysight) and subsequently

amplified using a commercial amplifier (A150, E&I). As previously (Deffieux et al., 2013), the output pressure maximum was set to 0.6 MPa, as measured in free field using a calibrated fibre-optic hydrophone (Precision Acoustics). To faithfully approximate the experimental pressure field, the coupling cone filled with agar was present during these measurements. The distribution of the pressure field was measured using a robotized moving stage (Aims III, Onda) and characterized in 1 (2) mm steps in the lateral (axial) dimensions (Fig. 1D). The animals were not sacrificed following the experiments and so the exact value of the pressure below the skull is not known. A previous study using the same stimulation set up and animals estimated the skull pressure transmission to be 58%±8% (Deffieux et al., 2013). During the experiment, the animals' hair was shaved and degassed ultrasound gel applied on the skin to mediate good acoustic coupling between the agar-filled coupling cone and the skin. The FEF target was localized using anatomical MRI images.

Characterization of decision curves

To investigate the effects of ultrasound on the animals' decisions in detail, we fitted the raw choice data (Fig. 2A) with a sigmoid function of four parameters—*slope*, *shift*, *vertscale*, and *vertshift*:

$$P(\Delta) = \mathit{vertscale}/(1 + \exp(-\mathit{slope}(\Delta - \mathit{shift}))) + \mathit{vertshift},$$

where Δ is the delay between the onset of the two targets and $P(\Delta)$ is the probability of choosing the rightward target (i.e., the individual points of each psychometric curve). The parameters were fitted to the choice data using non-linear minimization (function `fminsearch` in Matlab), minimizing the squared error between the fitted and the actual psychometric curves.

From this equation, the point of equal preference used in the analysis of Fig. 2B, Δ_{50} is determined as

$$\Delta_{50} = \frac{-1}{\mathit{slope}} \log\left(\frac{\mathit{vertscale}}{(0.5 - \mathit{vertshift})} - 1\right) + \mathit{shift}.$$

Acknowledgements

This work was supported by the NIH grant K99NS100986 (JK), Stanford Medicine Dean's Fellowship (JK), and Howard Hughes Medical Institute (JB, TM, WN). We thank Sania Fong and Shellie Hyde for technical and procedural assistance.

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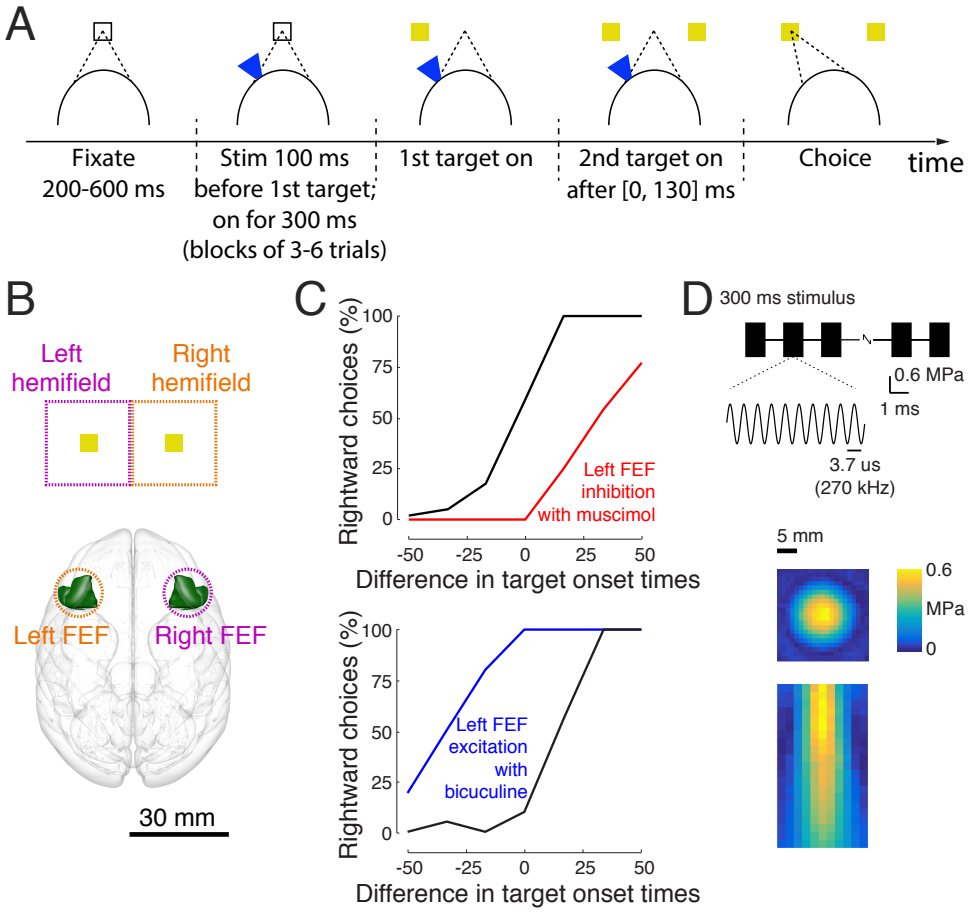

Figure 1. Task and Stimulus. A) Task. The subject fixates a central target. One target appears in the left or the right visual hemifield. After a brief random delay, a second target appears in the opposite hemifield. The subject is free to look at either target after the first target has been presented, and receives a liquid reward if he looks at a target within a 2 ° acceptance window. Ultrasound is applied in blocks of 3-6 trials, strictly interleaved with no stimulation blocks of the same duration, 100 ms prior to the appearance of the first target. B) Functional characterization of the visuomotor system. We delivered the ultrasound non-invasively (intact skull and skin) into the frontal eye fields (FEF). From anatomical and functional studies, it is known that left/right FEF preferentially represents targets in the right/left visual hemifield. The outline of the FEF was rendered using the Calabrese et al. (2015) atlas with Paxinos brain regions. C) Effects in previous studies. When a large amount of strong inhibitory/disinhibitory drugs is injected into left FEF in this task, animals show a strong ipsilateral/contralateral bias in this task (reproduced from Schiller and Tehovnik, 2003, with permission). These results are as expected given the contralateral nature of the visual hemifield representation (B), and are analogous when other nodes of the visuomotor network, such as the parietal area LIP, are perturbed. D) Stimulus. The ultrasound stimulus (0.6 MPa, 270 kHz, 300 ms duration) was pulsed at 500 Hz with 1 ms tone burst duration. The ultrasound was applied through a coupling cone filled with agar gel. The resulting pressure, measured in free field, is provided along the lateral (1 mm steps) and axial (2 mm steps) dimensions in color.

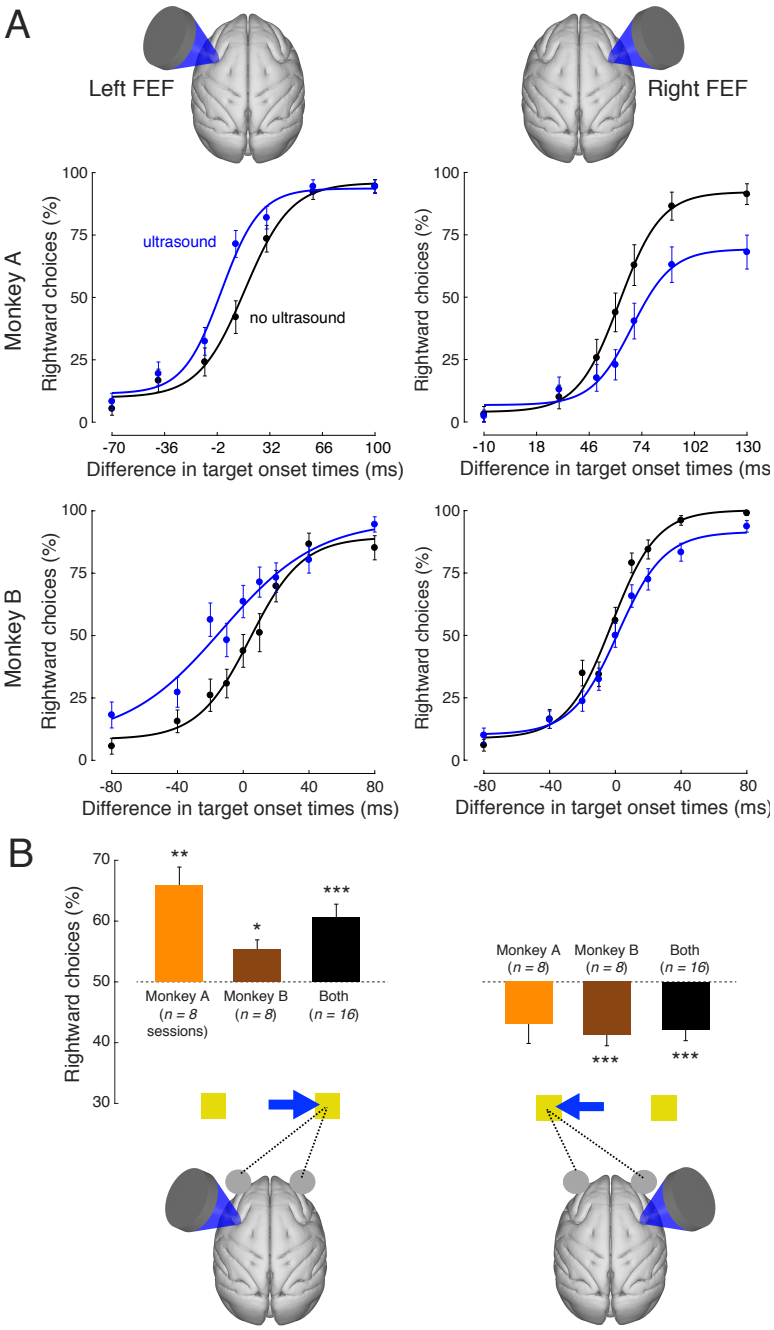

Figure 2. Ultrasonic stimulation of frontal eye fields biases visuomotor decisions. A) Single session examples. Mean (± s.e.m.) proportion of choices of the rightward target as a function of the difference in target onset times. Positive difference stands for the cases in which the rightward target appeared first. The black data points reflect choice behavior in the trials in which the animal was not stimulated, whereas the blue data points represent choice behavior in the stimulated trials. The data were fit with a four-parameter sigmoid function (see Materials and Methods). The data are presented separately for left and right FEF stimulation sessions (left and right columns; see illustration on top), and for monkey A and B (top and bottom row). B) Quantification of the effects for all sessions. The no stimulation data (black) of each session were first fit with a sigmoid function. Using the fit, we identified the time difference on the abscissa for which the animal chooses both targets in equal proportion (see Materials and Methods). At that point, we then assessed the proportion of rightward choices during the stimulated trials. The stars indicate the cases in which the mean effects statistically differ from equal preference (two-sided t-test; $*$: $p < 0.05$, $**$: $p < 0.01$, $***$: $p < 0.001$). The illustration on the bottom summarizes the polarity of these biasing effects.

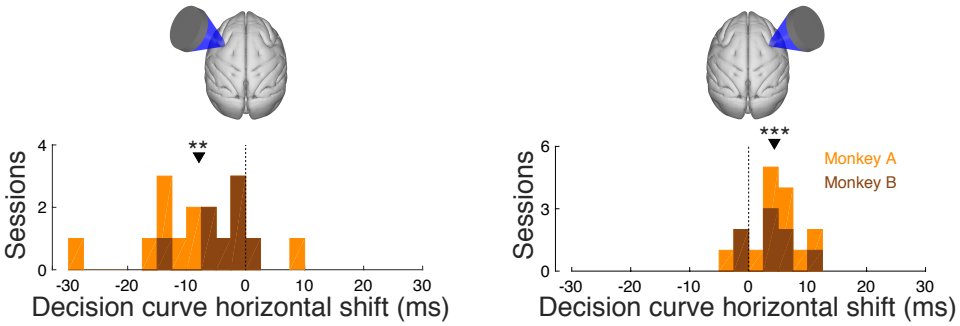

Figure 3. The effect constitutes a horizontal shift of the decision curves. The amount of horizontal shift of the decision curves by ultrasound (blue versus black in Fig. 2A). The data are presented separately for left and right FEF stimulation sessions (left and right panel), and separately for each session in each animal (effect histogram). The stars indicate the effect significance (two-sided t-test; $*$: $p < 0.05$, $**$: $p < 0.01$, $***$: $p < 0.001$).

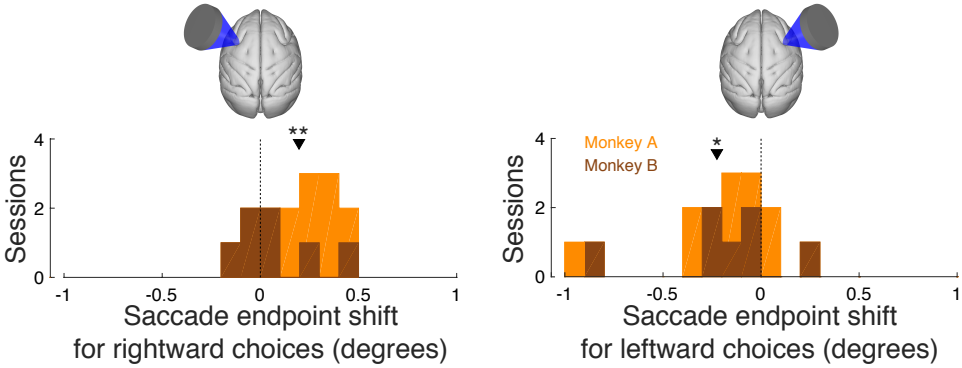

Figure 4. Ultrasound translates saccade endpoints further in the contralateral direction. The histograms show the change in saccade endpoints (visual degrees) following ultrasonic stimulation as a trial average over each session. The left/right panels specifically show the effects for contralateral (right/left) choices. The stars indicate the effect significance (two-sided t-test; $*$: $p < 0.05$, $**$: $p < 0.01$, $***$: $p < 0.001$).

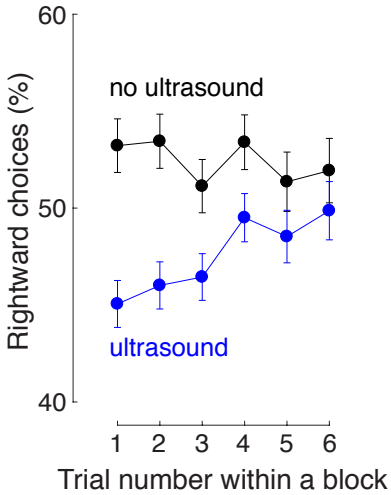

Figure 5. Effect dynamics. Mean $\pm$ s.e.m. choices of the rightward target as a function of trial number within a block, separately for no stimulation (black) and stimulation (blue) block of trials. Choices during right FEF stimulation were inverted so that they could be pooled together with left FEF choices to increase the signal-to-noise ratio of the analysis.

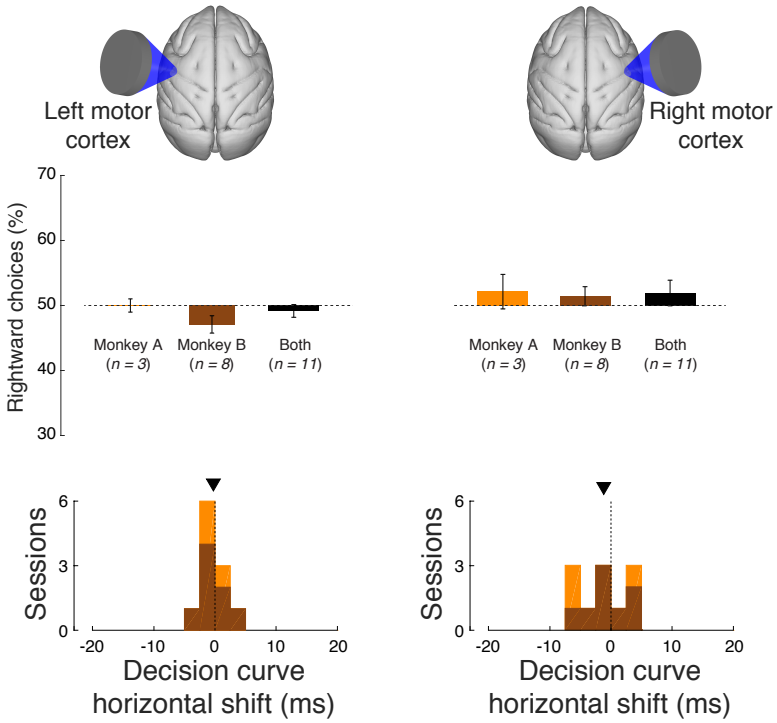

Figure 6. Stimulation of motor cortex had no effect on choice behavior. Same format as in Fig. 2 and Fig. 3, for ultrasound stimulation of left motor cortex (left panel) and right motor cortex (right panel).