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Supplemental Information

Driving Oscillatory Activity

in the Human Cortex

Enhances Motor Performance

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Supplemental Inventory

1. Supplemental Figures and Tables

Figure S1, related to Figure 3C

Figure S2, related to Figures 2A and 3A $\,$

Table S1, related to Results

Table S2, related to Results

Table S3, related to Results and Figure 3C

Table S4, related to Results

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2. Supplemental Experimental Procedures

3. Supplemental Analysis

4. Supplemental Results

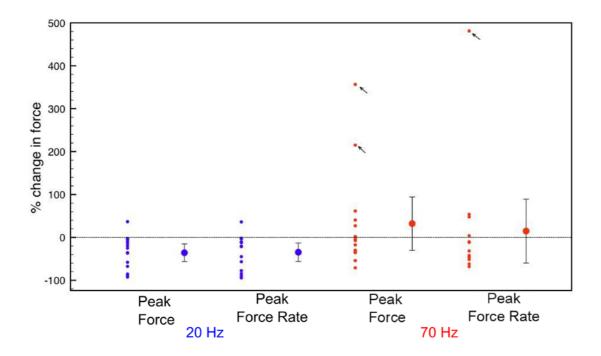


Figure S1. No-Go % Change with Outliers Included, Related to Figure 3C Individual % changes are shown for 20 Hz (blue) and 70 Hz (red) for peak force and peak rate along with means \pm 2 SEM. Outliers are identified with arrows.

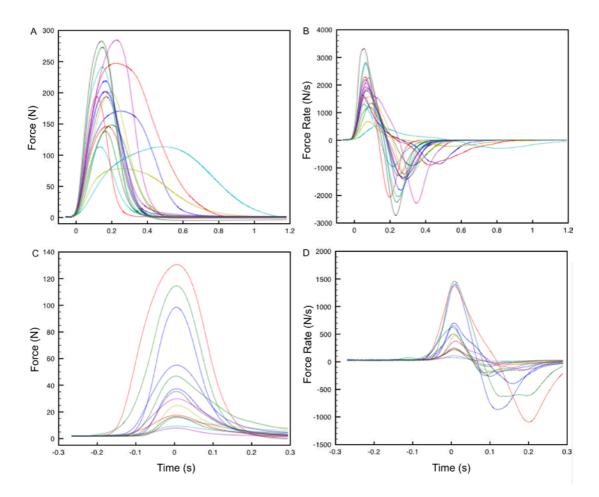


Figure S2. Individual Subject Averages during No-Stimulation, Related to Figure 2A and Figure 3A

Average traces per subject during no-stimulation trials in the 20 Hz session, shown for (A) go force aligned to onset, (B) go force rate aligned to onset, (C) no-go force aligned to peak, and (D) no-go force rate aligned to peak. Subjects are color matched across panels.

Condition	No Stimulation	Stimulation	Significance			
	(mean <u>+</u> SEM)		(paired t-test)			
Go reaction time	Go reaction time					
20 Hz	271 <u>+</u> 7.1 ms	268 <u>+</u> 8.3 ms	P=0.373			
70 Hz	266 <u>+</u> 11 ms	262 <u>+</u> 8.4 ms	P=0.416			
Go Peak force						
20 Hz	200 <u>+</u> 15 N	199 <u>+</u> 15 N	P = 0.198			
70 Hz	206 <u>+</u> 17 N	207 <u>+</u> 17 N	P = 0.546			
Percentage go trial	s rejected					
20 Hz	6.5 <u>+</u> 1.5	7.4 <u>+</u> 1.7	P=0.511			
70 Hz	5.8 <u>+</u> 0.9	6.1 <u>+</u> 1.0	P=0.705			
Force between fixation cross and pre-cue						
20 Hz	9.3 <u>+</u> 6.4 N	9.4 <u>+</u> 6.4 N	P=0.319			
70 Hz	10.0 <u>+</u> 6.2 N	10.1 <u>+</u> 6.2 N	P=0.655			
Force between pre-cue and response						
20 Hz	9.3 <u>+</u> 6.4 N	9.4 <u>+</u> 6.4 N	P=0.311			
70 Hz	10.0 <u>+</u> 6.2 N	10.1 <u>+</u> 6.2 N	P=0.580			

Table S1. Nonsignificant Differences for Go Trials, Related to Results

Table S2. Percentage of Errors of Commission and Total Number of No-Go Trials per Stimulation Frequency, Related to Results

Condition	No Stimulation	Stimulation	Significance (paired t-test)
Percentage relative to total number of trials			
20 Hz	44.5% <u>+</u> 6.1	41.6% <u>+</u> 7.0	P=0.336
70 Hz	45.3% <u>+</u> 5.3	45.0% <u>+</u> 5.5	P=0.947
Absolute number of trials			
20 Hz	13.3 <u>+</u> 1.7	12.8 <u>+</u> 1.7	P=0.758
70 Hz	13.0 <u>+</u> 1.4	12.6 <u>+</u> 1.5	P=0.664

Table S3. Related to Results and Figure 3C

70 Hz no-go trials, outliers removed. Gamma stimulation remains insignificant.

Parameter	% change	Significance
Peak Force	-10.1 + 11.2	P=0.386
Peak Rate	-21.0 + 11.1	P=0.072

Table S4. Related to Results

Significant differences between 20 Hz and 70 Hz stimulation. Go and no-go trials were different for all parameters.

	Significance (paired t-test)	
GO		
Initial velocity	P<0.001	
Peak velocity	P<0.001	
Time to peak	P=0.0037	
NO-GO		
Peak force	P = 0.0193	
Peak velocity	P = 0.0212	

Table S5. Nonsignificant Correlations, Related to Results

There was no correlation between percent changes during 20 Hz and 70 Hz stimulation, or between intensity of stimulation and performance.

Parameter 1	Parameter 2	r value	Significance		
% change for 20 Hz and 70 Hz stimulation					
Go trial initial rate	Go trial initial rate	-0.0356	P = 0.889		
with 20 Hz	with 70 Hz				
Go trial peak rate	Go trial peak rate	0.2359	P=0.346		
with 20 Hz	with 70 Hz				
Intensity of stimula	tion and % change ii	n performance			
Intensity of	Go initial rate with	0.0851	P=0.763		
Stimulation	20 Hz				
Intensity of	Go peak rate with	0.0929	P=0.742		
Stimulation	20 Hz				
Intensity of	Go initial rate with	-0.2125	P=0.447		
Stimulation	70 Hz				
Intensity of	Go peak rate	-0.1075	P=0.703		
Stimulation	with 70 Hz				
Intensity of	Go peak force with	-0.2917	P=0.291		
Stimulation	20 Hz				
Intensity of	Go peak force with	0.2987	P=0.280		
Stimulation	70 Hz				
Intensity of	Nogo peak force	0.1073	P=0.727		
Stimulation	with 20 Hz				
Intensity of	Nogo peak rate	0.0846	P=0.783		
Stimulation	with 20 Hz				
Intensity of	No-go peak force	-0.2589	P=0.442		
Stimulation	with 70 Hz				
Intensity of	No-go peak rate	-0.2405	P=0.476		
Stimulation	with 70 Hz				

Supplemental Experimental Procedures

During the first session, the subject's phosphene or scalp sensation threshold (whichever was lowest) to 20 Hz stimulation was determined, and an amplitude 50 μ A below this was selected for subsequent testing stimulation. Impedance during all stimulation sessions was always kept below 15 m Ω . Sub-threshold stimulation was confirmed by a forced-choice task comprising of 20 rounds of stimulation. One subject was always aware when stimulation was being applied and was therefore excluded from the study. A further two subjects experienced scalp sensations or phosphenes when stimulated at the selected intensity in the second session and so the intensity in the second session was lowered by a further 100 μ A. There was no significant difference between the mean stimulus amplitude of the two sessions (903 ± 111 μ A for beta and 915 ± 98 μ A for gamma stimulation, t_[17]=-0.128, p=0.901, paired t-test).

To assess the subject's perception of the stimulation, we employed a force-choice test prior to the experiment in which the subjects would attend to the task cues but instead of making a motor response they would indicate whether or not they thought the stimulation was active each time a response cue was presented (either red or green). The stimulation setting (ON or OFF) and their response (ON or OFF) was recorded for 20 trials. We then coded correct responses as 1 and incorrect responses as 0 and performed a binomial test on each dataset. The p value was always insignificant and ranged from 0.2 to 1, with the exception of one case which was 0.077. This confirmed that subjects were not aware of any active stimulation during the task.

Supplemental Analysis

For no-go trials, we sought to analyze all trials in which some erroneous force response (ie 'twitch') had been produced. To do this we calculated the standard deviation of the baseline period, and rejected any trial in which the peak force was below 5 times this standard deviation. This allowed us to include all trials in which a small deviation from baseline occurred, but reject trials in which there was no response whatsoever. In most go/no-go experiments , the response is defined by an 'all-or-none' behavior, such as a button push, thus giving rise to a fairly low error rate, as compared with ours (mean 45%). Here, we were most interested in capturing the changes in force performance due to stimulation, and thus designated a low cutoff. We subsequently aligned trials to peak force and peak force rate, as in go trials.'

Supplemental Results

The percentage change with gamma stimulation was significantly greater with respect to the initial rate of force development than peak velocity ($t_{[17]}$ =3.39, p=0.0035), but not so for beta stimulation($t_{[17]}$ =1.08, p=0.297).

Due to the faster development of force with gamma stimulation there was a 3.65 ± 0.86 % reduction in the time to achieve peak rate compared to no-stimulation ($t_{[17]} = -4.25$, p = 0.00053; drop from 81.3 ± 4.9 ms to 78.1 ± 4.3 ms , $t_{[17]} = 3.69$, p = 0.0018). No such change was apparent for beta stimulation (0.84 ± 0.63 %, $t_{[17]} = 1.327$, p =0.202). There were significant differences in initial and peak rate when comparing beta and gamma stimulation, confirming a differential effect between the two interventions (Table S4). The independence of the effects of the two stimulation frequencies was suggested by the lack of any across-subject correlation in either initial or peak rate percentage change between beta and gamma stimulation (Table S5). In addition, there was no correlation between intensity of stimulation and any behavioral change (Table S5). Finally, there were no differences in grip force levels before the response cue was presented, regardless of the ongoing stimulation at 20 Hz or 70 Hz. The lack of modulation in baseline resting force suggests that stimulation per se did not simply alter the resting level of muscle activity (Table S1).