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# Cerebellar control of visually-guided eye movements by the bilateral mass of activity in the caudal fastigial nuclei

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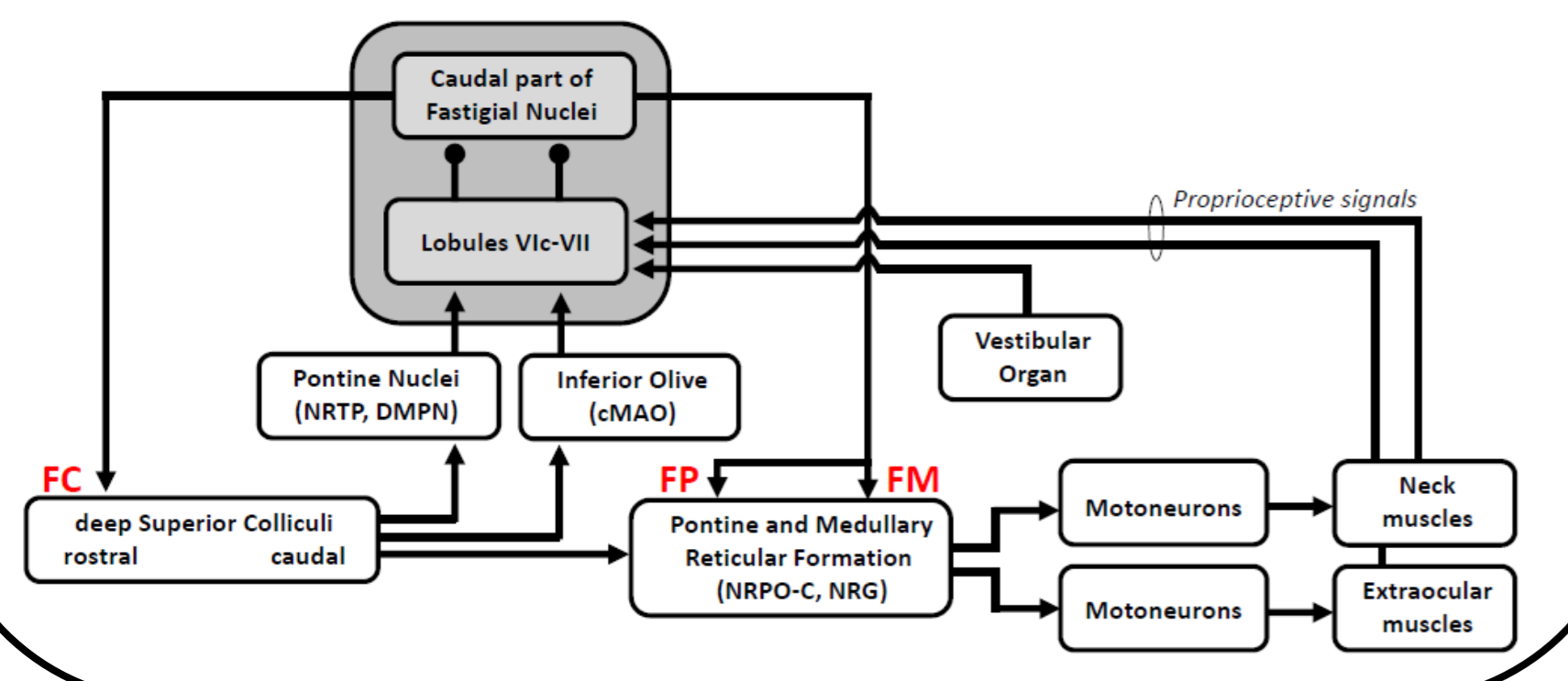
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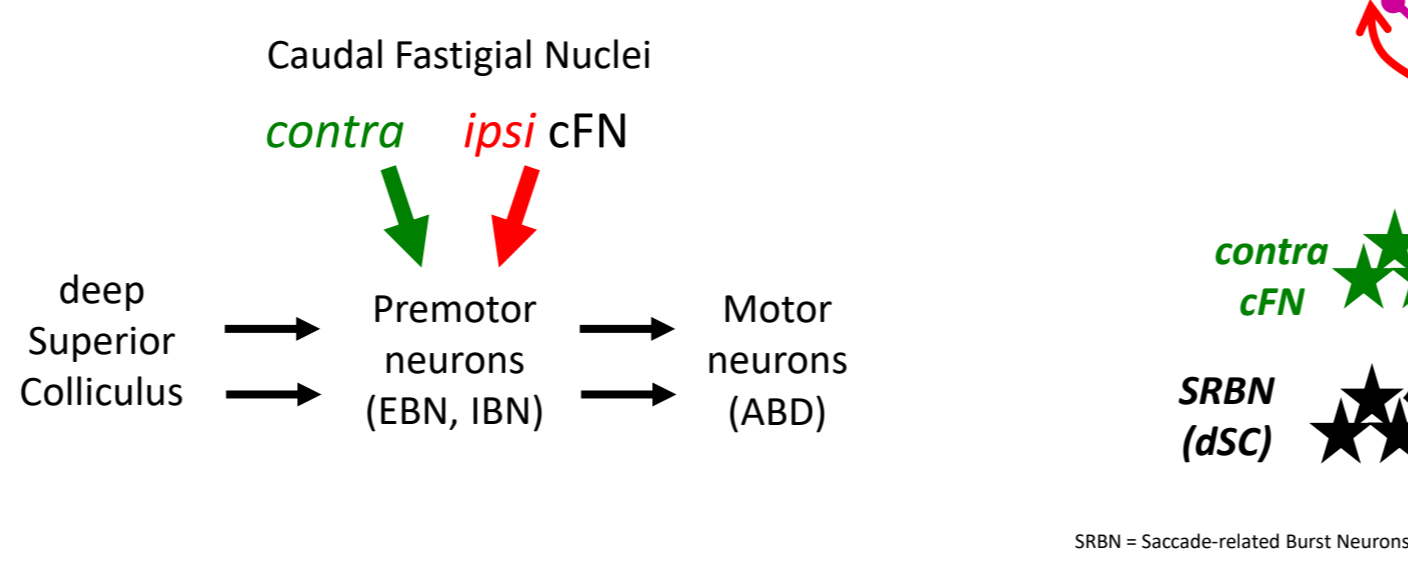


The caudal part of both fastigial nuclei (cFN) plays a crucial role in the ability to foveate a static or moving visual target. In the monkey, an unilateral inactivation of cFN by local injection of muscimol impairs the horizontal component of saccadic eye movements. An ipsilesional gaze offset is also observed while a target is being fixated or pursued. In the head-unrestrained monkey (and cat), the head exhibits an ipsilesional deviation relative to the trunk (cervical dystonia).

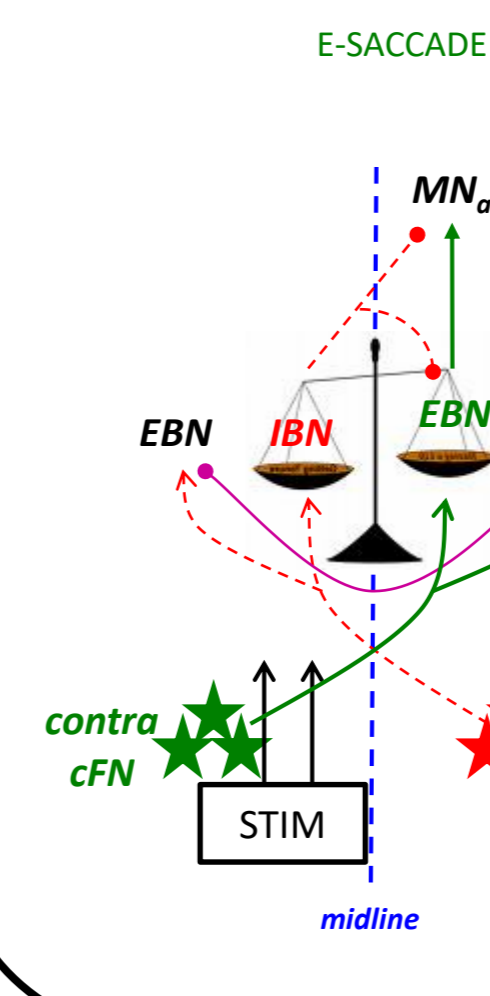


### BILATERAL HYPOTHESIS

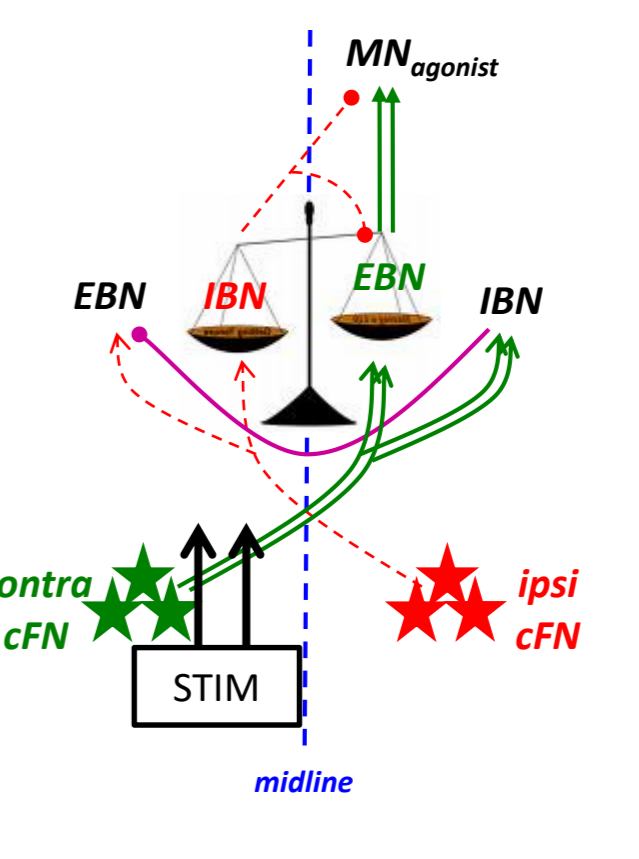
For every saccade (horizontal, oblique or vertical), the left and right cFN regulate the balance of activity between the excitatory and inhibitory input (from EBNs and IBNs) to the motor and internuclear neurons in the abducens nucleus.



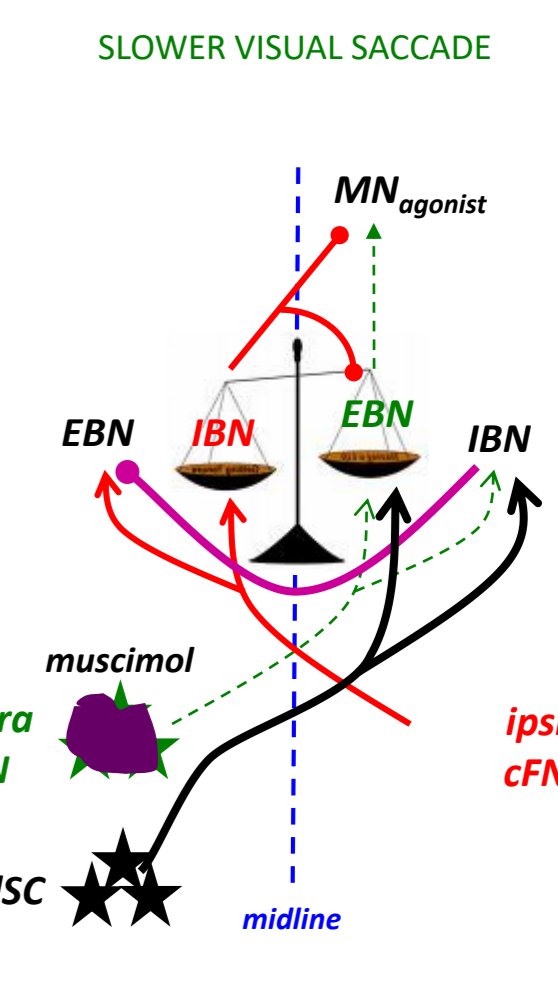
### ELECTRICAL MICROSTIMULATION



### FASTER E-SACCAD



### INACTIVATION



### ELEMENTS FOR A CONTROVERSY

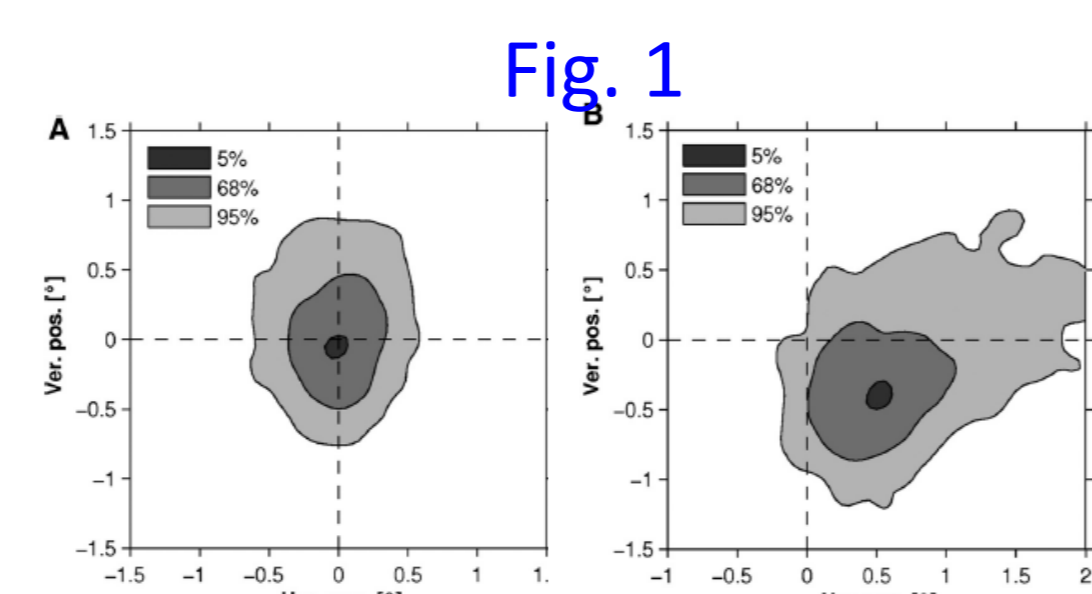
In a computational study, Eggert, Robinson & Straube (2016) reported more variability in saccade endpoints after cFN inactivation than before, and interpreted this effect as altered putative "noises" in brain activity: planning noise and/or signal dependent motor noise. Unfortunately, they did not consider the fact that after cFN inactivation, the monkeys have difficulties to foveate a visual target. The variability in saccade endpoints can indeed result from the fact that saccades did not start from the same starting positions (Table 1 and Fig. 1). Moreover, when we look at the literature, we do not find any particular increase in the variability of amplitudes. However, we find differences between experiments: the amount of dysmetria varies between the experiments (Fig. 2).

Table 1

Experiment	Injection	Session	n	Horizontal Eye Position			Vertical Eye Position		
				Mean	Range	Difference	Mean	Range	Difference
Q1	Left cFN (0.4 μM)	Control	378	-0.5 ± 0.6	[-1.0; 1.3]	-1.2*	0.0 ± 0.6	[-1.5; 1.8]	0.4*
Q2	Left cFN (0.4 μM)	Muscimol	614	-1.7 ± 1.0	[-4.9; 2.4]	-2.4*	0.4 ± 0.6	[-1.6; 3.0]	-0.1 NS
Q3	Left cFN (0.8 μM)	Control	605	0.4 ± 0.6	[-1.1; 2.4]	-2.4*	0.2 ± 0.7	[-2.2; 3.3]	-0.3*
Q4	Left cFN (0.8 μM)	Muscimol	502	-1.8 ± 1.6	[-6.6; 4.0]	-1.2*	0.1 ± 1.6	[-0.6; 6.1]	-0.3*
Q5	Right cFN (0.6 μM)	Control	782	0.0 ± 0.7	[-3.2; 3.1]	-1.2*	0.1 ± 0.7	[-2.5; 2.4]	-0.2*
Q6	Right cFN (0.6 μM)	Muscimol	1071	-1.2 ± 1.2	[-7.2; 2.7]	2.0*	-0.2 ± 0.6	[-3.0; 2.4]	-0.2*
Q7	Left cFN (1.0 μM)	Control	690	-0.2 ± 1.0	[-3.5; 2.9]	-1.4*	-0.3 ± 0.9	[-3.0; 3.1]	0.1 NS
Q8	Left cFN (1.0 μM)	Muscimol	962	1.8 ± 1.3	[-3.5; 6.4]	-1.4*	0.1 ± 0.8	[-1.1; 3.3]	-0.1 NS
Q9	Right cFN (0.6 μM)	Control	319	0.1 ± 0.7	[-1.1; 1.6]	-1.3*	-0.1 ± 0.4	[-1.2; 1.2]	-0.4*
Q10	Right cFN (0.6 μM)	Muscimol	378	-1.2 ± 0.8	[-3.4; 3.6]	-0.4*	-0.9 ± 0.4	[-2.2; 2.0]	-0.2*
Q11	Left cFN (0.8 μM)	Control	488	0.0 ± 0.8	[-2.3; 2.0]	0.8*	0.0 ± 0.5	[-1.2; 1.2]	-1.0*
Q12	Left cFN (0.8 μM)	Muscimol	425	0.1 ± 0.6	[-2.4; 1.7]	0.8*	0.0 ± 0.4	[-2.2; 0.7]	-1.0*
Q13	Right cFN (0.8 μM)	Control	496	0.5 ± 0.7	[-1.5; 2.3]	-0.3*	-0.1 ± 0.5	[-1.6; 2.1]	-0.1*
Q14	Right cFN (0.8 μM)	Muscimol	609	-0.3 ± 0.8	[-3.2; 2.5]	0.3*	-0.1 ± 0.5	[-1.6; 1.6]	0.2*
Q15	Right cFN (1.2 μM)	Control	88	0.3 ± 0.8	[-2.0; 2.5]	0.3*	-0.1 ± 0.4	[-1.2; 1.2]	0.2*
Q16	Right cFN (1.2 μM)	Muscimol	154	0.6 ± 1.0	[-1.6; 3.8]	0.1 ± 0.8	0.1 ± 0.8	[-1.4; 2.5]	0.2*

Difference was tested with a Student's t-test for independent samples; NS: no statistically significant difference; \*P < 0.05. LED, light-emitting diode; cFN, caudal fastigial nucleus.

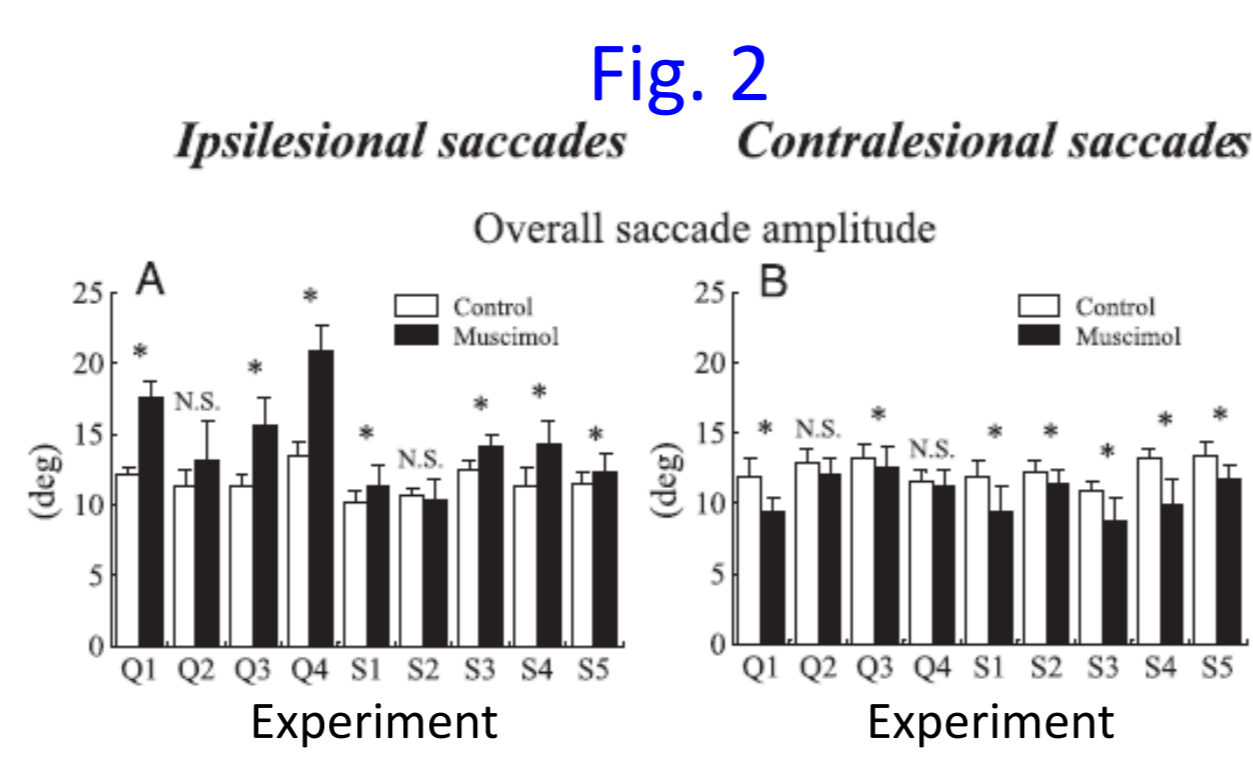
Fig. 1



Spatial distribution of dwell times before (A) and after (B) inactivation of the right FOR. Each area encloses the 5, 68, and 95% (from darkest to lightest) of the total fixation time during an experiment. After muscimol injection in the right FOR, the primary saccade (either hypometric or hypermetric) brings the eyes to the ipsilesional side. A contralateral saccade is prompted for improving target foveation, but its amplitude is smaller than required to accurately place the target image on the same region of the fovea as before the injection. Ipsilesional saccades can also be observed even before gaze reaches the target. Consequently, gaze dwells longer in the ipsilesional side.

from Guerrasio, Quinet, Büttner & Goffart J Neurophysiol (2010)

Fig. 2



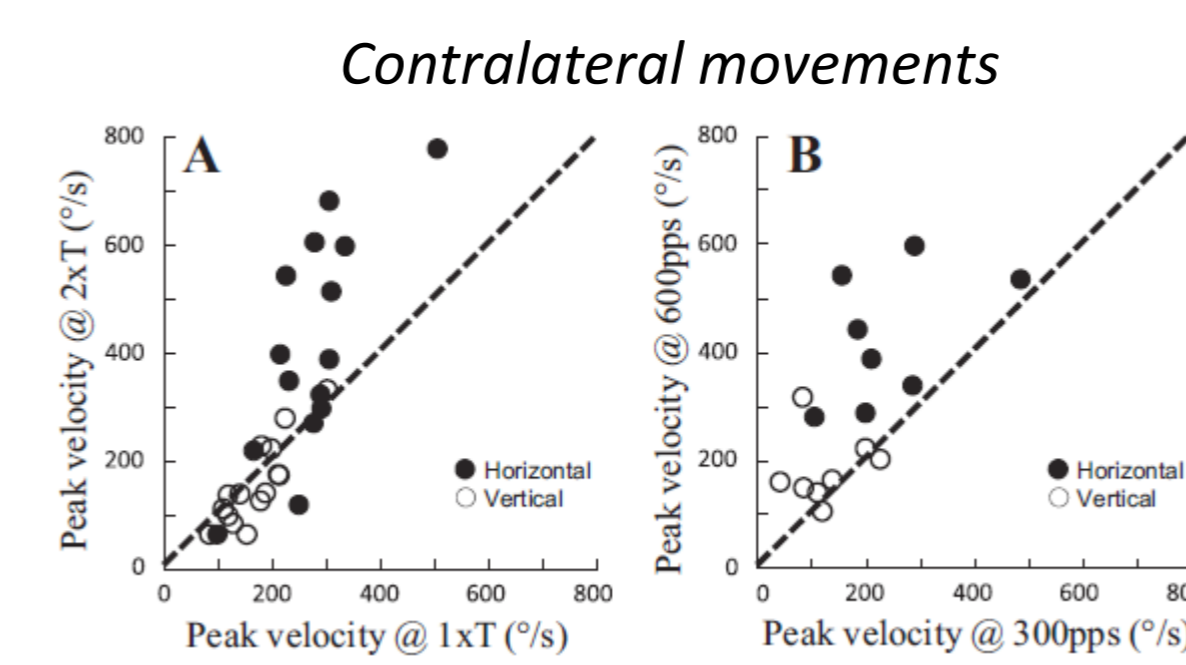
Mean amplitude of horizontal (ipsilesional) (A) and contralateral (B) saccades recorded before (●) and after (■) muscimol injection in the cFN. Saccades were initiated from the central fixation LED and aimed at a 12° eccentric target LED. The error bars represent the SDs. N.S.: no statistically significant difference; \*P < 0.05 (Mann-Whitney U test).

from Goffart, Chen & Sparks J Neurophysiol (2004)

### NEURONAL RECRUITMENT HYPOTHESIS

From the demonstration that larger current enhances the size and velocity of saccades evoked by electrical cFN microstimulation (Fig. 3), we propose that the variability of saccade endpoints reflects the variable size of the active population of cFN neurons contributing to the movement generation. Indeed, if during the pharmacological experiments, the perturbation is not exactly centered in cFN, the number of inactivated neurons should increase as the muscimol diffuses, resulting in a dysmetria that increases with time.

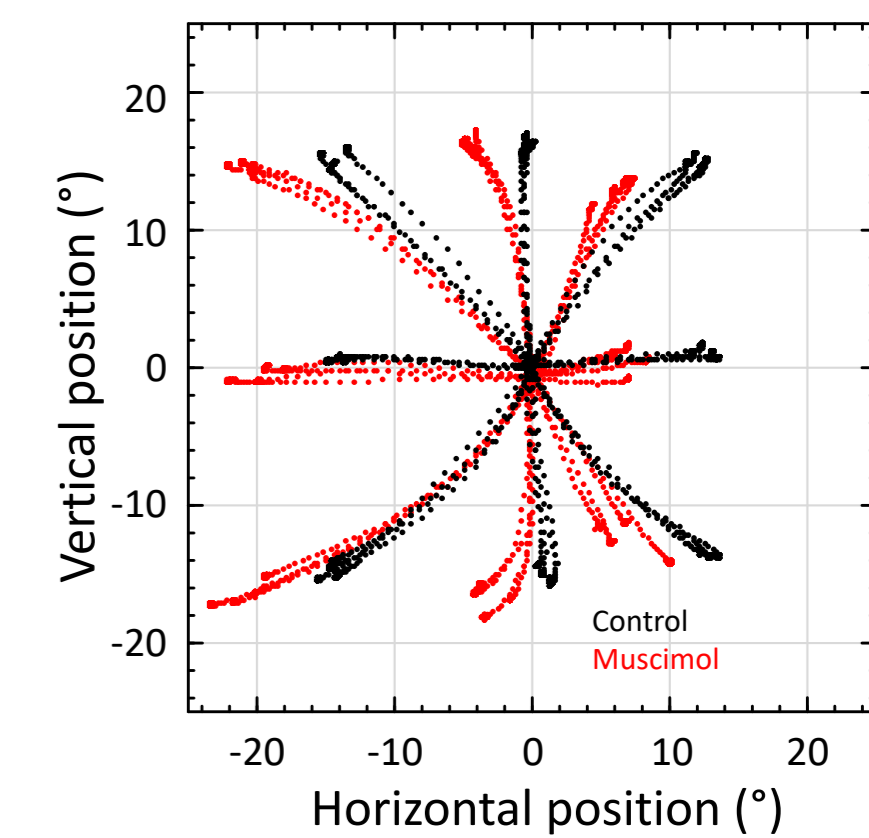
Fig. 3



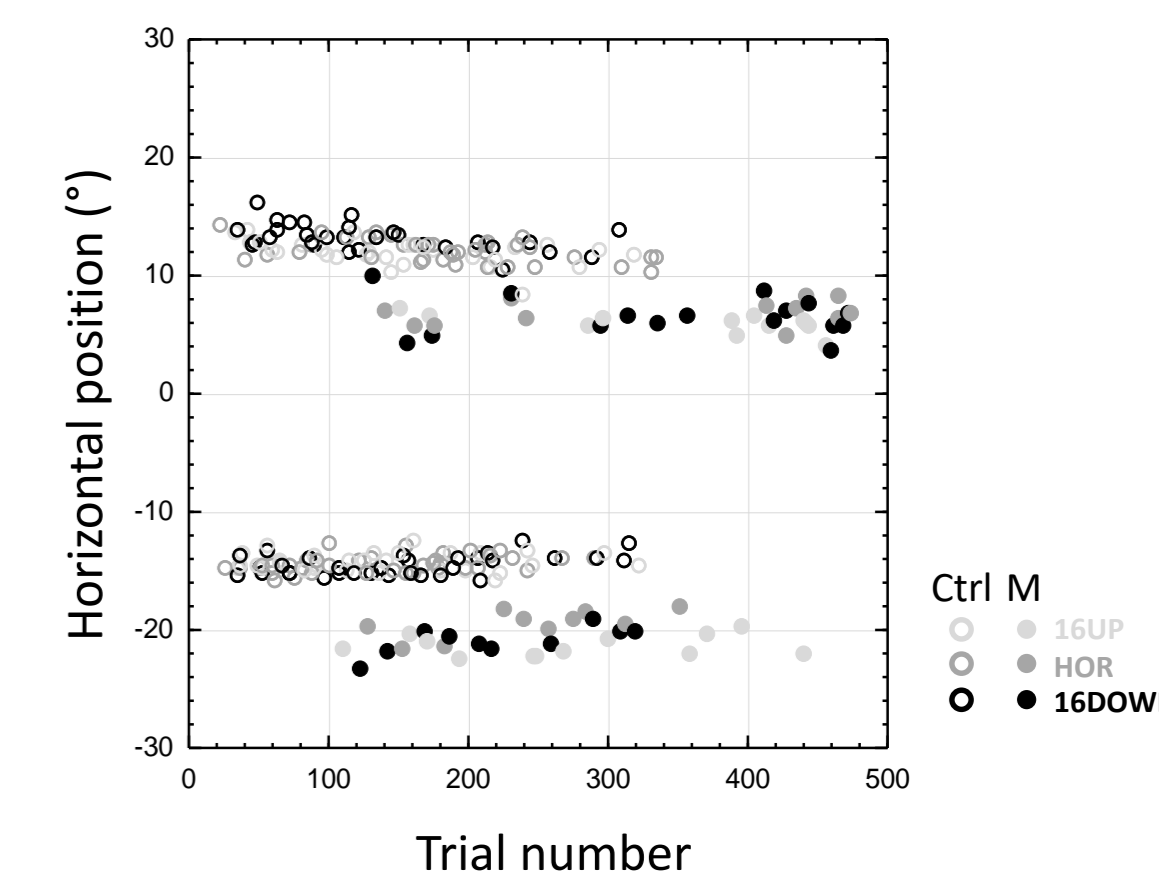
Effect of increased fastigial activity on peak velocity of contralateral saccades. The effects of doubling the stimulation current (A) and frequency (B) on the average horizontal (filled circles) and vertical peak velocities (open circles) are shown. In A, the stimulation current was increased from 17 to 27 (15 tested sites), whereas in B, the stimulation frequency was increased from 300 to 600 pps (8 sites).

from Quinet & Goffart J Neurophysiol (2015)

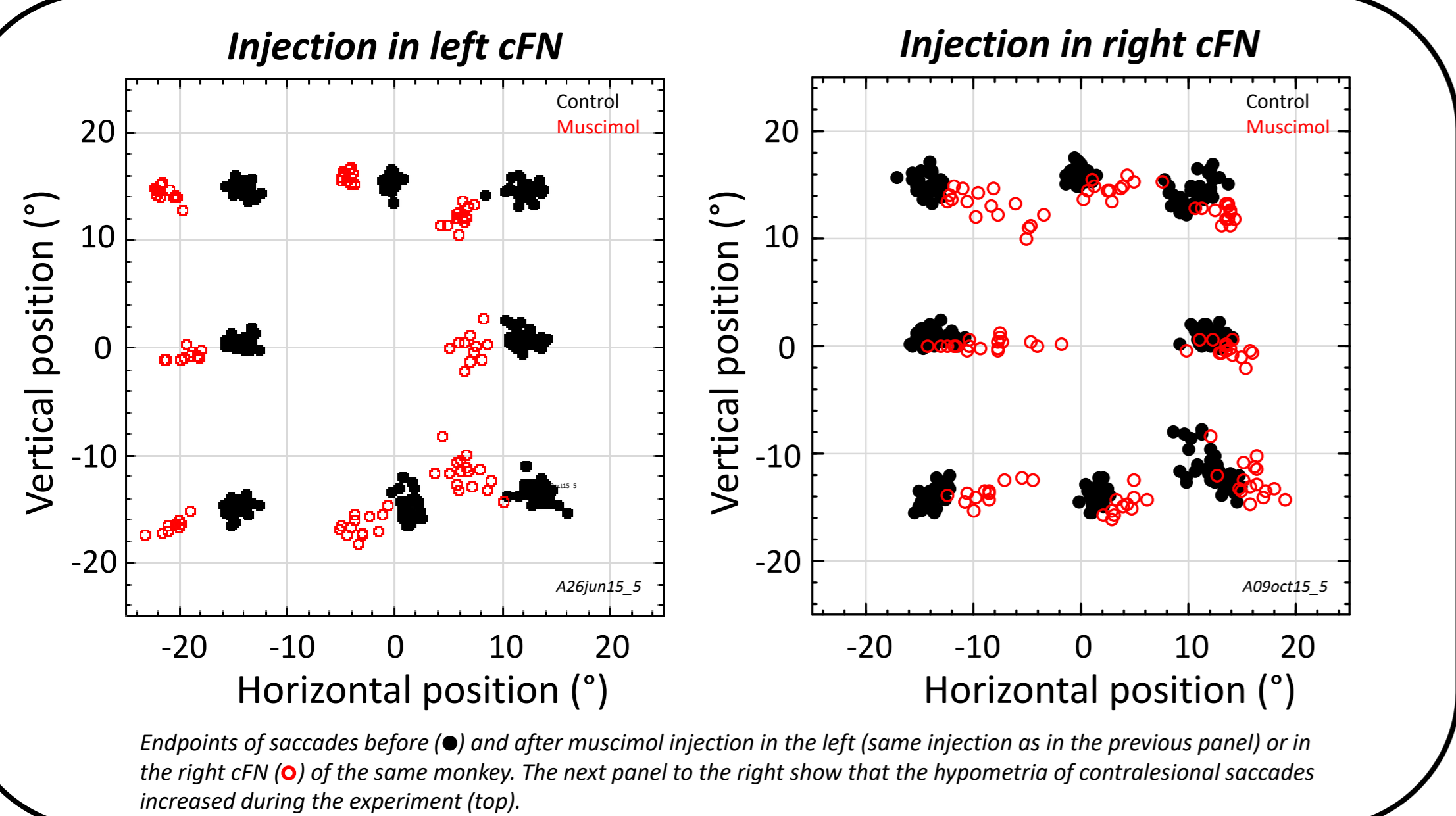
### TYPICAL EXPERIMENT



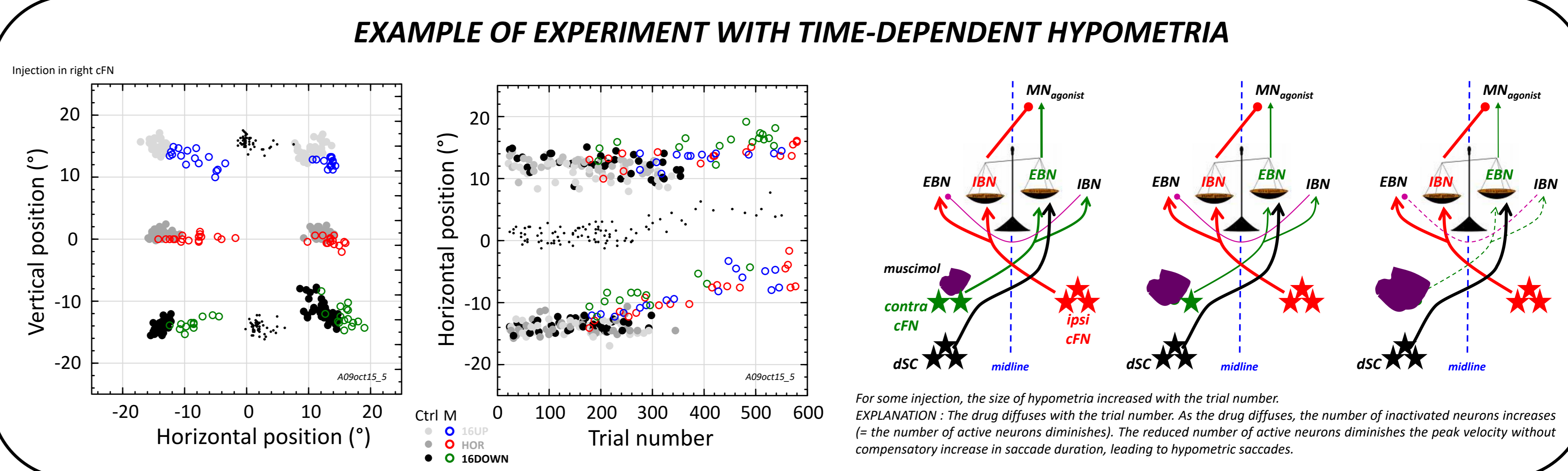
After muscimol injection in the left caudal fastigial nucleus, the horizontal component is hypermetric for ipsilateral (leftward) saccades and hypometric for contralateral (rightward) saccades. Vertical saccades also exhibit a leftward deflection in their trajectory.



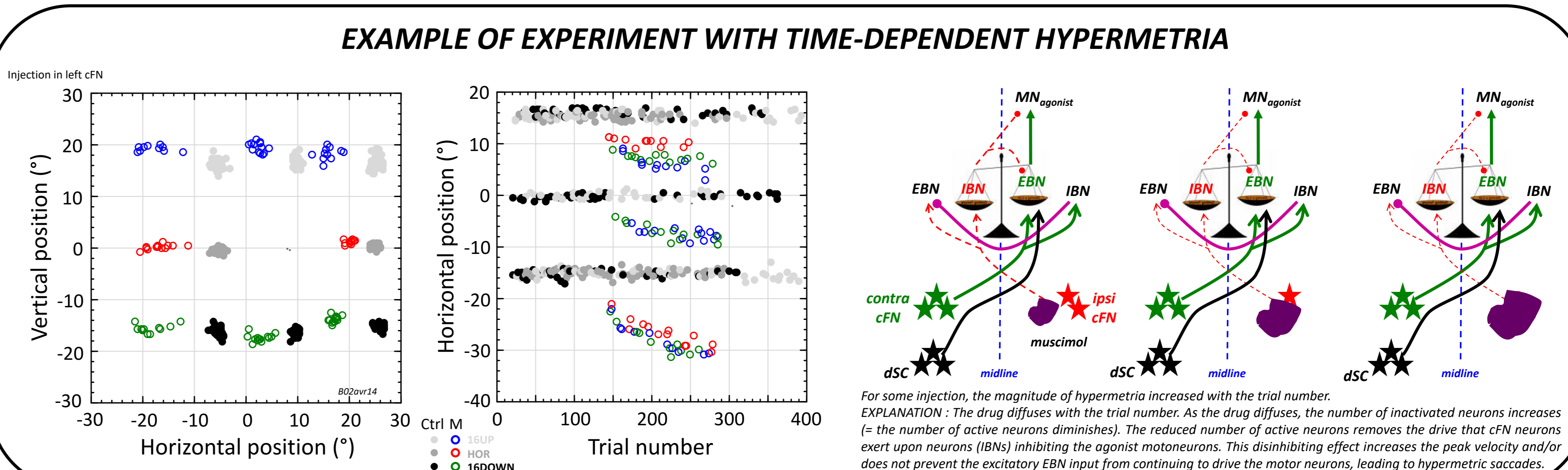
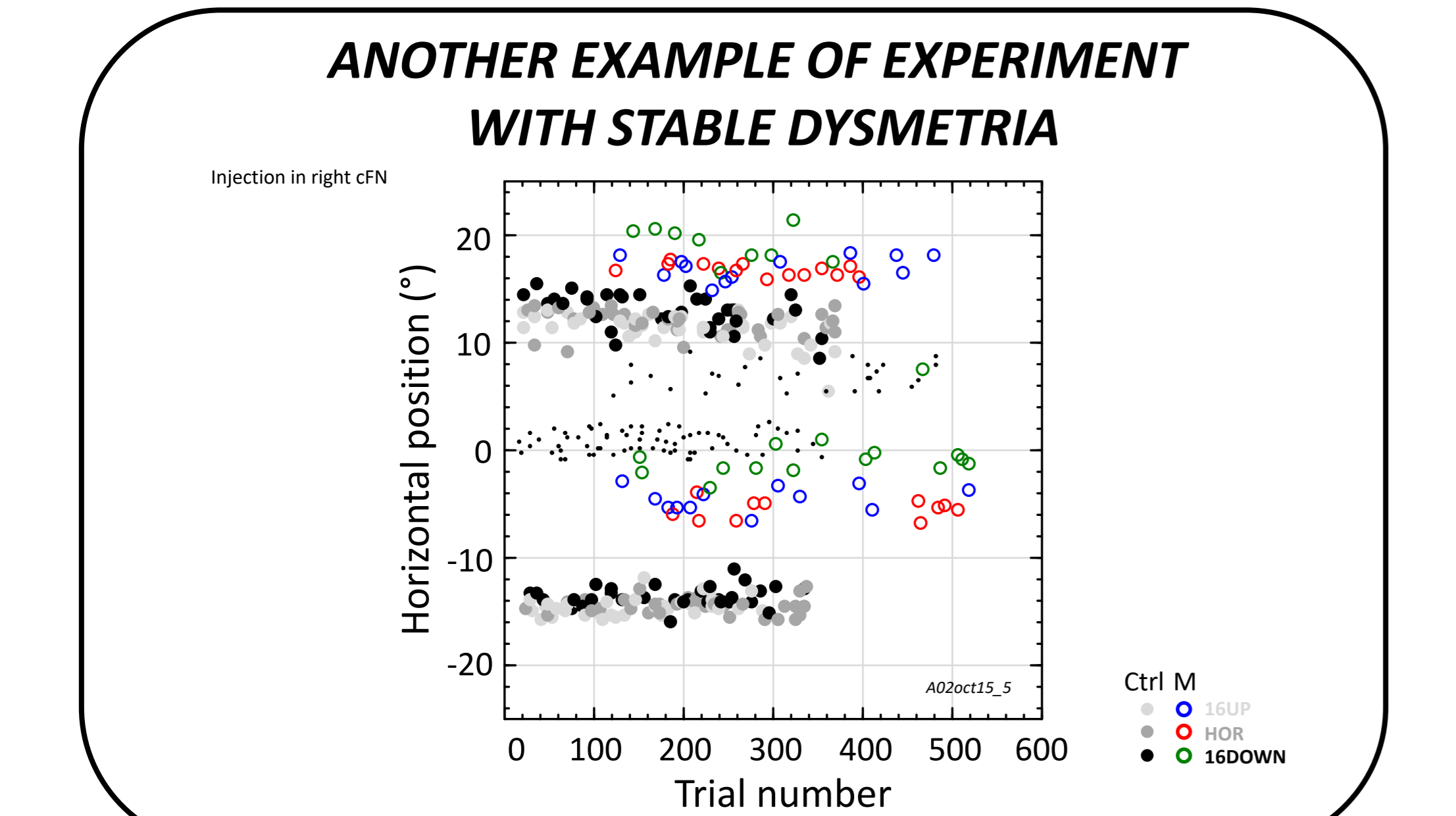
The endpoints of contralateral saccades (positive values) and ipsilesional saccades (negative values) do not change across trials.



Endpoints of saccades before (●) and after muscimol injection in the left (same injection as in the previous panel) or in the right cFN (○) of the same monkey. The next panel to the right show that the hypometria of contralateral saccades increased during the experiment (top).



For some injection, the size of hypometria increased with the trial number. EXPLANATION: The drug diffuses with the trial number. As the drug diffuses, the number of inactivated neurons increases (= the number of active neurons diminishes). The reduced number of active neurons diminishes the peak velocity without compensatory increase in saccade duration, leading to hypometric saccades.



For some injection, the magnitude of hypermetria increased with the trial number. EXPLANATION: The drug diffuses with the trial number. As the drug diffuses, the number of inactivated neurons increases (= the number of active neurons diminishes). The reduced number of active neurons removes the drive that cFN neurons exert upon neurons (IBNs) inhibiting the agonist motoneurons. This disinhibiting effect increases the peak velocity and/or does not prevent the excitatory EBN input from continuing to drive the motor neurons, leading to hypermetric saccades.

### CONCLUSION

There is no need to appeal to various kinds of noise in the brain to explain the variability of saccades.

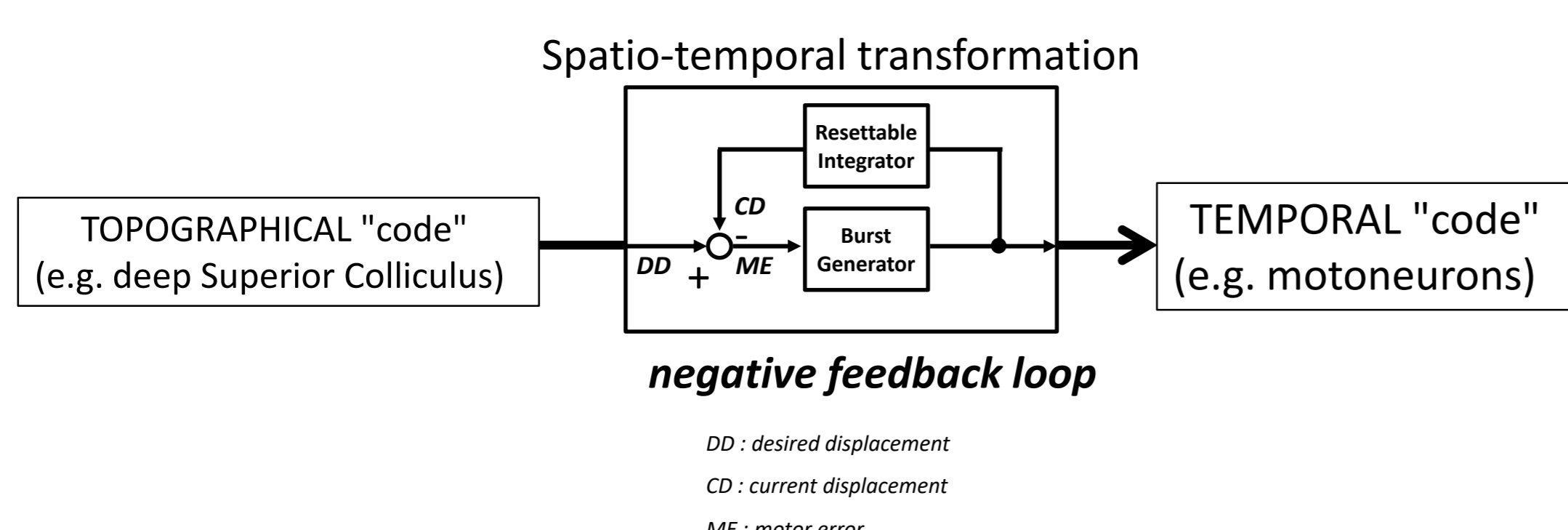
The number of active neurons contributes to the dynamics of saccades

The different evolutions of ipsilesional and contralateral dysmetria suggest that these two deficits are unrelated

Between the topographical and temporal "codes", instead of appealing to a negative feedback loop that reduces a mismatch between internal signals encoding kinematic parameters (desired versus current displacement as estimated from the integration of velocity commands), we propose that a saccade is the outcome of a process that restores symmetry of activities between commands exerting opposite movement tendencies.

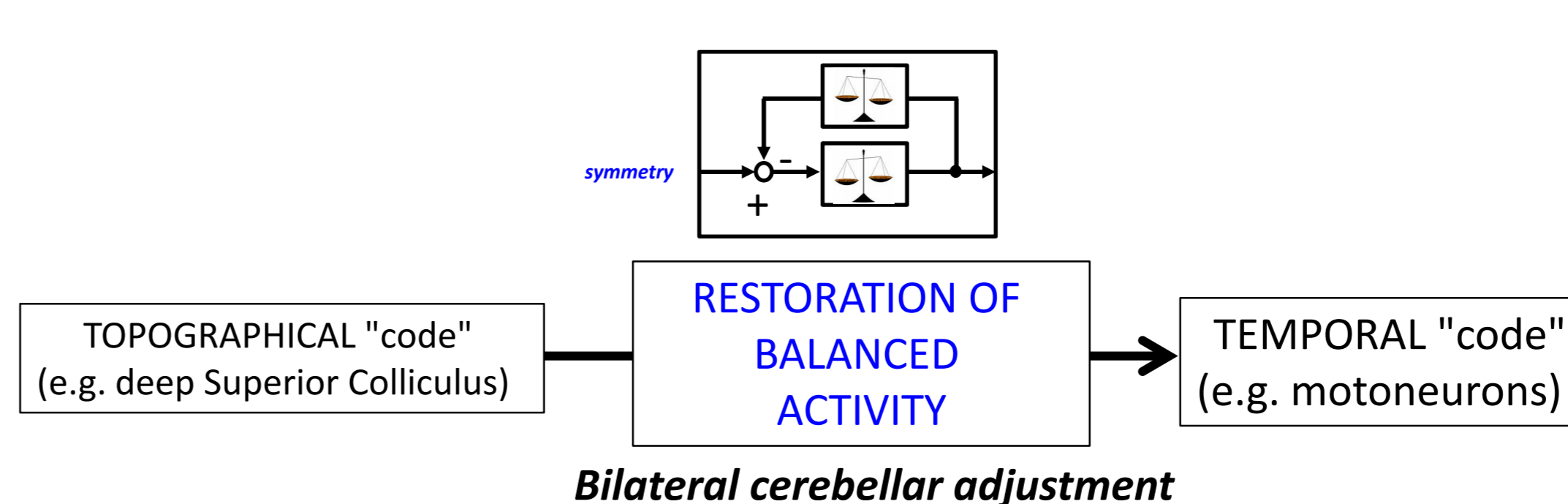
The orienting movement is saccadic for no teleological purpose; it is saccadic merely because the flow of activity from the retinal to the motor neurons involves neurons whose pattern of firing is bursting and also because of fast-twitch properties of extraocular muscles.

### CURRENT FRAMEWORK



DD: desired displacement  
CD: current displacement  
ME: motor error

### ALTERNATIVE FRAMEWORK



Bilateral cerebellar adjustment