Abstract
The transient visual response of feline dorsal lateral geniculate nucleus (dLGN) cells was studied under control conditions and during the application of repetitive transcranial magnetic stimulation at 1 Hz (rTMS@1Hz) on the primary visual cortex (V1). The results show that rTMS@1Hz modulates the firing mode of Y cells, inducing an increase in burst spikes and a decrease in tonic firing. On the other hand, rTMS@1Hz modifies the spatiotemporal characteristics of receptive fields of X cells, inducing a delay and a decrease of the peak response, and a change of the surround/center amplitude ratio of RF profiles. These results indicate that V1 controls the activity of the visual thalamus in a different way in the X and Y pathways, and that this feedback control is consistent with roles associated with each cell type.

Introduction
Early research on the role of visual cortical feedback has suggested both a facilitating [1,2] and an inhibitory effect on thalamic neurons [3,4]. Other studies have attributed different functions to cortical input, such as the ability to synchronize the neurons of the thalamus, to interfere in the spatial processing of the image or to enhance the receptive field (RF) center-surround antagonism of thalamic cells [4-6].

To date a variety of methods have been used to reveal the role of cortical feedback to the dLGN including pharmacological manipulation, cooling and aspiration of cortical tissue. Recently, repetitive transcranial magnetic stimulation (rTMS) has emerged as an alternative method. It is a noninvasive technique, its effects are fast and its application, depending on the stimulation parameters, allows excitation or depression of cortical activity [7–9]. Moreover, it has been shown that rTMS is a valuable tool to study the role of the corticofugal feedback since it is able to transiently reduce the activity of visual cortical neurons [10,11]. For instance, rTMS at the frequency of 1 Hz applied to the visual cortex of the cat, produced a significant reduction in visually driven responses recorded simultaneously in layer 6 of V1 (the source of the feedback) and in the dLGN [10] (see also methods section).

It has been shown that visual cortex influences receptive field organization in the thalamus [12]. An immediate question is whether this feedback is able to modulate the spatio-temporal properties of dLGN receptive fields as could be expected from what has been said so far and from previous data showing that cortico-geniculate feedback seems to improve the temporal accuracy of signal transmission [13,14]. On the other hand, it is already known that corticofugal feedback can influence the firing pattern in dLGN cells [12,13–17] and that thalamic cells in general, including those in the dLGN, exhibit two distinct response patterns, burst and tonic firing. However it is not known whether this influence operates in an unspecific manner at the level of the dLGN or, conversely, it acts differentially depending upon the cell type, which would represent the presence of, at least, two (X and Y) physiologically different descending lines of control.

This paper aims to analyze the contribution of feedback from V1 to the spatio-temporal characteristics of RFs and response mode of dLGN cells during visual stimulation. rTMS was applied on the occipital cortex of anesthetized cats to temporarily block the cortical activity [10]. We have chosen to use a specific type of visual stimulus (pseudo-random sequence of bright and dark bars flashed throughout the receptive field) appropriate to map the spatiotemporal RFs of dLGN cells [10] and valuable to evoke robust transient responses of spike bursts.

Results
A population of 51 dLGN neurons, 18 X, 28 Y and 5 unclassified cells were recorded. All cells RFs were located within 12° of the area centralis.

Burst analysis
We calculated the number of spikes per burst and the percentage of spikes in bursts, and found no significant changes after applying rTMS@1Hz when all the dLGN cells were grouped together (X, Y and unclassified), or by dividing the population by polarity (ON - OFF) or laterality (ipsi-and contralateral). However, significant differences between the X and Y subpopulations were evident, with rTMS affecting Y cells but not X cells.

Burst analysis for X cells
Figure 1A shows the distribution of the percentage of spikes in burst during control condition. Statistical analysis revealed that
rTMS@1Hz did not alter either the burst or tonic response, as it is shown in Figure 1B for spikes in burst.

Burst analysis for Y cells

Burst properties of Y cells were not affected by the application of rTMS. However, differences were found depending on their visual response prior to rTMS application. Figure 2A shows the distribution of the percentage of spikes in bursts for Y cells during control conditions. A visual analysis of Figure 2A shows that there are two populations of neurons that generate bursts with different rate. To corroborate this subdivision of the sample, a two-phase clustering was performed, yielding a cut-off value around 40–50%, which divides the sample into neurons with low and high percentages of bursts in the response shown in Figure 2A (dashed line). In addition, an intra-burst analysis revealed that the subgroup with a high rate in burst response tended to generate bursts with a greater number of spikes than cells with a lower rate (Figure 2B, p<0.05, two samples t-test). This result suggests that these groups could constitute two different populations of neurons, henceforth each group was analyzed separately.

Neurons with a low percentage of bursts, i.e. the proportion of spikes in bursts was below 40–50% (n = 18), showed a significant increase in the percentage of spikes in bursts when rTMS@1Hz was applied, as it is shown in Figure 3A (21.1±2.2% to 24.0±2.2%, P<0.05, paired t-test) and therefore a decrease in the percentage of tonic spikes (78.9±2.2% to 76.0±2.2%, P<0.05, paired t-test, Figure 3B). Considering each burst as an event, rTMS@1Hz application induced an 11.0% increase in the number of bursts (from 143.2±17.7 to 162.5±19.0, P<0.05, paired t-test, Figure 3C).

Figure 3G shows an example of an ON Y cell with a low rate of spikes in bursts during control (29%, black line). Application of rTMS@1Hz increased the percentage of spikes in bursts to 38% (gray line).

Neurons with a high percentage of bursts, i.e. the proportion of spikes in bursts were more than 40–50% of the total response (n = 10), did not show significant changes with rTMS@1Hz application (see Figures 3D–F).

Intervalogram analysis

Only spikes separated in the range from 0 to 20 ms were considered for the analysis because almost all interspike intervals during the transient phase of dLGN cells response fell into this range (see Material and Methods). According to the previous results, intervalogram analyses were developed separately for X and Y subpopulations.

Intervalogram analysis for X cells

Average intervalograms for X neurons population were calculated for both, control condition and when rTMS@1Hz was applied. Figure 4 shows the scatter plot of the difference between these two intervalograms. Dots indicate an increase (red dots) or decrease (blue dots) in the number of a determined interspike interval and dot size represents the degree of change. During the presentation of visual stimuli (vertical black bar on left) the main changes induced by rTMS are associated with a decrease in the number of spikes separated from 1 to 3 ms. Elsewhere in the intervalogram there are more subtle changes generally related to a decrease of interspike intervals.

Intervalogram analysis for Y cells

The sample of Y cells was subdivided depending on their rate of bursts response considering our previous bursts analysis. Y neurons with a low rate of spikes in bursts (smaller than 40–50%) increased the amount of interspike intervals within a range of 1 to 5 ms during rTMS application (Figure 5A). In contrast, neurons with a high percentage of bursts seem to be less affected by rTMS@1Hz with a scatter plot that reflects changes of slighter magnitude (Figure 5B).

Spatiotemporal response analysis

We then studied the spatial and temporal profile of the dLGN cells visual response using the reverse correlation technique. No significant changes were observed when considering the whole...
population (X, Y and unclassified). The analysis was then performed separately on X and Y cells as follows.

Spatiotemporal response analysis for X cells

The application of rTMS@1Hz induced changes in the spatiotemporal response of X cells. Figure 6A is an example of an ON X cell showing a decrease in peak response and an increase in peak latency after applying rTMS@1Hz. The cell also showed a change in the surround/center amplitude ratio (Figure 6B). Figure 7 shows the mean changes induced by rTMS@1Hz in cell responses for the population of X neurons. There is a decrease in the maximum response of 10.9% (from 23.0 ± 1.4 to 20.5 ± 1.4, P < 0.05, paired t-test, Figure 7A), a slight increase in latency (42.0 ± 1.6 to 43.0 ± 1.6, P < 0.05, paired t-test, Figure 7B) and an increase of 5.8% in the surround/center Gaussian amplitude ratio (from 14.9 ± 4.0% to 20.7 ± 5.0%, P < 0.05, paired t-test, Figure 7C).

Spatiotemporal response analysis for Y cells

The application of rTMS@1Hz did not induce significant changes in the response of Y neurons, either taking all the population of neurons or subdividing the sample according to the percentage of bursts (not shown). It is worthy of notice the fact that, in contrast to the X cell population, a group of Y neurons (n = 11) did show a decrease in latency. The remaining neurons presented an increase (n = 9) or no change in this parameter (n = 8). None of these groups was identified as neurons with a high or low percentage of bursts.

Discussion

We have applied rTMS@1Hz to explore the influence of the corticofugal pathway in the transient phase of dLGN cells visual response. We have chosen this low-frequency stimulation protocol because it has been shown to be a fast and reliable method to transiently depress the excitability of the cerebral cortex [10,19] (see example in the method section). Moreover, we have previously used this protocol to demonstrate that corticogeniculate downflow selectively affects the sustained component of the visual response of dLGN cells with no effect on bursts discharge [10]. In the present work we show that the visual cortex exerts a differential control on
burst frequency and spatiotemporal RF organization depending on the cell type. This apparent contradiction may be explained taking into account some differences in the visual stimulation protocol. In previous works, to study the whole visual response of ON dLGN cells, we initially presented a black background and subsequently a white dot centered on the RF, and vice versa for OFF dLGN cells [10]. This proved to be an optimal way to analyze the effect of rTMS in the sustained phase of the response, but it had some disadvantages when considering the transient phase. For a relay dLGN neuron, prolonged exposure to a stimulus with a non-preferred contrast (black for an ON cell) produces a “rebound” response at the end of the stimulus mainly constituted by spikes in bursts [20–22]. These bursts may be driven by the inhibition produced by dLGN interneurons whose receptive field position coincides with the relay neuron, but with an opposite polarity (ON or OFF interneurons) [22]. Thus, an initial black background presented to an ON dLGN neuron preconditions their Vm to a more hyperpolarized level, by disynaptic inhibition via OFF interneurons, which in turn favors the generation of bursts once the visual stimulus appears. In the present study special care was taken to maintain a stable state in the inhibition that directly depends on retinal input [22]. To do this, a gray background (mean luminance of 45 cd/m²) for black and white bars series was used to keep the same background activity for interneurons and relay cells.

To analyze spikes in bursts, we applied the criteria based on intracellular recordings in cat dLGN neurons [23]. These are the most widely accepted criteria for burst analysis of extracellular recordings [12,24–26]. Additionally, we tested a more restrictive selection to prevent the inclusion of high frequency tonic spikes, considering each burst as a train comprised at least of three spikes [10,17,27]. Finally, we applied both criteria but considering 6 ms as the maximum intraburst interspike interval [28]. All criteria gave similar results which reinforces our conclusions.

Effects of rTMS on transient phase response

Previous results [29] showed that cortical cooling induces a decrease in the peak response of the transient phase of ON X cells (a decrease of 18%) whilst the peak value of Y cells remained unchanged. Our study adds new information, showing that the cortex selectively influences the spatiotemporal response of X neurons producing, together with a decrease in peak response, a slight increase in peak latency of around 1 ms; both phenomena are probably associated with an imbalance in the center/periphery amplitude ratio. Moreover, burst analysis showed that Y cells, but not X cells, increase their burst rate and reduce tonic spikes when the cortex is inactivated. This behavior was observed in those Y cells characterized by having a burst rate at control conditions of less than 40–50%. This splits our Y cell sample into two groups that also differ in their spikes per burst distribution. Since the number of spikes per burst increases at higher levels of hyperpolarization [30], all these data indicate that cortical blockade influences primarily neurons with some prior level of depolarization. Thus, when cortical input is reduced, dLGN neuron’s Vm becomes more negative, increasing the number of spikes per burst. Hence, it is reasonable to hypothesize that Y cells with a high response rate in bursts reflect a lower pre-rTMS cortical activity, a condition in which the TMS seems to generate less effect on the cortex [11]. As far as we know, this clustering of Y cells has not been reported previously, although a study focused on dLGN cells burstiness suggests certain subdivision of Y cells depending on the "burst fraction" of their response [31].

Differences of cortical effects between X and Y cells could be due to some differences of connectivity patterns in both pathways. Besides the fact that both groups of neurons receive inputs from different types of retinal neurons, there is evidence that the X pathway could be exposed to higher local inhibitory control than the Y pathway [32,33] and that each pathway could have independent interneurons contacting only with the associated relay neurons [34,35]. Moreover, in visual cortex both pathways remain partially segregated in V1; reviewed in [36], but see [37]. These results suggest the existence of a difference in corticothalamic connectivity, as it has been recently shown by others in monkey [38].

Regarding corticothalamic synapses, neuroanatomical studies have revealed that axons coming from area 17 make contact with

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**Figure 4.** Scatter plot obtained subtracting average intervalogram of rTMS from control condition for X dLGN cells. Blue and red dots represent decrease and increase of interspike intervals (X axis) along the time (Y axis) respectively. Dot size depicts the magnitude of change and black bar shows the duration of visual stimulus. Asterisk indicates the interspike interval range for spikes in burst.

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A: Y dLGN cells with spikes in burst percentage < 40–50 %, n = 18

B: Y dLGN cells with spikes in burst percentage > 40–50 %, n = 10

Figure 5. Scatter plot for Y cells. Scatter plot of changes on intervalograms induced by rTMS@1Hz for Y dLGN cells with low (A) and high (B) content of bursts.
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Figure 6. Example of a dLGN X-on cell whose STRF profile was modified by rTMS@1Hz. Circles represent real values normalized to control and curves their mathematical fits. A, Transient response elicited by a white bar presented at the RF center shows a decrease in peak value (from 80 to 60%) and an increase in latency (from 39 to 41 ms) when rTMS was applied. B, Construction of RF profile by fitting a difference of Gaussians (DoG) function to real values. rTMS@1Hz modified the rate between Gaussian amplitudes by a decrease of the center (upper thin curve) and an increase of the periphery (lower thin curves).
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Figure 7. Effect of rTMS on X cells. rTMS@1Hz decreased (A) and delayed (B) the peak transient response and induced an increase in the rate between periphery and center amplitudes (C) in dLGN X-cells (bin = 2 ms). Vertical error bars represent the standard error of the mean. doi:10.1371/journal.pone.0017041.g007

Experimental preparation
Six adult cats of either sex were prepared following standard procedures used in our laboratory for extracellular recordings in the visual pathway [17,46]. Briefly, animals were anesthetized with isoflurane (5% for induction, 1.5–2% for surgery, and 0.5–1% for maintenance) in nitrous oxide (70%) and oxygen (30%). The trachea was cannulated, an intravenous line inserted, and a craniotomy performed for thalamic recordings. To prevent eye movements, animals were paralyzed with gallamine triethiodide (loading dose of 40 mg, maintenance 10 mg/kg/h i.v.) and held in a stereotaxic frame. End-tidal CO₂ levels, electrocardiogram (EKG) waveform and intersystolic interval, and the electro-
cientific procedures were performed with the appropriate permit from the University of A Coruña Ethics Committee (CE-UDC30/1/09). The study was approved by the University of A Coruña Ethics Committee (CE-UDC30/1/09).

**Magnetic stimulation**

The rTMS was carried out with a MagStim Rapid system (The MagStim Company Ltd, Whitland, UK) equipped with 2 boosters and applied on the occipital cortex of cats via a figure-of-eight coil (mean diameter: 25 mm). The midpoint of the coil was centered over the thalamus (Horsley-Clark antero-posterior: 0 to –6, Medio-lateral: 0) directly touching the exposed bone. The coil was fixed by a mechanical arm at an angle of about 60° (wings located laterally) with the handle pointing up and backwards. For this study, all rTMS parameters were optimized to produce cortical suppression in the region of cortex below the coil [10,47–49]. The stimulation frequency utilized was one pulse per second, applied during a period of two minutes, hereinafter referred to as the rTMS@1Hz protocol. The intensity was set to 50% which, according to the data supplied by the manufacturer (http://www.magstim.co.uk), induces 1.5 T on the cortical surface (3 mm from the coil), and an electric field strength of 220 V/m, higher than those reported in other studies [50] but which effect, shown in the results, is consistent with those described by these authors.

The statistical significance of the changes induced by magnetic stimulation was determined by using paired and unpaired t-test or Wilcoxon test. The hypothesis of normality was checked using the Kolmogorov-Smirnov test. Results were deemed to be significant when P<0.05. Cluster analysis was carried out with the two-phase modality and by applying the Akaike criterion.

**Recording procedures**

Tungsten microelectrodes (FHC) of 8–12 MΩ, mounted on a micromanipulator (FHC, Brunswick, ME) were introduced in the dLGN following stereotaxic coordinates (Horsley–Clark A6 L9). Neurons responding to visual stimulation of both eyes (bimedular) with big RFs were classified as perigeniculate nucleus cells and were not considered. Neurons giving only monocular responses with a defined RF were considered for further recording and analysis. Cell type was identified, X or Y, on the basis of standard tests including the test of linearity of spatial summation, size and eccentricity of the receptive field, type of response to points of light, and presence of shift effect [51–54]. The visual response of the cell was recorded during four minutes: two minutes for control condition followed by two minutes for the simultaneous application of the rTMS@1Hz protocol. The entire protocol was run once per cell. The inter-recording interval was dependent upon the time devoted to search, isolate and characterize each cell, but usually was more than one hour.

**Visual stimulus**

The one-dimensional version of the reverse correlation algorithm [18,55–58] (see below Reverse correlation analysis) was applied to evaluate the dLGN cells visual response. This protocol allows the quick generation of a RF map during the application of rTMS (the period was limited by the activation of the temperature-sensitive protection of the equipment). The transient response elicited in dLGN neurons was used to characterize the spatiotemporal dynamics of the RF as well as its response mode (see Burst analysis). In particular, the visual stimulus consisted in two groups of 20 white and 20 dark bars presented on a gray background and in a pseudorandom sequence. Each group of bars was arranged orderly and centered on the RF cell under study, as it is shown in Figure 8A. Stimulation was adapted to the preferred spatial frequency of the cells by changing the bar width. Bars had a length of 12° and a width of 0.4° and 1°, covering an area of +/- 6° of elevation and +/- 4° of azimuth respectively. Each bar was displayed for 40 ms with no direct correlation with the magnetic stimulation.

**Reverse correlation analysis**

A diagram summarizing the procedure we used is shown on figure 8. Visual stimuli were displayed in an area centered over the dLGN cell’s RF (Fig. 8A). Each spike was associated with the visual stimulus that preceded it for an interval of 200 ms, then a peristimulus time histogram (PSTH) for each bar was generated (Fig. 8B). PSTHs were ordered according to the position of the visual stimulus that elicited each one and normalized with respect to the maximum total value. As a result, spatial response profiles across the receptive field were obtained (Fig. 8C), both for ON stimulus (white bars) and OFF stimulus (black bars). A composite spatiotemporal receptive field (STRF) was then obtained subtracting the OFF from the ON response (Fig. 8D). Thus, for an ON cell, regions of the STRF excited by the bright and dark bars are shown as positive and negative values respectively. Figure 8E-F shows the spatial and temporal profiles obtained by slicing the STRF through the peak value as it is shown by arrows in Fig. 8D. Filled circles in Fig. 8E-F represent real values and solid curves show the mathematical fit for the spatial (difference of two Gaussian functions) [18,59–61] and temporal (difference of two Gamma functions) [18,62,63] profiles. This process was applied to all STRFs to quantitatively characterize and compare different recordings. All curve fittings were performed with the cftool function (MATLAB version R2006b, The MathWorks).

**Control experiments (visual cortical responses to local rTMS)**

To gain insight into the effect of rTMS on V1, in previous control experiments we were able to record visual-driven activity simultaneously from cortical and dLGN cells during rTMS stimulation at 1 Hz. Electrodes to record cortical spikes were either advanced during the experiment at an oblique angle (avoiding the stimulating coil) or cemented in place (tip located approximately in layer VI) prior to positioning of the TMS coil.

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**TMS Effects on Transient Response of LGN Cells**

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Visual stimulus was the same one used to study dLGN cells; the orientation of the bars was the optimal for each cortical cell. We did not pretend to systematically map the spatial extent of the cortical suppression zone; however, it is likely, given the size of the cat brain and the spatial resolution of rTMS, that our stimulation effectively covered the primary visual cortex and that the effect was uniform across this region. The histogram in figure 9 shows the visual response of both a dLGN neuron and a deep layer cortical cell (presumably a layer 6 neuron), before (blue line), during rTMS (red line) and after rTMS (green line).

Intervalogram analysis

The intervalogram method was introduced by Funke and Worgotter [44] to study the sustained phase of cat dLGN cell responses. Basically, an intervalogram shows the variation of the distribution of interspike intervals in the cell response. Figure 10A shows schematically the response of a cell to one bar displayed on the RF center. Firstly, in a time window of 50 ms (see below) starting at $t = 0$, a first interspike interval histogram [INTH] was obtained for the spikes within the window. The second INTH was obtained for a time window displaced 5 ms after the first window (5 ms corresponds to the 10% of the total duration of the window according to the criterion of Funke and Worgotter) [44]. Remaining INTH’s were obtained moving the window every 5 ms to complete the total period of analysis (250 ms). Arranging INTH’s in a two dimensional array allowed us to detect changes on interval distribution over time whenever the cell was stimulated. Due to the visual stimulus properties, intervalograms illustrate changes during the transient phase of the response. Since each visual stimulus generated an intervalogram, an average intervalogram for a given number of stimuli was obtained. Figure 10B shows an average intervalogram for a Y OFF cell obtained after the presentation of 72 black bars on the RF center. Figure 10C shows the first mean INTH of that intervalogram, obtained immediately after the onset of the visual stimulus. The cell responded preferentially with spikes separated 3 ms (arrow). Figure 10D shows the mean 9th INTH, calculated from a time window starting at the end of the visual stimulus (40 ms); even though the preferred frequency response is maintained, the 9th INTH is more distributed than the first INTH i.e., spikes with longer intervals begin to appear.

In general, when intervalograms are used, some details must be taken into account. There is a compromise between the time window size and the characteristics of the stimulus used. A large time window has the advantage of sampling a higher number of spikes and thus generating INTHs with a higher resolution, but it is not optimal to analyze transient responses. In contrast, smaller time windows generate intervalograms with a greater temporal resolution, but with fewer spikes used to compute each INTH. We chose 50 ms because it was optimal to analyze the transient phase of dLGN cell responses. Regarding the analysis of intervalograms, due to the fact that each point in the intervalogram represents how many times a value of interspike interval is repeated in a certain time window, the peak value of intervalograms before and after applying rTMS can move in a 2D plane i.e., a cell can either change its preferred firing frequency, or the instant it reaches the peak, or both parameters at once. The latter is a problem when applying statistics and it is the reason to read intervalograms in a qualitative way.

Burst analysis

Burst analysis was carried out to unveil changes in the response mode of dLGN neurons. Trains of action potentials preceded by a silence period of at least 30 ms and whose interspike intervals were
**Figure 9. Example of the effect of rTMS on cortical cells.** Top panel: rTMS stimulation produced a reduction in the peak response (red curve) compared to the pre rTMS values (blue curve). Green curve depicts the recovery of the response (bin = 4 ms). The cell was recorded in the deeper layers of V1. The effect of rTMS on a dLGN cell recorded simultaneously is also shown (bottom panel).

**Figure 10. Construction of intervalograms.** A. Spikes elicited by a dark bar (VS) presented on RF center of a Y off dLGN cell. Interval distributions are computed for small time windows of 50 ms shifted 5 ms each other along the time axis. An inter-spike interval histogram (INTH) is calculated for each window (1, 2, 3, … 40) by adding all interval distributions obtained from each dark bar presented on RF center of the dLGN cell. INTHs are plotted as a colour-scaled horizontal pixel line and all the intervalogram is normalized to its maximum value. B. Average intervalogram for all Y dLGN cells. The temporal resolution for the horizontal axis is 1 ms and the maximum value is limited by the time window, i.e. 50 ms. The temporal resolution for the vertical axis is determined by the interval between time windows, i.e. 5 ms and maximum value is 200 ms. C. First INTH of the average intervalogram showed in B. D. INTH from the average intervalogram at the end of the VS.
less than 4 ms were considered as bursts [23]. All other spikes were considered as tonic response.

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Author Contributions

Conceived and designed the experiments: NE JM CL JC. Performed the experiments: NE JM CL JC. Analyzed the data: NE JM CL JC. Wrote the paper: NE JM CL JC.
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