Dynamics of Intrinsic Dendritic Calcium Signaling during Tonic Firing of Thalamic Reticular Neurons

Patrick Chausson, Nathalie Leresche, Régis C. Lambert

To cite this version:

Patrick Chausson, Nathalie Leresche, Régis C. Lambert. Dynamics of Intrinsic Dendritic Calcium Signaling during Tonic Firing of Thalamic Reticular Neurons. PLoS ONE, 2013, 8 (8), pp.e72275. 10.1371/journal.pone.0072275 . hal-01348949

HAL Id: hal-01348949
https://hal.science/hal-01348949
Submitted on 26 Jul 2016

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L’archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d’enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.
Dynamics of Intrinsic Dendritic Calcium Signaling during Tonic Firing of Thalamic Reticular Neurons

Patrick Chausson1,2, Nathalie Leresche1,2, Régis C. Lambert1,2*

1 UMR 7102 CNRS, Paris, France, 2 UPMC, Université Paris 6, Paris, France

Abstract
The GABAergic neurons of the nucleus reticularis thalami that control the communication between thalamus and cortex are interconnected not only through axo-dendritic synapses but also through gap junctions and dendro-dendritic synapses. It is still unknown whether these dendritic communication processes may be triggered both by the tonic and the T-type Ca\(^{2+}\) channel-dependent high frequency burst firing of action potentials displayed by nucleus reticularis neurons during wakefulness and sleep, respectively. Indeed, while it is known that activation of T-type Ca\(^{2+}\) channels actively propagates throughout the dendritic tree, it is still unclear whether tonic action potential firing can also invade the dendritic arborization. Here, using two-photon microscopy, we demonstrated that dendritic Ca\(^{2+}\) responses following somatically evoked action potentials that mimic wake-related tonic firing are detected throughout the dendritic arborization. Calcium influx temporally summates to produce dendritic Ca\(^{2+}\) accumulations that are linearly related to the duration of the action potential trains. Increasing the firing frequency facilitates Ca\(^{2+}\) influx in the proximal but not in the distal dendritic compartments suggesting that the dendritic arborization acts as a low-pass filter in respect to the back-propagating action potentials. In the more distal compartment of the dendritic tree, T-type Ca\(^{2+}\) channels play a crucial role in the action potential triggered Ca\(^{2+}\) influx suggesting that this Ca\(^{2+}\) influx may be controlled by slight changes in the local dendritic membrane potential that determine the T-type channels’ availability. We conclude that by mediating Ca\(^{2+}\) dynamic in the whole dendritic arborization, both tonic and burst firing of the nucleus reticularis thalami neurons might control their dendro-dendritic and electrical communications.

Citation: Chausson P, Leresche N, Lambert RC (2013) Dynamics of Intrinsic Dendritic Calcium Signaling during Tonic Firing of Thalamic Reticular Neurons. PLoS ONE 8(8): e72275. doi:10.1371/journal.pone.0072275

Copyright: © 2013 Chausson et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Competing Interests: The authors have declared that no competing interests exist.

* E-mail: regis.lambert@snv.jussieu.fr

Introduction
Back-propagation of action potentials (APs) in the dendritic tree has been extensively studied in a number of neurons where it has been shown to determine the neuronal firing pattern, to contribute to dendritic integration and to support long-term and spike-timing dependent plasticity [1]. Moreover, by evoking widespread Ca\(^{2+}\) signals throughout the dendritic arborization, back-propagating APs are also likely candidate for dendritic neurotransmitter release at dendro-dendritic synapses [2] (see also review by [3]). The physiological consequences of dendro-dendritic synapse activation have been analyzed in details in the olfactory bulb [4] and, although far less investigated, dendro-dendritic synapses involving GABAergic neurons have also been described in the thalamus. In the lateral geniculate nucleus, interneurons express GABAergic vesicles in dendritic appendages and a majority of the interneuron synapses on the thalamocortical neurons are made by dendritic boutons [5,6]. Combining whole-cell recordings and either two-photon Ca\(^{2+}\) imaging [7] or voltage-sensitive dye imaging [8], it has been shown that APs of interneurons back-propagate with high fidelity through the dendrites contributing to GABA release and feedforward inhibition of thalamocortical neurons.

In the Nucleus Reticularis Thalami (NRT), a GABAergic nucleus that controls the communication between the thalamus and the cortex and plays a crucial role in the generation of the synchronized activities within the thalamocortical loop during sleep, neurons are interconnected not only through axo-dendritic synapses but also through gap junctions and dendro-dendritic synapses [9,10]. NRT neurons present two modes of discharge according to the state of vigilance. During wakefulness, NRT neurons discharge tonically but they switch to a high frequency bursting mode underlined by a low-threshold Ca\(^{2+}\) spike (LTS) during low vigilance and sleep [11]. Generation of this LTS is due to the recruitment of the T-type Ca\(^{2+}\) channels following their de-inactivation by hyperpolarization. It has been recently shown that activation of T-type Ca\(^{2+}\) channels underlying the high frequency burst firing actively propagates throughout the dendrites suggesting that the bursting mode of firing may support dendro-dendritic communication [12]. However, whether tonic AP firing can also invade the dendritic arborization remains unclear.

Here, using two-photon microscopy, we investigated the dynamics of intrinsic dendritic Ca\(^{2+}\) signaling across the NRT dendritic tree. We found that dendritic Ca\(^{2+}\) responses following somatically evoked APs can be detected in the dendritic arborization of the NRT neuron in thalamic slices at physiological temperature. In the more distal compartment of the dendritic tree, T-type Ca\(^{2+}\) channels play a crucial role in the action potential triggered Ca\(^{2+}\) influx suggesting that this Ca\(^{2+}\) influx may be controlled by slight changes in the local dendritic membrane potential that determines the T-type channel availability. There-
Figure 1. Dendritic Ca^{2+} responses evoked by tonic APs or LTS in NRT neurons. A1. Dendritic ΔCa^{2+} triggered by somatically evoked train of 15 APs at 10 Hz (top record). Application of 0.5 μM TTX fully blocked the APs and the ΔCa^{2+} (bottom trace). A2. Dendritic ΔCa^{2+} triggered by a somatically evoked LTS. An enlarged record of the LTS is presented in inset. Application of 3 μM TTA-P2 fully blocked the LTS and the ΔCa^{2+}. B. ΔCa^{2+} recorded in three different dendrites of the same neuron. B1. Dendritic ΔCa^{2+} evoked by LTS (left column) and 1, 3, 5 and 10 APs at 10 Hz (right column) in the different dendrites. Examples of the somatically recorded voltage responses are shown underneath the ΔCa^{2+} recorded in location 1 (an enlarged record of the LTS response is illustrated in inset). ΔCa^{2+} recorded in response to a single back-propagating AP is presented at higher magnification in inset. B2. Plot of amplitude of ΔCa^{2+} evoked by back-propagating APs (n = 10 neurons). The amplitude was normalized to the ΔCa^{2+} evoked at the time of the 20th AP of the trains. Note the almost linear summation of the ΔCa^{2+} evoked by the first 5 back-propagating APs. In A and B the scanned dendritic regions are indicated by boxes on the maximal intensity Z projection of the neurons presented in the left panels.
doi:10.1371/journal.pone.0072275.g001
fore, we conclude that both tonic and burst firing trigger intracellular Ca\textsuperscript{2+} increase throughout the NRT neuron arborization potentially linking neuronal firing to the dendritic integration and communication processes.

**Methods**

**Ethical Approval**

Ethical approval was obtained for all experimental protocols from the Departmental Direction of Veterinary Services, Paris. All procedures involving experimental animals were carried out in accordance with the EU Council Directive 86–609. Every effort was made to minimize animal suffering and the number of animals used. For removal of tissues, animals were deeply anesthetized with inhaled isoflurane and immediately sacrificed.

**Preparation of Brain Slices and Recordings**

Brains were excised from 12–18 day old Wistar rats. A block of tissue containing the thalamus was removed, placed in a cold (<4°C) oxygenated (95%O\textsubscript{2}/5%CO\textsubscript{2}) solution of artificial cerebrospinal fluid (aCSF) (in mM): 125 NaCl, 2.5 KCl, 0.4 CaCl\textsubscript{2}, 1 MgCl\textsubscript{2}, 1.25 Na\textsubscript{2}HPO\textsubscript{4}, 26 NaHCO\textsubscript{3}, 25 glucose, and 1 kynurenic acid (pH 7.3; osmolarity 310 mOsm). The block of tissue was glued, ventral surface uppermost, to the stage of a vibroslice (Leica VT1200S), and 220–300 μm thick horizontal slices containing the ventrobasal nucleus and the NRT were prepared using the internal capsule and the medial lemniscus as landmarks. Slices were stored in an oxygenated incubation chamber containing aCSF of the above composition, but without kynurenic acid and with 2 mM CaCl\textsubscript{2}, for at least 1 h before being transferred to the recording chamber, where they were perfused (2.5 ml/min) continuously with an oxygenated recording solution of the same composition. Experiments were conducted at 32°C.

Using the patch-clamp technique (Axopatch 200B; Clampex 10, Molecular Devices), whole-cell recordings in current clamp mode were performed in NRT neurons visualized with an Olympus BX51WI (x60 lens). Recordings were filtered by a 4-pole Bessel filter set at a corner frequency of 1 kHz and digitalized at 10 kHz and latter analyzed using Matlab R2009b (The MathWorks, Inc.).

Electrodes were filled with the following solution (in mM): 140 methanesulfonic acid, 4 MgCl\textsubscript{2}, 10 HEPES, 4 Na-ATP, 15 phosphocreatine, 150 units/ml creatine phosphokinase; pH 7.3, osmolarity 280 mOsm. Alexa Fluor 594 and Oregon Green 488 Bapta-1 (Molecular Probes) were added with a final concentration osmolarity 280 mOsm. Alexa Fluor 594 and Oregon Green 488 delivered at 20–30 s intervals.

**Calcium Imaging**

Images were obtained with a custom-built 2-photon laser scanning microscope. Two-photon excitation of the fluorescent dyes was performed by a femtosecond Ti:Saphir laser (Mai Tai HP, Spectra-Physics, Mountain View, CA) tuned to 800 nm. Image acquisition was controlled by MPScope software [13]. Before imaging, neurons were loaded with indicators for 30 min. The fluorescent signals from Alexa Fluor 594 and Oregon Green were acquired simultaneously across dendrites at selected regions of interest (ROI) (acquisition frequency 40–60 frame/s) by two high gain photomultiplier tubes (PMT Hamamatsu H9305-03, Hamamatsu photonics, Japan). At the end of the recordings Z series of 160–190 images (512 pixels, 0.26 μm/pixel) were taken with 0.5 μm focal steps to construct a two-dimensional maximum intensity projection of each neuron.

**Data Analysis**

Recordings were analyzed off-line using Matlab R2009b (The MathWorks, Inc.). Morphological distances were approximated by measuring along the dendrite from the beginning of the dendrite to the ROI of interest on the two-dimensional maximum intensity projection of each neuron. Fluorescence signals were measured by integration of the signal over a region of interest. The reported change in fluorescence (ΔG/R) was calculated as the change in fluorescence (Gpeak) from baseline (G0, average of the 400 ms period before stimulus) of the Ca\textsuperscript{2+}-sensitive indicator (Oregon Green 480 Bapta-1) normalized to the average fluorescence of the Ca\textsuperscript{2+}-insensitive indicator (Ravg, Alexa Fluor 594): ΔG/R = (Gpeak−G0)/Ravg [14]. The peak response was calculated as the combined average of 10–20 trials to maximize signal-to-noise ratio.

Quantitative data in the text and figures are given as mean ± s.e.m. Two-tailed t-tests were used to compare the values of ΔCa\textsuperscript{2+} evoked by LTSs. As action potentials ΔCa\textsuperscript{2+} values do not follow a normal distribution (Shapiro test), a non-parametric Wilcoxon-Mann Whitney test was used to assess significance. Differences were considered significant for p<0.05.

**Drugs**

TTA-P2, 3,5-Dichloro-N-[4-(2,2-dimethyl-tetrahydro-pyran-4-ylmethyl)-4-fluoro-piperidinx-4-ylmethyl]-benzamide, was made up as 10 mM stock solution in dimethyl sulphoxide, kept at −20°C until use [15,16]. TTA-P2 was provided by Merck and Co., Inc. TTX was obtained from Latoxan, 6-cyano-7-nitroquinoxaline-2,3-dione (CNQX) from Sigma.

**Results**

Combining whole-cell patch-clamp recordings with two-photon laser-scanning microscopy, we investigated NRT neuron Ca\textsuperscript{2+} dynamics in response to somatically generated tonic APs and LTSs-elicited high-frequency burst discharges. Burst discharges were triggered from a membrane potential of −75 to −80 mV by short current injections of minimal amplitude. Similarly, trains of tonic APs were evoked by injecting brief depolarizing current pulses at a frequency of either 10 or 40 Hz from a membrane potential set between −50 and −60 mV where T-type Ca\textsuperscript{2+} channels are almost fully inactivated (see methods). Individual neurons were loaded through the patch pipette with both the Ca\textsuperscript{2+}-sensitive fluorophore and a Ca\textsuperscript{2+}-insensitive indicator to visualize the dendritic tree.

Not only burst discharges but also single AP or trains of APs were able to transiently raise the intracellular Ca\textsuperscript{2+} concentration (ΔCa\textsuperscript{2+}) in every NRT neuron dendritic branch (Figure 1B). Blocking sodium channels with TTX, or the T-type Ca\textsuperscript{2+} channels with the specific antagonist TTA-P2 [15] resulted in a full block of the ΔCa\textsuperscript{2+} evoked by single/tonic APs or burst firing, respectively (Figure 1A). As illustrated in Figure 1B, single AP triggered a small but clear increase in fluorescence in every dendrites that were monitored on the same neuron. The AP associated ΔCa\textsuperscript{2+} were present in both primary and secondary dendrites (Figure 1B1 and...
Figure 2. Back-propagating APs produce robust ΔCa\(^{2+}\) in distal dendrites of the NRT neurons. A. Top traces: ΔCa\(^{2+}\) recorded in response to a somatically evoked burst of 45 APs at 10 Hz (left trace). Amplitude of the ΔCa\(^{2+}\) first increased in the most proximal part of the dendrite (compare 10 and 40 mm) before decreasing in the more distal dendritic compartment. Note however that a noticeable ΔCa\(^{2+}\) is still present at the more distal location (120 mm). Bottom traces: ΔCa\(^{2+}\) recorded in response to a somatically evoked LTS (left trace). Note the increase in ΔCa\(^{2+}\) with dendritic distance. B. Summary of ΔCa\(^{2+}\) grouped by dendritic location. LTS: n = 19, 13 and 8; APs: n = 21, 15 and 6 for distances <60, 60–120 and >120 μm, respectively. *, p<0.05, n.s.: p>0.05. C1. ΔCa\(^{2+}\) recorded in response to a somatically evoked burst of 45 APs at 10 (black traces) and 40 Hz (red traces). Increasing the firing frequency produced a clear enhancement in ΔCa\(^{2+}\) at proximal and intermediate locations but not at the more distal...
see also Figure 2C) and dendritic Ca\(^{2+}\) influxes temporally summated during 10 Hz trains of APs. Quantification of \(\Delta\text{Ca}^{2+}\) showed a linear relationship for the first 4–5 APs, reaching a plateau level around the 20th AP of the train (Figure 1B2). This plateau was not due to the saturation of the Ca\(^{2+}\) sensitive dye since a LTS-elicited burst discharge always evoked a larger fluorescence increase at the same dendritic location.

To assess the extent of AP back-propagation in the dendritic tree of NRT neurons, the magnitude of the responses evoked by trains of APs at increasing distances from the soma was measured (Figure 2). Over the first tens of microns of the dendrite (60 μm), the amplitude of the \(\Delta\text{Ca}^{2+}\) evoked by 10 Hz trains of APs either increased by 36±19% (n = 5, Figure 2), reaching a maximum between 20 and 40 μm, or remained stable (n = 8), when compared to the most proximal signal measured around 10 μm. In 6 neurons Ca\(^{2+}\) influxes could be monitored in both proximal and distal regions (120 to 200 μm) along the same dendrite. Although the AP evoked Ca\(^{2+}\) influx decreased in the distal dendrite, the Ca\(^{2+}\) response was still 54±10% of the maximal proximal response (Figure 2A). This mean value reflects a heterogeneous behavior as out of these 6 neurons, 4 of them presented a fairly small attenuation of \(\Delta\text{Ca}^{2+}\) with a distal signal corresponding to 67±9% of the Ca\(^{2+}\) influx observed in the proximal compartment while in the 2 remaining neurons, this percentage dropped to 35 and 23%, respectively. On average pooled data from all our results grouped by dendritic location (180 μm). C2. Summary of \(\Delta\text{Ca}^{2+}\) evoked in the same neurons by trains of 45 APs at 10 and 40 Hz grouped by dendritic location, <60 μm: n = 8; 60–120 μm: n = 7; >120 μm: n = 4. **: p<0.01, *: p<0.05, n.s.: p>0.05. D. Contribution of sodium APs to LTS-evoked dendritic \(\Delta\text{Ca}^{2+}\). D1. LTS (left traces) and associated dendritic \(\Delta\text{Ca}^{2+}\) (middle traces) measured at 110 μm from the soma in control condition and after TTX application. After blockade of the sodium APs, note the clear increase in amplitude and duration of the LTS and the lack of change in dendritic \(\Delta\text{Ca}^{2+}\). An enlargement of \(\Delta\text{Ca}^{2+}\) in both conditions is presented in inset. Right traces: at the same dendritic location a clear \(\Delta\text{Ca}^{2+}\) was evoked by a train of 45 APs at 10 Hz. Therefore, the lack of changes in LTS-evoked \(\Delta\text{Ca}^{2+}\) following TTX application did not result from the inability of APs to back-propagate in this distal dendrite. D2. Top plot shows the maximal fluorescence signal evoked in distal dendrites by LTS in control condition and after TTX application for each neuron. Note that TTX has no clear effect on calcium entry (mean ΔG/R: control 1.2±0.1, TTX: 1.08±0.13, p>0.05). Bottom plot presents the duration of LTSs in control condition and after TTX application (control: 76.1±2.8 ms, TTX: 107.4±5.6 ms, p>0.05). In A and C the scanned dendritic regions are indicated by boxes on the maximal intensity Z projection of the neurons.

doi:10.1371/journal.pone.0072275.g002

Figure 3. Contribution of T-type current to AP back-propagation. A. APs (A1) and associated dendritic \(\Delta\text{Ca}^{2+}\) (A2) recorded in control condition and after TTA-P2 application (3 μM). Same neuron as in figure 2A. Note that TTA-P2 did not modify the tonic discharge of APs. Enlargements of the first and last APs of the train are presented in insets. In contrast, the block of the T-type current decreased the amplitude of \(\Delta\text{Ca}^{2+}\) and almost fully abolished Ca\(^{2+}\) entry in the distal dendritic region. B1–2. Examples of two neurons in which TTA-P2 had no effect on the \(\Delta\text{Ca}^{2+}\) recorded in proximal dendrites. Same calibration as in A2. C. Plots of the maximal fluorescence signal evoked by trains of APs in control condition and after TTA-P2 application for each neuron. The left graph shows that TTA-P2 has little effect on calcium entry monitored in proximal dendrites (mean ΔG/R: control 0.31±0.04, TTA-P2:0.25±0.04, p>0.05). Right graph: same measurements performed on distal dendrites showed a clear decrease upon TTA-P2 application (mean ΔG/R: control 0.25±0.7, TTA-P2 0.11±0.05, p<0.005).
doi:10.1371/journal.pone.0072275.g003
(n = 24 neurons, Figure 2B), clearly show that trains of APs can back-propagate up to the distal dendritic compartment in NRT neurons maintained at physiological temperature.

Increasing the frequency of firing to 40 Hz enhanced the ΔCa\textsuperscript{2+} plateau measured at proximal and intermediate dendritic locations (20–60 μm, 152±5% n = 8; 60–120 μm, 197±7% n = 7) (Figure 2C). As predicted from the temporal summation of Ca\textsuperscript{2+} influxes during AP trains, the increase in firing frequency also induced a faster Ca\textsuperscript{2+} accumulation (mean time constant: 742±120 ms and 278±40 ms for 10 and 40 Hz, respectively; n = 6). Surprisingly, accelerating tonic firing frequency from 10 to 40 Hz had no clear effect on the amplitude of ΔCa\textsuperscript{2+} in the more distal area (>120 μm; 104±1%; n = 4; Figure 2C) suggesting a strong low-pass filtering of the dendritic tree for back-propagating AP trains.

As previously reported by Crandall et al., [17], LTS mediated burst firing produced robust Ca\textsuperscript{2+} responses that increased along the dendrite (Figure 2A, 2B). In the proximal part of the dendrites (<60 μm) the ratio of ΔCa\textsuperscript{2+} due to trains of 10 Hz action potentials versus LTS could be close to 1 or even larger in some neurons (1.21±0.12; n = 5), while in the remaining neurons it was around 0.5 (0.45±0.04; n = 15). However due to the inverse spatial relationship between the magnitudes of ΔCa\textsuperscript{2+} evoked by an LTS and back-propagating APs, respectively, this ratio rapidly dropped to a value of 0.2 for dendritic distances greater than 60 μm (0.16±0.02; n = 18), increasing the difference between LTS- and AP-associated Ca\textsuperscript{2+} dynamics in the distal dendritic compartments.

Since trains of APs can back-propagate up to distal dendritic regions we then tested whether the high frequency bursts of APs contribute to the overall ΔCa\textsuperscript{2+} observed during LTS mediated firing. Although during bath application of 0.5 μM TTX, ΔCa\textsuperscript{2+} tend to decrease by 15.2±5.2 and 10.8±4.2% in the proximal (n = 5) and distal (n = 7) dendritic regions, respectively, these decreases were not significant (p > 0.05) (Figure 2D). However the TTX effects are complex and blocking AP firing resulted in a larger and prolonged LTS (48.7±4.8% increase in total duration, n = 7; Figure 2D). Such change in LTS waveform is likely due to the suppression of the strong AP after-hyperpolarization, may affect LTS associated ΔCa\textsuperscript{2+} and impair isolation of the specific Ca\textsuperscript{2+} entry due to back-propagating APs.

As 1-type Ca\textsuperscript{2+} channels are assumed to be the major source of Ca\textsuperscript{2+} entry in the dendrites of the NRT neurons [17], we then investigated whether activation of these channels may also support the Ca\textsuperscript{2+} signal associated with the back-propagation of APs in distal dendrites. To test this hypothesis, we compared the back-propagation of AP trains in control condition and in the presence of TTA-P2. TTA-P2 did not modify the tonic firing evoked by trains of short current pulses and had no discernible effect on either the AP waveform or its after-hyperpolarization (Figure 3A1). In the more proximal dendrites TTA-P2 had either little or no effect in 5 out of 7 neurons (Figure 3B, 3C) while in the 2 remaining neurons, the ΔCa\textsuperscript{2+} was decreased by 30 and 40% (Figure 3A2, 3C). When considering the whole neuronal population TTA-P2 induced a mean non-significant decrease of 19.1±5.7% of the Ca\textsuperscript{2+} entry in the proximal dendrites (n = 7).
contrast, although at our holding potential set between −50 and −60 mV the T-type Ca\(^{2+}\) channel population should be almost fully inactivated, the selective T-type current antagonist markedly decreased ΔCa\(^{2+}\) monitored at dendritic location more distal than 80 μm (63.8±9.3%, n = 6) (Figure 3A2, 3C).

During the course of this study we patch-clamped three neurons in which no LTS could be evoked by injecting a depolarizing current step from hyperpolarized membrane potential (−80 to −90 mV) or after strong long-lasting hyperpolarizations that maximally recruit the T-type channels (Figure 4). However, a short train of tonic APs at a frequency around 10 Hz was observed at the break of the hyperpolarizing pulse associated to a clear ΔCa\(^{2+}\). Both the ΔCa\(^{2+}\) and the rebound tonic firing were blocked by TTA-P2, demonstrating that T-type Ca\(^{2+}\) currents are present in non-bursting NRT neurons where they participate to the generation of a tonic rebound activity. These neurons did not present a peculiar morphology when compared to other NRT neurons or a specific localization within the NRT.

Discussion

The main finding of this study is that a tonic discharge of APs that mimics the firing of NRT neurons during wakefulness evokes a transient rise in intracellular Ca\(^{2+}\) that can be detected up to the distal regions of their dendrites (200 μm) and that T-type channels play a crucial role in this distal dendritic Ca\(^{2+}\) influx.

Using a similar approach, Crandall et al [17] previously reported that a somatically-evoked LTS produces a transient Ca\(^{2+}\) response along the NRT neuronal dendrites, that increases in magnitude at greater distances from the soma (up to 200 μm). However, in contrast to our data, single, or trains of, APs produced a minimal Ca\(^{2+}\) influx that was restricted to the most proximal dendrites (15 to 25 μm). A main difference between these two studies is the temperature at which experiments were performed, with Crandall and al. [17] being carried out at room temperature while we used a more physiological temperature (32–34°C). It is well known that temperature drastically modifies the biophysical properties of ionic channels, including the T-type Ca\(^{2+}\) channels [18], and therefore should condition both active and passive properties of the dendritic tree with major consequences on its ability to back-propagate APs and/or to generate Ca\(^{2+}\) influxes. In addition, different tonic firing patterns were used in the two studies. Crandall et al. [17] investigated the back-propagation of a short train of 4 APs at 100 Hz that mimics the burst of APs crowning an LTS rather than the tonic firing of NRT neurons. During waking, NRT neurons have a mean firing rate of 7–22 Hz in head-restrained cat [19] and of about 40 Hz in freely moving cats [20]. Moreover, in anesthetized cats, NRT neurons fire tonically at a frequency of 30 Hz in response to sensory stimulation [21]. To match the firing of NRT neurons during wakefulness, our protocols consisted of 15 to 45 APs at a frequency of either 10 or 40 Hz. Using these protocols, we showed that increasing the firing frequency facilitates Ca\(^{2+}\) influx in the proximal but not in the distal dendritic compartment suggesting that the dendritic arborization acts as a low-pass filter for back-propagating APs. This filtering ability should also dampen the propagation of short high-frequency trains of APs such as the ones used in Crandall et al., [17]. Accordingly we did not observe any consistent decrease in Ca\(^{2+}\) influx when blocking the high frequency burst of APs with TTX during a somatically evoked LTS. However, this result should be considered with caution since blocking APs firing induces marked changes in the LTS waveform that may mask a decrease in ΔCa\(^{2+}\) due to the loss of the high-frequency back-propagating APs.

In agreement with previous studies [22,23,24], we report that a small proportion of NRT neurons does not display high-frequency burst. Such inability to trigger LTS following somatically imposed membrane hyperpolarization is likely due to a reduced density of T-type channels in these cells, at least in their somatic and proximal dendritic compartments. However, transient rebound tonic activities, fully blocked by TTA-P2, were observed in these neurons demonstrating the presence of a minimal amount of T-type channels. This surprising rebound tonic activity exceeds the duration of T-type current mediated depolarization suggesting the activation of an additional transient depolarizing conductance, such as the Ca\(^{2+}\) activated non-selective cationic (CAN) current [25,26]. Nevertheless, nearly all NRT neurons express a high density of T-type channels and our results indicate that back-propagation of APs depends in part upon activation of these channels since a clear decrease of ΔCa\(^{2+}\) was observed following TTA-P2 application. The contribution of T-type current to the Ca\(^{2+}\) influx associated to AP back-propagation was not constant along the entire dendritic tree since the TTA-P2 effect was more pronounced in distal than in proximal/intermediate dendritic compartment. This spatially gradual effect may reflect an attenuation of the amplitude of the AP-evoked depolarization along the dendrite that would become too small to activate high-voltage activated Ca\(^{2+}\) currents in the distal dendritic compartments. Alternatively, the expression of the high-voltage activated Ca\(^{2+}\) channels may be restricted to the more proximal dendrites and the T-type channels may constitute the only active Ca\(^{2+}\) channels expressed in distal dendritic branches. Accordingly, morphological data confirm the dense distribution of T-type channels on the dendritic membrane of NRT neurons [27], and experimental and computational data suggested a higher density of T-type currents in distal dendrites [20]. The contribution of T-type channels to the Ca\(^{2+}\) influx evoked by back-propagating APs is at first glance surprising since tonic activity was triggered from a holding potential set between −50 to −60 mV where T-type channels are thought to be almost fully inactivated [29,30,31]. However the membrane potential of the distal dendritic compartment may not be fully controlled by the somatic electrode [32]. As a consequence, a hyperpolarizing gradient of the membrane potential may exist between the soma and the distal dendrites with a similar increasing gradient of deinactivated T-type channels. Nevertheless, because in some NRT neurons T-type currents clearly participated to the Ca\(^{2+}\) influx in the first tens microns of the dendritic branch (see Figure 3A2), it is unlikely that the T-type current activation can be fully explained by a lack of control of the dendritic membrane potential by our somatic electrode. We have previously demonstrated that, due to the high-density of T-type channels expressed in thalamic neurons, the small fraction of deinactivated channels at depolarized potential is responsible for a physiologically meaningful current. Indeed, although the open probability of the channels is low at −60 mV, the number of available channels in both thalamocortical and NRT neurons is sufficient to generate a window current that contributes to the resting membrane potential of these neurons [15]. Moreover, in thalamocortical neurons maintained at depolarized potentials, we recently demonstrated that the drastic increase in open probability induced by sensory excitation inputs induces a significant T-type current that boost the post-synaptic excitation potential [33]. Therefore, back-propagating APs may similarly recruit the deinactivated channels population that is present at depolarized potentials in the NRT neurons to generate a clear Ca\(^{2+}\) influx.
should be closely controlled by both excitatory and inhibitory synaptic inputs that influence local dendritic potential. Conversely, dendritic Ca$^{2+}$ influxes triggered by APs may control integration of the incoming inputs in NRT neurons. Indeed, among the classical roles attributed to back-propagating APs is their function in synaptic plasticity. Glutamatergic synapses impinging on NRT neurons with an organized pattern, as thalamocortical inputs mainly innervate the soma and proximal dendrites [34,35,36] while the corticothalamic synapses mostly contact the intermediate or distal dendritic compartments [37]. GABAergic synapses arising from intra-NRT connections impinge on the whole dendritic arborization [38,39]. Therefore, by showing clear Ca$^{2+}$ influxes not only in the proximal and intermediate dendritic regions but also in the more distal dendritic branches, our results suggest that back-propagating APs evoke local signals at corticothalamic, thalamocortical and intra-NRT GABAergic post-synaptic areas that may trigger short or long-term modification. In addition, as highlighted in the introduction, back-propagating APs could not only contribute to the integration of incoming synaptic inputs but may also participate to the activation of dendro-dendritic synapses [2,3]. Finally, NRT neurons are strongly coupled by electrical synapses composed of connexin36 proteins [40]. It was recently shown that L/Ts induce an LTD of the electrical coupling between NRT neurons [41]. Although delayed and of reduced amplitude, this LTD could also be induced by paired sodium spikes evoked at depolarized potentials, and it is highly likely that the increase in intracellular Ca$^{2+}$ through back-propagating APs is required to trigger plasticity of the electrical synapses. Therefore, by mediating Ca$^{2+}$ dynamic in the whole dendritic arborization, tonic firing of NRT neurons may control their dendro-dendritic and electrical communications as well as the integration of their synaptic inputs.

Acknowledgments

We wish to thank Drs. V.N. Urbele and J.J. Renger from Merck and Co., Inc for their gift of TTA-P2.

Author Contributions

Conceived and designed the experiments: NL RCL. Performed the experiments: PC RCL. Analyzed the data: PC NL RCL. Wrote the paper: NL RCL PC.

References

1. Kampa BM, Letzkus JJ, Stuart GJ (2007) Dendritic mechanisms controlling spiketriggering-dependent synaptic plasticity. Trends in neurosciences 30: 436–463.
2. Egger V, Svoboda K, Mainen ZF (2003) Mechanisms of lateral inhibition in the olfactory bulb: efficiency and modulation of spike-evoked calcium influx into granule cells. J Neurosci 23: 7531–7550.
3. Kennedy MJ, Ehlers MD (2011) Mechanisms and function of dendritic excytosis. Neuron 69: 856–875.
4. Lagier S, Panzaneli P, Russo RE, Nissant A, Bathellier B, et al. (2007) GABAergic inhibition at dendrodendritic synapses tunes gamma oscillations in the olfactory bulb. Proceedings of the National Academy of Sciences of the United States of America 104: 7259–7264.
5. Famiglietti EV Jr (1970) Dendro-dendritic synapses in the lateral geniculate nucleus of the cat. Brain research 20: 181–191.
6. Montero VM (1986) Localization of gamma-aminobutyric acid (GABA) in type 3 cells and demonstration of their source to F2 terminals in the cat lateral geniculate nucleus: a Golgi-electron-microscopic GABA-immunocytochemical study. The Journal of comparative neurology 234: 228–245.
7. Ascone-Goycoolea C, Benowitz SD, Regehr WG (2008) Active dendritic conductances dynamically regulate GABA release from thalamic interneurons. Neuron 57: 420–431.
8. Casale AE, McCormick DA (2011) Active action potential propagation but not initiation in thalamic interneuron dendrites. J Neurosci 31: 18329–18332.
9. Pinault D, Smith V, Doscheus M (1997) Dendrodendritic and axoaxonic synapses in the thalamic reticular nucleus of the adult rat. J Neurosci 17: 3213–3233.
10. Montero VM (1984) Ultrastructure and synaptic relations of neural elements containing glutamic acid decarboxylase (GAD) in the perigeniculate thalamic neurons and their modulation by light. Neuroscience 104: 247–264.
11. Steriade M, McCormick DA, Sejnowski TJ (1993) Thalamocortical oscillations and synchronization in neocortex. J Neurosci 13: 456–474.
12. Crandall SR, Cox CL (2001) Local dendrodendritic inhibition regulates fast dendritic spines. Neuron 33: 439–452.
13. Williams SR, Stuart GJ (2000) Action potential backpropagation and somatodendritic signaling in NRT Neurons. J Neurosci 32: 2513–2522.
14. Nguyen QT, Tsai PS, Kleinfeld D (2006) MIPScope: a versatile software suite for multiphoton microscopy. Journal of neuroscience methods 156: 351–359.
15. Sakhare BL, Oertner TG, Svoboda K (2002) The life cycle of Ca(2+)/ions in dendritic spines. Neuron 33: 439–452.
16. Dreyfus EM, Teichert A, Errington AC, Renger JG, Shih HS, et al. (2010) Selective T-type calcium channel block in thalamic neurons reveals channel redundancy and physiological impact of IT/widow. J Neurosci 30: 99–109.
17. Shipke WB, Bawor JC, Yang ZQ, Lindsey CW, Yang FY, et al. (2000) Design, synthesis, and evaluation of a novel 4-aminoethyl-4-thiophosphorodine as a T-type Ca$^{2+}$ channel antagonist. J Med Chem 43: 6992–6995.
18. Crandall SR, Goverdiah G, Cox CL (2010) Low-threshold Ca$^{2+}$ current amplitudes distal dendritic signaling in thalamic reticular neurons. J Neurosci 30: 15419–15429.
19. Ilincu M, McKay BE, Snutch TP, McRory JE, Turner RW, et al. (2006) Temperature dependence of T-type calcium channel gating. Neuroscience 142: 1031–1042.
20. Barioniuco G, Benoit O, Tempier P (1981) Evidence for two types of firing pattern during the sleep-waking cycle in the reticulata thalamic nucleus of the cat. Experimental neurology 72: 496–501.
21. Amzica F, Nunez A, Steriade M (1992) Delta frequency (1–4 Hz) oscillations in thalamic reticular neurons and their modulation by light. Neuroscience 51: 285–294.
22. Contreas D, Carro Dossi R, Steriade M (1992) Bursting and tonic discharges in two classes of reticular thalamic neurons. J Neurophysiol 68: 973–977.
23. Brunton J, Charpak S (1997) Heterogeneity of cell firing properties and opioid sensitivity in the thalamic reticular nucleus. Neuroscience 78: 303–307.
24. Bartha BH, Nunes ML (2007) Evaluation of sleep habits in children with epilepsy. Epilepsy & behavior : E&B 11: 60–64.
25. Bal T, McCormick DA (1993) Mechanisms of oscillatory activity in guinea-pig nucleus reticularis thalami in vitro: a mammalian pacemaker. J Physiol (Lond) 461: 669–691.
26. Bledtyn KL, Hughes SW, Toth TI, Cope DW, Crunelli V (2006) Neuronal basis of the slow (<1 Hz) oscillation in neurons of the nucleus reticularis thalami in vitro. J Neurosci 26: 2474–2486.
27. Liu XD, Murray KD, Jones EG (2011) Low-threshold calcium channel subunit Cav3.3 is specifically localized in GABAAergic neurons of rodent thalamus and cerebral cortex. The Journal of comparative neurology 519: 1181–1193.
28. Destexhe A, Contreras D, Steriade M, Sejnowski TJ, Huguenard JR (1996) In vivo, in vitro, and computational analysis of dendritic calcium currents in thalamic reticular neurons. J Neurosci 16: 169–185.
29. Huguenard JR, Prince DA (1992) A novel T-type current underlies prolonged Ca$^{2+}$-dependent burst firing in GABAAergic neurons of rat thalamic reticular nucleus. J Neurosci 12: 3094–3107.
30. Jokovic PM, Cossey DF, Todorovic SM (2007) Inhibition of T-type calcium current in the reticular thalamic nucleus by a novel neuroactive steroid. Annals of the New York Academy of Sciences 1122: 83–94.
31. Perez-Reyes E (2003) Molecular physiology of low-voltage-activated t-type calcium channels. Physiol Rev 83: 117–161.
32. Williams SR, Stuart GJ (2000) Action potential backpropagation and somato-dendritic distribution of ion channels in thalamocortical neurons. J Neurosci 20: 1307–1317.
33. Delercq EZ, David F, Behuret S, Sadoc G, Shiin HS, et al. (2012) T-type calcium channels consolidate tonic action potential output of thalamic neurons to neocortex. J Neurosci 32: 12228–12236.
34. Montero VM (1989) Ultrastructural identification of synaptic terminals from cortical axons and from collateral axons of geniculocortical relay cells in the perigeniculate nucleus of the cat. Experimental brain research Experimentelle Hirnforschung Experimentation cerebrale 75: 65–72.
35. Ohara PT (1988) Synaptic organization of the thalamic reticular nucleus. Journal of electron microscopy technique 10: 283–292.
36. Ohara PT, Lieberman AR (1985) The thalamic reticular nucleus of the adult rat: experimental anatomical studies. Journal of neurocytology 14: 365–411.
37. Liu XR, Jones EG (1999) Predominance of corticothalamic synaptic inputs to thalamic reticular nucleus neurons in the rat. The Journal of comparative neurology 414: 67–79.
39. de Biasi S, Frassoni C, Spreafico R (1986) GABA immunoreactivity in the thalamic reticular nucleus of the rat. A light and electron microscopical study. Brain research 399: 143–147.

40. Landisman CE, Long MA, Beierlein M, Deans MR, Paul DL, et al. (2002) Electrical synapses in the thalamic reticular nucleus. J Neurosci 22: 1002–1009.

41. Haas JS, Zavala B, Landisman CE (2011) Activity-dependent long-term depression of electrical synapses. Science 334: 389–393.