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RESEARCH ARTICLE

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NKCC1 controls GABAergic signaling and neuroblast migration in the postnatal forebrain

Sheyla Mejia-Gervacio^{1,2*}, Kerren Murray^{1,2}, Pierre-Marie Lledo^{1,2*}

Abstract

From an early postnatal period and throughout life there is a continuous production of olfactory bulb (OB) interneurons originating from neuronal precursors in the subventricular zone. To reach the OB circuits, immature neuroblasts migrate along the rostral migratory stream (RMS). In the present study, we employed cultured postnatal mouse forebrain slices and used lentiviral vectors to label neuronal precursors with GFP and to manipulate the expression levels of the Na-K-2Cl cotransporter NKCC1. We investigated the role of this Cl⁻ transporter in different stages of postnatal neurogenesis, including neuroblast migration and integration in the OB networks once they have reached the granule cell layer (GCL). We report that NKCC1 activity is necessary for maintaining normal migratory speed. Both pharmacological and genetic manipulations revealed that NKCC1 maintains high [Cl⁻]_i and regulates the resting membrane potential of migratory neuroblasts whilst its functional expression is strongly reduced at the time cells reach the GCL. As in other developing systems, NKCC1 shapes GABA_A-dependent signaling in the RMS neuroblasts. Also, we show that NKCC1 controls the migration of neuroblasts in the RMS. The present study indeed indicates that the latter effect results from a novel action of NKCC1 on the resting membrane potential, which is independent of GABA_A-dependent signaling. All in all, our findings show that early stages of the postnatal recruitment of OB interneurons rely on precise, orchestrated mechanisms that depend on multiple actions of NKCC1.

Background

The subventricular zone (SVZ) and the rostral migratory stream (RMS) contain neural progenitors that proliferate and migrate to the olfactory bulb (OB) throughout life [1]. In the OB, the precursors differentiate into two classes of interneurons: granule cells (GCs) and periglomerular cells (PGCs) [2-4], which populate deep and superficial layers of the OB, respectively.

The migration of neuroblasts in the RMS is a highly regulated process, and may determine the rate of integration of newborn neurons in the OB. A series of chemical factors have been identified as regulators of neuroblast migration in the RMS, including adhesion molecules [5], extracellular matrix molecules [6], and molecules with repulsive/attractive functions [7-9]. Moreover, migration in the SVZ-OB system is sensitive to the action of neurotransmitters [10,11], providing a basis for neural activity-dependent regulation of

postnatal neurogenesis. In this regard, a previous study showed that neuroblast migration in the RMS is reduced by the tonic depolarizing action of gamma-aminobutyric acid (GABA) acting on GABA_A receptors [12,13].

GABA_A receptor activity affects the cell membrane potential, producing depolarizing or hyperpolarizing influences depending on the Cl⁻ gradient across the cell membrane, principally determined by the intracellular concentration of Cl⁻ ([Cl⁻]_i) [14-16]. [Cl⁻]_i is tightly regulated by the activity of ionic transporters such as Na⁺-K⁺-2Cl⁻ (NKCC1) or K⁺-Cl⁻ (KCC2) [17-19], which, following the driving force of Na⁺ and K⁺, have the tendency to increase or decrease, respectively, the [Cl⁻]_i at rest.

NKCC1 is frequently expressed in developing central nervous system regions and its activity has been associated with depolarizing actions of GABA [19-23]. It is presently unknown if NKCC1 regulates the [Cl⁻]_i in the RMS. This is an important issue since, in several regions, depolarizing GABA signaling appears early in development and can modulate neuronal excitability to promote neuronal maturation [21,24-27].

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In the present study, we investigated whether NKCC1 shapes the action of GABA on the maturation of OB neuronal precursors and its influence on cellular motility. Our results indicate that the activation of GABA_A receptors induces a strong positive driving force in migrating neuroblasts, which diminishes upon arrival of the cells at the GC layer (GCL). We identified NKCC1 as the uptake mechanism maintaining elevated [Cl⁻]_i in migrating neuroblasts, thus shaping the action of GABA_A receptors. Moreover, we report that the activity of NKCC1 is necessary to maintain normal migration speed and this process occurs independently of GABA_A-signaling since NKCC1 controls the resting state of excitability in RMS neuroblasts.

Results

Forebrain organotypic slice cultures

We developed an organotypic slice culture model that maintains the anatomical organization of the SVZ-OB system (Figure 1A) to study the migration and the first steps of integration of neuronal precursors in the OB circuit at early postnatal periods. For this study, the slices were obtained from 7-day-old mice and maintained for up to 12 days *in vitro* (div). Immunohistochemical staining performed at 6 div with antibodies directed against a neuronal precursor marker (doublecortin (DCX); $n = 3$ slices; Additional file 1) and more mature neuronal markers (NeuN, tyrosine hydroxylase (TH) and GABA; $n = 3$ slices for each; Additional files 1 and 2) were in agreement with the expected distribution patterns observed at similar postnatal ages, as visually compared to the Allen Institute for Brain Science atlas for P14 animals (and P7 when available) for TH and DCX. Similarly, our labeling corresponded to previously published data from adult slices for GABA in the RMS [12] and NeuN in the OB [28,29]. Together, these observations suggested that overall OB organization was preserved in our slice model.

A total of 132 cultured slices were transduced at 1 div with lentiviral vectors to induce the expression of GFP and to manipulate their levels of NKCC1 expression in SVZ precursor cells. After 1 to 2 days post-injection (dpi), strong GFP labeling was visible at the injection site (Figure 1A,B). Several GFP⁺ cells were distributed along the RMS shortly after vector injection (>3 dpi; Figure 1C). These labeled cells displayed typical morphology of migrating neuroblasts, with a small and elongated cell body (6 to 8 μ m), including leading and trailing processes [30]. The first GFP⁺ cells reached the GCL 5 days after vector injection (Figure 1D). Six days after injection, scattered cells were observed throughout the bulb, including in the glomerular layer (Figure 1E). After reaching the appropriate layer, the morphology of GFP⁺ cells changed, with cells transforming into more

mature OB interneurons including mainly GC and less numerous PGCs.

The use of lentiviral vectors precludes precise birth-dating of newborn neurons. Nonetheless, from morphological and physiological points of view, the population of neuroblasts studied was homogeneous during the course of our analysis (around 10 days). We used the measurement of input membrane resistance (IR) as an index of cell maturation [31] and recorded GC in addition to RMS neuroblasts for comparison. In the RMS, we did not find a correlation between the IR and the number of dpi ($n = 15$ from 11 slices; $P = 0.36$; Additional file 3A), while a significant decrease in IR over time ($P = 0.05$) was seen for GFP⁺ cells recorded in the GCL ($n = 21$ from 19 slices; Additional file 3B), consistent with the expected maturation of newborn GCs.

Efficacy of a short hairpin RNA for NKCC1

To test the potential effects of NKCC1 on neuroblast migration and their electrophysiological properties, we used two different strategies. The first consisted of the pharmacological blockade of NKCC1 using the broad-spectrum blocker bumetanide [32,33]. To check for potential unspecific effects of bumetanide, we repeated all our measurements in cells transduced with a well-characterized RNA interference sequence to knock down the expression of NKCC1 [21,23,34].

To validate the use of the short hairpin RNA (shRNA) sequence designed and characterized by Ge *et al.* [21], we performed immunohistochemical staining for NKCC1 on RMS neuroblasts. We compared the staining in cells transduced with either a lentiviral vector encoding GFP and a non-encoding control sequence (hereafter called the control) or a vector encoding GFP and the shRNA for NKCC1 (hereafter called shNKCC1). Two sets of experiments were done. First, we compared the number of neuroblasts coexpressing GFP and NKCC1 in slices transduced with either the control or the shNKCC1 encoding sequences after 1 and 6 dpi (Figure 2). Second, we compared the number of GFP⁺/NKCC1⁺ cells in both the RMS and the GCL at 6 dpi transduced with either one of the sequences (Additional file 4). In both sets of experiments, we found a significant difference between groups using the Kruskal-Wallis test: values for each experiment were $H = 33.82$ ($P = 0.1 \times 10^{-4}$) and $H = 38.13$ ($P = 0.1 \times 10^{-4}$). Subsequent Mann-Whitney comparisons between the groups showed a significantly higher proportion of neuroblasts positive for NKCC1 in slices transduced with the control sequence compared to those transduced with shNKCC1 at 6 dpi: values changed in the first experiment from 0.69 ± 0.05 ($n = 85$ cells) to 0.13 ± 0.07 ($n = 41$ cells), $P = 1 \times 10^{-3}$ and in the second experiment from 0.80 ± 0.09 ($n = 58$ cells) to 0.25 ± 0.11 ($n = 63$ cells), $P = 1 \times 10^{-4}$.

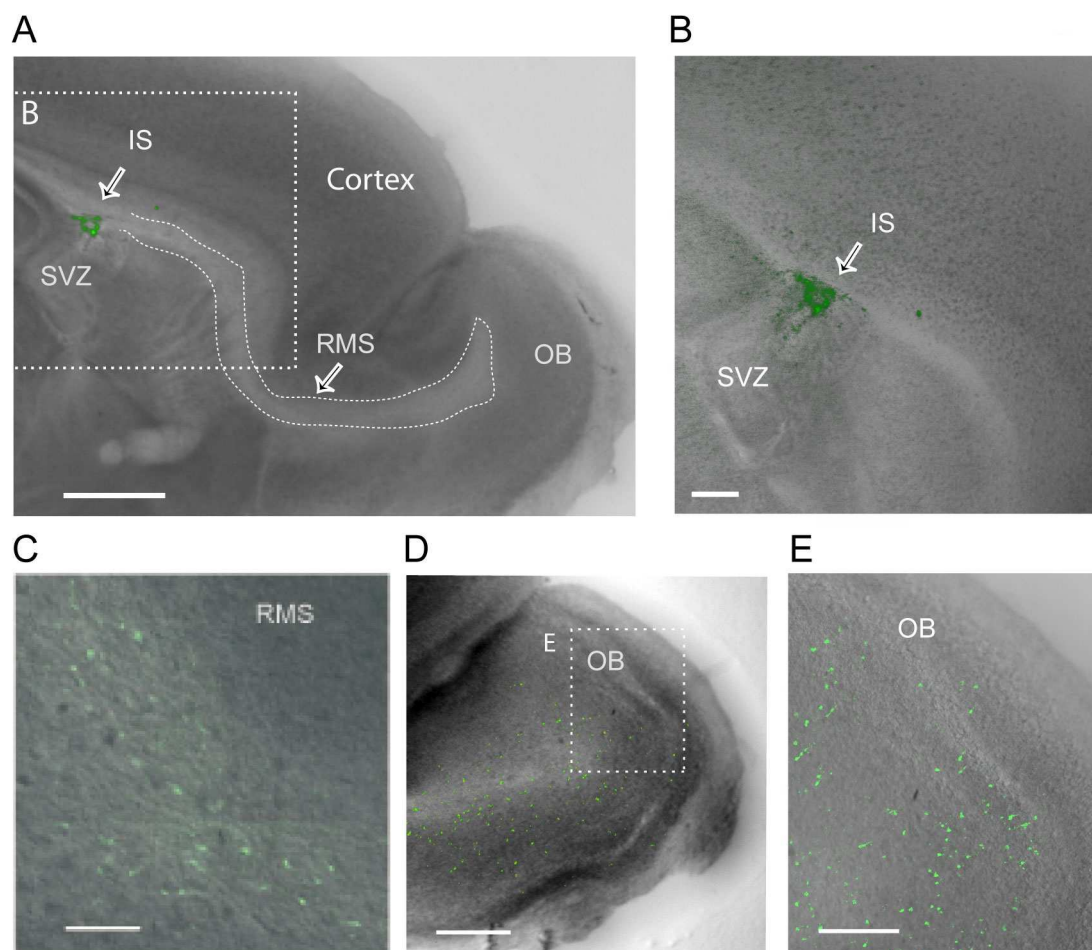


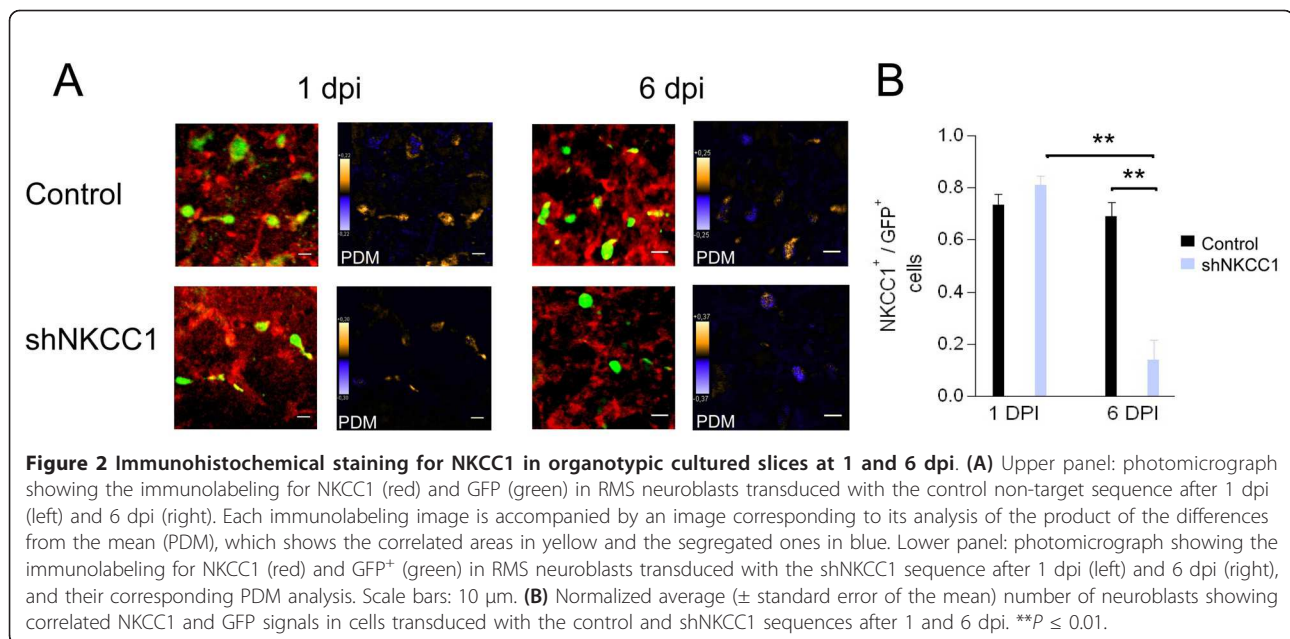
Figure 1 The injection of lentiviral vectors in the SVZ of organotypic slice cultures gives rise to GFP⁺ migrating neuroblasts in the RMS and labeled neurons in all OB layers. **(A)** Photomicrograph of a cultured mouse forebrain sagittal section containing the SVZ-RMS pathway, showing the viral injection site (IS) in the SVZ at 2 days post-injection (dpi). Scale bar: 800 μ m. Inset. Approximate location of the image shown in (B). **(B)** Detail of the IS displayed in (A) at higher magnification. Scale bar: 200 μ m. **(C)** Photomicrograph showing GFP⁺ cells migrating in the RMS at 5 dpi. Scale bar: 100 μ m. **(D)** Photomicrograph of the entire OB showing the arrival of GFP⁺ cells at 5 dpi. Scale bar: 500 μ m. The outline represents the approximate location of the magnification in (E). **(E)** Photomicrograph of the area outlined in (D), magnified to show GFP⁺ cells scattered throughout all OB layers. The same slice culture used in (D) is shown at 11 dpi. Scale bar: 200 μ m.

In contrast, the proportion of NKCC1⁺ GCs was small in control slices (0.22 ± 0.05 ; $n = 27$ cells) and did not significantly change in shNKCC1 slices at 6 dpi (0.05 ± 0.03 ; $n = 17$ cells; $P = 0.09$). In slices injected with the shNKCC1-encoding vector, we observed a significant reduction in the proportion of GFP⁺/NKCC1⁺ neuroblasts when comparing results at 1 and 6 dpi (from 0.81 ± 0.03 to 0.13 ± 0.07 ; $n = 22$ and 41 cells; $P = 1 \times 10^{-3}$). Meanwhile, no difference was observed between 1 and 6 dpi in slices injected with the control vector (0.73 ± 0.03 and 0.69 ± 0.05 ; $n = 67$ and 85 cells; $P = 0.25$). The significance reported in this section takes into account the adjustment for P -values using the Bonferroni method.

Our immunohistochemical studies show that the shRNA interference strategy used in this study is efficient to reduce the expression of NKCC1 in neuroblasts, and coincides with its previously proven efficacy in the hippocampus [21,34] and the cortex [23]. Moreover, our results show that the expression of NKCC1 is reduced when the newborn cells leave the RMS to integrate the GCL.

NKCC1 activity is necessary to maintain normal migration speed in RMS neuroblasts

We explored whether the diminution in the activity of NKCC1 could impair the migration of neuroblasts in the RMS. For this, we compared the tangential migration of cells transduced with the non-target shRNA-GFP



(control) against control cells in slices incubated in 10 μ M bumetanide 1 hour before and during the time lapse recording. A similar approach was employed with cells transduced with the shNKCC1-GFP virus. We calculated the average migration speed and the total displacement (ΔX) along the horizontal axis of the RMS in 202 cells (from 9 slices) for controls; 61 cells (4 slices) incubated in bumetanide, and 83 cells (from 8 slices) transduced with shNKCC1, in 7 independent experiments. Control neuroblasts migrated with an average speed of 42.2 ± 2.0 μ m/h. The migration speed became significantly slower in cells incubated in bumetanide (32.4 ± 4.7 μ m/h; $P = 0.02$) or transduced with shNKCC1 (31.9 ± 3.1 μ m/h, $P = 0.007$) (Figure 3A; Additional files 5 and 6).

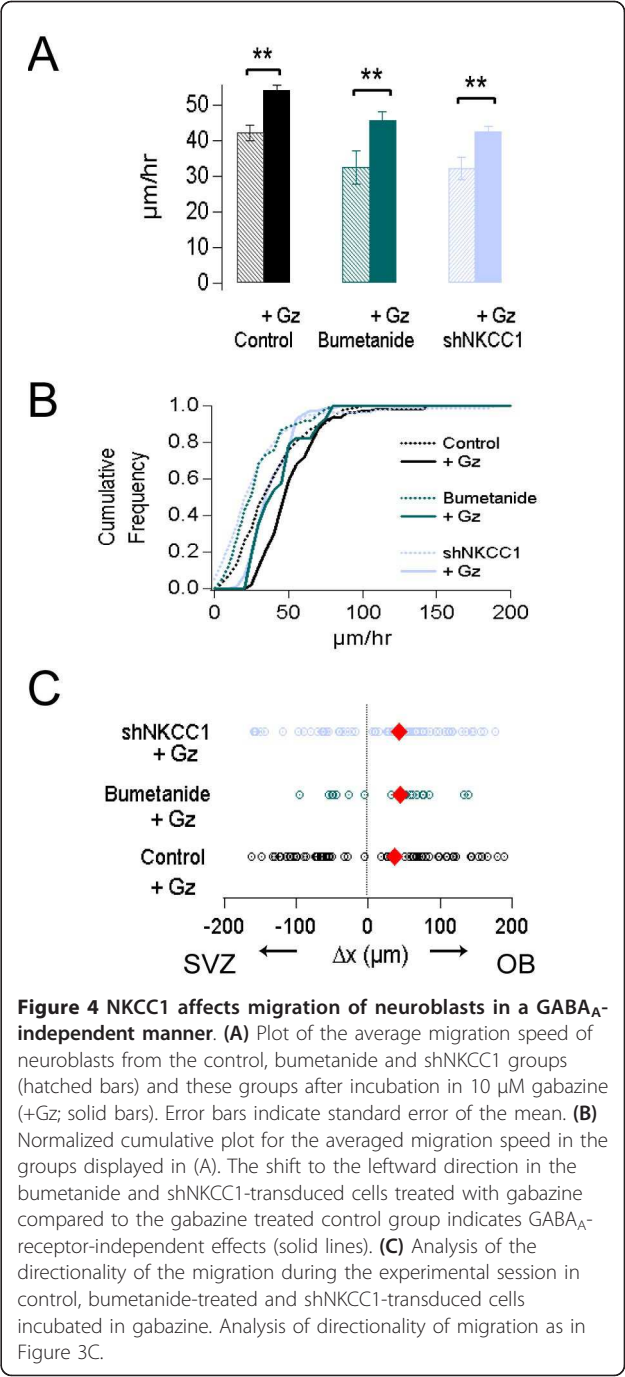
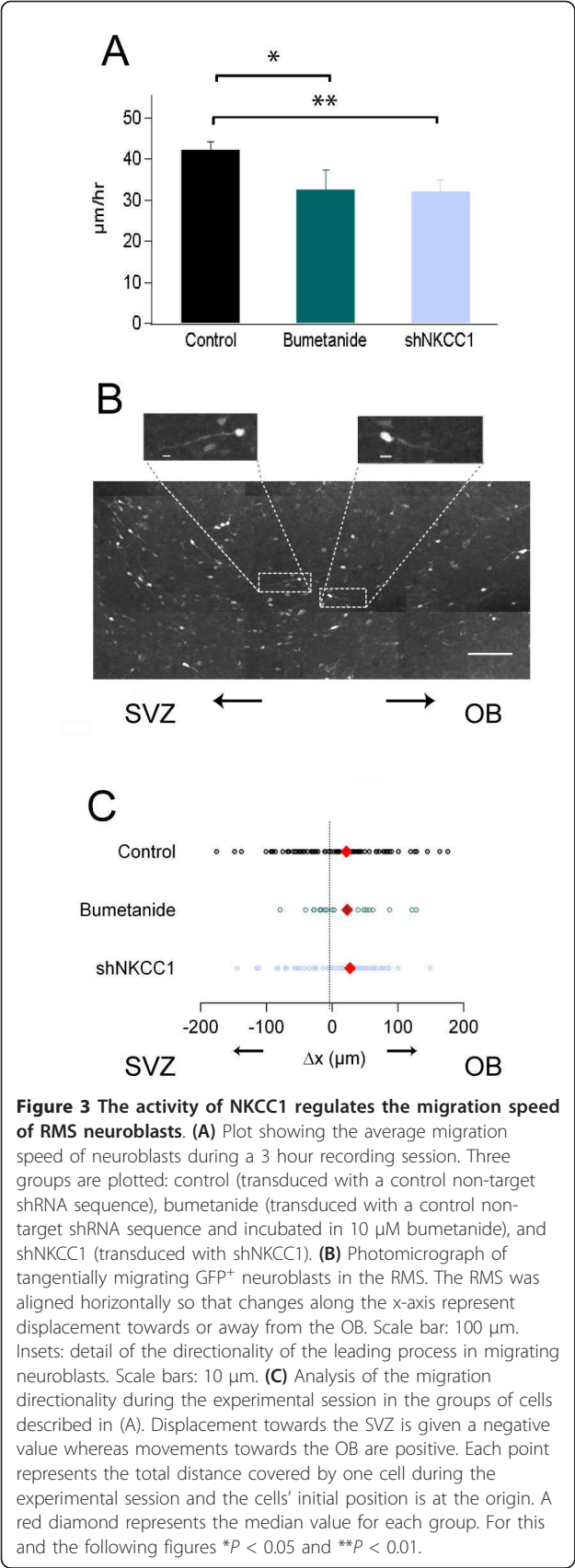
Figure 3B illustrates RMS neuroblasts migrating back and forth with respect to the OB. To investigate whether NKCC1 was required for determining the net tendency of the cells to migrate towards the OB, we compared the total displacement of cells along the x-axis, and used it as an index for directionality (Figure 3C). The initial cell position was set to zero and the values of displacement were considered negative when cells moved towards the injection site (that is, the SVZ) and positive when cells headed towards the bulb. We found no significant differences in the values of Δx between control cells and cells incubated in bumetanide or transduced with the shNKCC1 ($P = 0.9$ and 0.6 , respectively).

NKCC1 modulates the migration speed in a GABA_A-independent manner

GABA acting on GABA_A receptors reduces the migration of neuroblasts [12]. In order to test whether the

effects observed by reducing the activity of NKCC1 were due to changes in the action of GABA_A receptors, as a consequence of a potential drop in $[Cl^-]_i$, we analyzed the effects of blocking GABA_A receptors with 10 μ M gabazine in slices transduced with the control sequence, in slices transduced with the control sequence and treated with bumetanide, and in slices transduced with the shNKCC1 sequence.

We analyzed 69 cells transduced with the shNKCC1 sequence (from 4 slices), 43 cells treated with bumetanide (from 3 slices) and 112 cells transduced with the control sequence (from 5 slices); all were recorded in the presence of gabazine during the experiments. Our results were compared using a two-way ANOVA, which showed significant differences between the group injected with the control sequence, the group injected with control sequence and treated with bumetanide and the shNKCC1 sequence group ($P = 1 \times 10^{-6}$), as well as a significant effect of gabazine ($P = 1 \times 10^{-6}$). The differences between individual groups were then evaluated using a Tukey posthoc test. The comparisons show an increase in the average migration speed due to the GABA_A receptor blockade in the three groups (Figure 4A). The average speed of control cells treated with gabazine was significantly faster than in the non-treated control group (53.9 ± 1.5 compared to 42.2 ± 2 μ m/h; $P = 1 \times 10^{-3}$; Figure 4A). Gabazine also significantly accelerated the speed of migration when comparing bumetanide groups (from 34.4 ± 2.8 to 45.5 ± 2.5 μ m/h; $P = 0.01$) and shNKCC1-treated cells (from 31.9 ± 3.1 to 42.5 ± 1.6 μ m/h; $P = 0.01$). The cumulative distribution curve shifted to the right in all the groups treated



with gabazine, compared with their respective controls (Figure 4B). The Kolmogorov-Smirnov test revealed no significant difference in the distribution of the average speed between control groups and bumetanide or shNKCC1 groups treated with gabazine (*P* = 0.99 and 0.13). Overall, the analysis shows a very significant effect of NKCC1 pharmacological and genetic blockade as well as a significant effect of GABA_A-receptor blockers on the speed of migration of neuroblasts. Since the

interaction between these two variables was not statistically significant ($P = 0.78$), we conclude that the role of NKCC1 on the maintenance of migration speed is independent of the modulation exerted by GABA_A receptors.

To explore the effects of GABA_A signaling on the directionality of the migration, we analyzed the displacement of the neuroblasts along the x-axis, as explained in the previous section. We did not observe any effect on direction after the blockade of GABA_A receptors in any of the groups treated with gabazine - that is, control, bumetanide and shNKCC1 groups - compared to their respective non-treated groups ($P = 0.8$, 0.56 and 0.7 , respectively; Figure 4C). Altogether, our results show that neither NKCC1 nor the GABA_A-mediated activity are important in determining the directionality of RMS migrating neuroblasts.

Concerning the speed of migration, our experiments show that activation of GABA_A receptors diminishes the migration speed in a way compatible with previous observations [12]. Moreover, the experiments performed in the presence of gabazine showed that the effects of NKCC1 on neuroblast migration do not depend on GABA_A signaling.

A tonic GABAergic current is active in both migrating neuroblasts and maturing GCs

Our whole-cell recordings show that migrating neuroblasts do not receive spontaneous synaptic inputs, confirming previous studies [13]. The lack of synaptic inputs in neuroblasts holds even after treatment with alpha-latrotoxin to trigger massive neurotransmitter release (data not shown, $n = 3$ cells from 3 slices) [35,36]. In contrast, newly formed GCs displayed spontaneous synaptic activity, which started at 6 dpi and significantly increased at 10 dpi.

Due to the lack of synaptic activity in neuroblasts, the action of GABA on the migration speed must depend on a tonic influence of this neurotransmitter. This mechanism has been suggested [12] but the tonic currents have never been recorded. Thus, we recorded the spontaneous activity of GFP⁺ cells (5 to 11 dpi) held at -60 mV before and after applying the GABA_A receptor blocker gabazine ($10 \mu\text{M}$). GFP⁺ cells were either migrating neuroblasts located in the RMS (Figure 5A) or, for comparison, neurons located in the GCL (Figure 5B). In the GCL we paid particular attention to avoid recording from migrating GFP⁺ cells, most likely representing newborn PGCs *en route* to the glomerular layer. Our results confirm the tonic action of GABA_A receptors in neuroblasts and extend the observation to GCs. Moreover, they show that $10 \mu\text{M}$ gabazine, the same concentration used in our migration studies, efficiently blocks the tonic GABA_A-mediated currents, which became evident from a diminution in holding current of 4.0 ± 0.7 pA in migrating

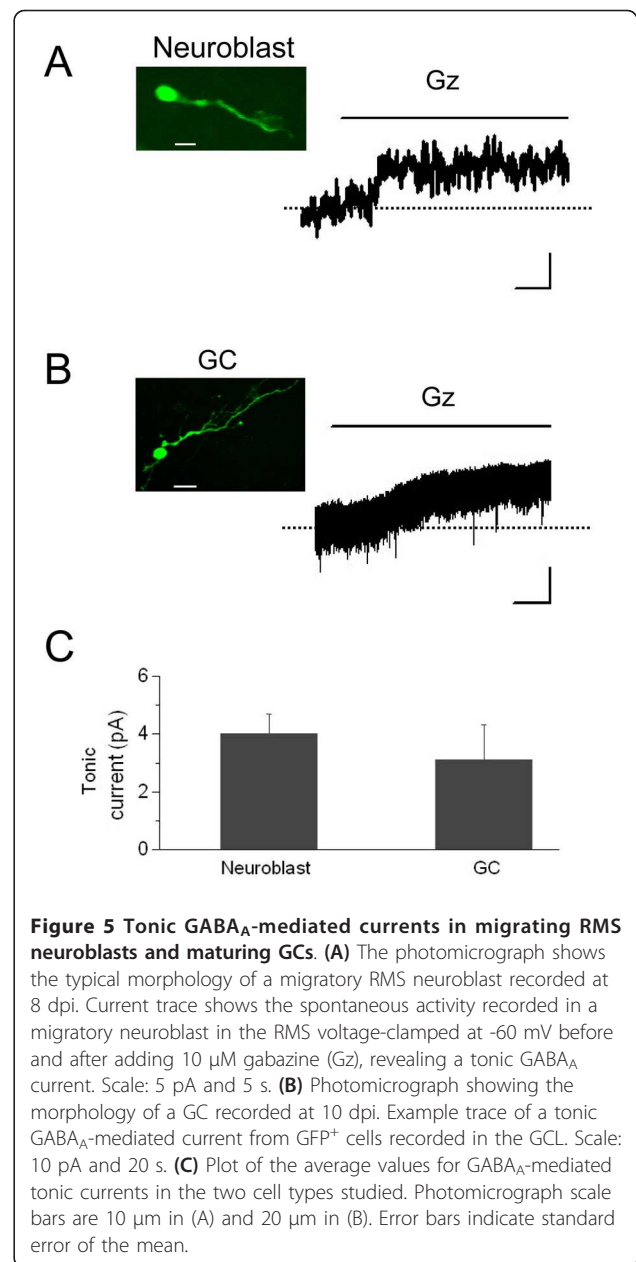


Figure 5 Tonic GABA_A-mediated currents in migrating RMS neuroblasts and maturing GCs. (A) The photomicrograph shows the typical morphology of a migratory RMS neuroblast recorded at 8 dpi. Current trace shows the spontaneous activity recorded in a migratory neuroblast in the RMS voltage-clamped at -60 mV before and after adding $10 \mu\text{M}$ gabazine (Gz), revealing a tonic GABA_A current. Scale: 5 pA and 5 s. (B) Photomicrograph showing the morphology of a GC recorded at 10 dpi. Example trace of a tonic GABA_A-mediated current from GFP⁺ cells recorded in the GCL. Scale: 10 pA and 20 s. (C) Plot of the average values for GABA_A-mediated tonic currents in the two cell types studied. Photomicrograph scale bars are $10 \mu\text{m}$ in (A) and $20 \mu\text{m}$ in (B). Error bars indicate standard error of the mean.

neuroblasts ($n = 10$ cells from 9 slices; $P = 9\text{e-}4$, paired t -test) and of 3.1 ± 1.2 pA in GCs ($n = 6$ cells from 6 slices; $P = 0.02$, paired t -test) (Figure 5C) [37].

NKCC1 controls E_{GABA} in RMS neuroblasts but not in GCs

In order to understand the effects observed on neuroblast migration, we decided to determine whether and, if so, how the activity of NKCC1 affects the action of GABA_A receptors and the overall excitability of the neuroblasts. For this we measured E_{GABA} (the reversal potential for GABA_A mediated responses) in RMS cells transduced with the control sequence and incubated in $10 \mu\text{M}$ bumetanide or in cells transduced with the

shNKCC1 sequence. As a reference, we repeated our measurements in newborn cells at a more mature stage, that is, in GFP⁺ GCs.

Gramicidin-perforated patch-clamp recordings were carried out to determine $[Cl^-]_i$ and the strength of GABA_A-mediated responses. A voltage-ramp from -80 to +20 mV was applied in the presence and absence of a focal application of 10 μ M muscimol (pressure ejected near the cell soma). The resulting currents were subtracted to calculate E_{GABA} (Figure 6Ai,Bi). Our results show that NKCC1 is an important modulator of GABA action in neuroblasts, since the diminution of its activity by either of the means tested produced a significant hyperpolarization of E_{GABA} . E_{GABA} in migrating RMS neuroblasts shifted from -29.8 ± 1.7 mV ($n = 6$ cells from 6 slices) in control to -49.7 ± 2.8 mV in bumetanide ($n = 9$ from 6 slices; $P = 0.01$) and to -56.7 ± 2.3 mV in shNKCC1-treated neuroblasts ($n = 8$ from 7 slices; $P = 4 \times 10^{-4}$) (Figure 6Bii).

In contrast, new GCs had E_{GABA} values significantly more hyperpolarized than neuroblasts in the control (-57.5 ± 4.1 mV; $n = 13$ from 13 slices; $P = 6.9 \times 10^{-5}$; Figure 6Aii) and this value was not sensitive to either bumetanide -58.6 ± 3.6 mV ($n = 12$ from 10 slices; $P = 0.99$) or shNKCC1 treatment (-52.8 ± 3.3 mV; $n = 6$ from 5 slices; $P = 0.94$) (Figure 6Bii).

We controlled for possible bias in our measurements due to the concentration of Cl^- in the patching solution. The comparison of E_{GABA} values between independent groups of cells using pipette solutions containing either a low or a high $[Cl^-]$ (solutions 1 and 2 in Materials and methods) did not show any difference ($P = 0.47$) so the results were pooled. Furthermore, an additional control using immunohistochemical labeling for KCC2 showed that the hyperpolarization of E_{GABA} in neuroblasts transduced with shNKCC1 is not due to a compensatory expression of KCC2, since no positive KCC2 labeling was observed in either GFP⁺ neuroblasts or GFP⁺ GCs (Additional file 7).

We used E_{GABA} values to estimate $[Cl^-]_i$ in both neuroblasts and GCs in the control and after treatments (Figure 6C). According to the effects on E_{GABA} , the calculated values of $[Cl^-]$ decreased from 43.5 ± 3.1 mM in control neuroblasts to 21.4 ± 2.3 mM ($P = 6.14 \times 10^{-5}$) in bumetanide- and 15.9 ± 1.3 mM ($P = 1.09 \times 10^{-5}$) in shNKCC1-treated cells. In newly formed GCs the values of $[Cl^-]$ in bumetanide- (16.4 ± 2.6 mM; $P = 0.99$) and shNKCC1-treated cells (17.2 ± 2.8 mM; $P = 1$) were not significantly different from those in the control (17.5 ± 2.7 mM). The lack of effect of the treatments diminishing NKCC1 activity on GCs agrees with the low expression of NKCC1 observed in the immunohistochemical analysis (Additional file 4B,C).

The values of E_{GABA} and $[Cl^-]$ were constant in each cell type for the duration of our study as no significant

correlation between E_{GABA} and dpi was observed in Spearman correlation tests ($r = -0.11$ for neuroblasts and $r = -0.22$ for GCs; Additional file 8).

NKCC1 controls the resting membrane potential in RMS neuroblasts

To understand the net effect of the activation of conductance on a cell's membrane potential, it is useful to determine the value of the potential at rest. The resting membrane potential (RMP) was measured in the cell-attached mode as the equivalent of the reversal potential of K⁺ conductances using isotonic intra- and extracellular concentrations of this ion [38]. We did verify that, in our system, this method was sensitive to predictable changes in RMP by measuring the reversal of K⁺ currents before and after raising the extracellular $[K^+]$ to 10 mM. This manipulation produced an average 3.4 ± 1.6 mV depolarization in the four cells tested.

We then measured the RMP in migrating neuroblasts and GCs in control slices. Remarkably, we did not find significant differences in RMP values between these cell groups: -71.5 ± 3.6 mV ($n = 6$ from 5 slices) and -71.0 ± 4.4 mV ($n = 6$ from 6 slices; $P = 0.93$, *t*-test), respectively. Nonetheless, the comparison of RMP in neuroblasts treated with bumetanide or shNKCC1 showed a depolarization due to the diminution of NKCC1 activity. The measured values were -59.8 ± 3.2 mV ($n = 6$ from 5 slices; $P = 0.07$) for bumetanide and -58.7 ± 2.9 mV ($n = 9$ from 5 slices; $P = 0.02$) in cells expressing shNKCC1 (Additional file 9). In contrast, in GCs the diminution of NKCC1 activity did not produce significant changes in RMP compared to control; values of -67.0 ± 1.0 mV ($n = 2$ from 2 slices; $P = 0.90$) in bumetanide- and -69.4 ± 6.1 mV ($n = 5$ from 5 slices; $P = 0.97$) in shNKCC1-treated cells.

We used the E_{GABA} values from muscimol-mediated currents and the RMP values to calculate the driving force for GABA-mediated responses. In neuroblasts, both the hyperpolarization of E_{GABA} as well as the depolarization of about 11 mV in their RMP, observed in bumetanide and shNKCC1 groups, resulted in an important decrease in the driving force for GABA responses, with the strongest effects on the shNKCC1-treated group. The calculated driving force value in control neuroblasts (41.7 ± 2.8 mV) was significantly diminished to 10.2 ± 2.9 mV ($P = 2.84 \times 10^{-5}$) in bumetanide- and to 2.0 ± 2.6 mV ($P = 1.03 \times 10^{-5}$) in shNKCC1-treated cells. Moreover, as expected from the lack of effect of the diminution of NKCC1 activity on both E_{GABA} and RMP in GCs, we did not observe any effect on the driving force between control GCs (13.5 ± 4.2 mV) and the bumetanide- (10.81 ± 3.6 mV; $P = 0.98$) or shNKCC1-treated groups (16.6 ± 4.8 mV; $P = 0.99$) (Figure 6Aiii).

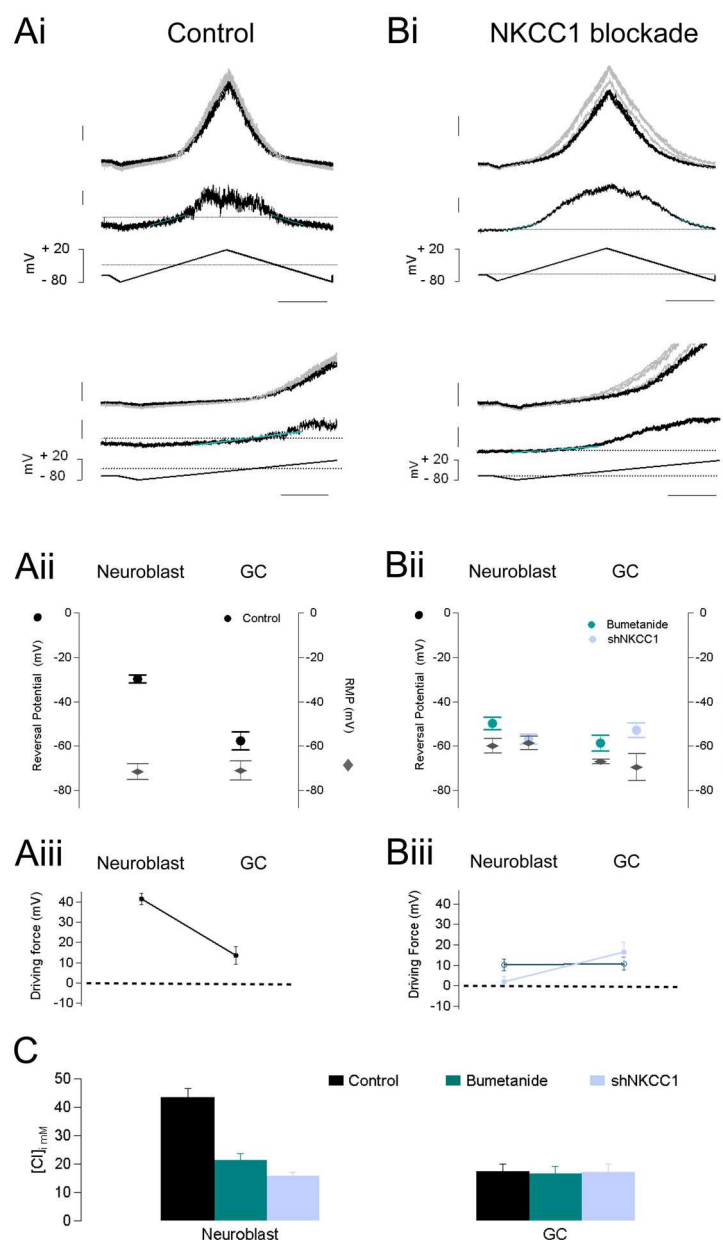


Figure 6 NKCC1 controls the reversal potential of GABA_A responses and resting membrane potential in migrating neuroblasts but not in GCs. (A) Reversal potential of muscimol-induced currents and resting membrane potential (RMP) in cells injected with control shRNA. (Ai) Upper panel: current traces showing the responses to a voltage-ramp protocol (bottom trace) applied to a RMS neuroblast recorded with the gramicidin-perforated patch technique. The ramp was induced in the absence (black) and presence (gray) of muscimol puffed near the soma of the recorded cell. The reversal potential was determined by fitting a straight line (light blue) to the subtracted current (middle trace). Scale top to bottom: 50 pA, 20 pA, 200 ms. Lower panel: high magnification version of the trace in the upper panel. Scale: 100 pA, 40 pA, 100 ms. (Aii) Average reversal potentials (left axis; dots) of the muscimol responses in migrating neuroblasts and GCs. The RMP (diamonds) is indicated on the right axis of the same plot. (Aiii) Driving force for muscimol responses in control neuroblasts and GCs. **(B)** Reversal potential to muscimol-induced current and RMP in cells with impaired NKCC1 function. (Bi) Current traces elicited by a voltage ramp applied to a migrating neuroblast transduced with shNKCC1 (as described in (A)). Scale: 100 pA, 50 pA, 200 ms. Lower panel: higher magnification of the upper panel trace. Scale: 100 pA, 60 pA, 100 ms. (Bii) Reversal potential (left axis) and RMP (right axis) measured in migrating neuroblasts and GCs after incubating slices with 10 μ M bumetanide (dark blue) and in cells transduced with shNKCC1 (light blue). (Biii) Driving force of GABA-induced currents, for each cell type, after bumetanide treatment (dark blue) or in shNKCC1-expressing cells (light blue). **(C)** Plot of calculated average \pm sem $[Cl^-]_i$ values in migrating neuroblasts and maturing GCs in control conditions (black), after incubation with bumetanide (dark blue), and in cells transduced with shNKCC1 (light blue).

A comparison between cell types - neuroblasts versus GCs - in the control condition shows a significant decrease in the driving force for GABA_A-mediated responses in newborn interneurons when reaching the bulb ($P = 5.28 \times 10^{-5}$; Figure 6Aiii,Biii). Nonetheless, GABA maintained a depolarizing action on both cell types. Our results indicate that NKCC1 expression is diminished in GCs from the time of their arrival at the OB. Therefore, in sharp contrast to neuroblasts, neither the action of GABA nor the values of RMP depend on NKCC1 activity in developing GCs.

Finally, we verified whether gabazine was able to change the RMP in neuroblasts, as would be expected from the measured driving forces for muscimol responses. In control cells, gabazine induced a 2.3 ± 0.8 mV ($n = 3$ cells) hyperpolarization of the RMP while only a 0.45 ± 0.04 mV hyperpolarizing shift was observed in shNKCC1 neuroblasts ($n = 2$). The lesser effect of gabazine on RMP in shNKCC1 cells supports the diminished driving force for muscimol observed in these cells. Moreover, the lesser effect on RMP observed in shNKCC1 cells might be related to the decreased effect of gabazine on the acceleration of neuroblast migration in the bumetanide and shNKCC1 cells (Figure 4).

Discussion

In the present study we characterize the role of NKCC1 on the regulation of the migration and excitability of migratory neuroblasts and developing OB interneurons in early postnatal tissue. We found that NKCC1 activity is important for the neuroblasts to maintain a normal migratory speed since two independent manipulations employed to impair NKCC1 function, pharmacological blockade with bumetanide and transduction with shNKCC1, produced a diminution of the migration speed (Figure 3A). Meanwhile, the manipulation of NKCC1 did not affect the directionality of migration (Figure 3C), suggesting that these two aspects are regulated by different mechanisms.

We performed electrophysiological measurements to determine the effects of NKCC1 on the excitability of migratory neuroblasts. Our results show that, similar to other developing neurons [19-21], the activity of NKCC1 maintains a high concentration of $[Cl^-]_i$ in RMS neuroblasts (Figure 6Bii). In addition, we demonstrate that the activity of this cotransporter is necessary to keep a hyperpolarized RMP in neuroblasts (Figure 6Bii), and thus to control the resting excitability in these cells in a manner independent of the regulation of Cl^- fluxes initiated by neurotransmission. This result supports previous studies showing modifications of RMP and neuronal excitability after manipulation of Cl^- cotransporters in different types of neurons [20,39-41]. It is likely that changes in the ionic concentration of Na^+ contributed

by the activity of Cl^- cotransporters alter the homeostasis and resting potential of the cells. Along these lines, manipulations depleting the internal concentration of this ion have been shown to depolarize the RMP of neonatal neurons [39]. Possible explanations relating the alterations in Na^+ fluxes and modulations of RMP could involve diminished electrogenic activity of ionic carriers such as Na^+/K^+ ATPase, thus producing a depolarization of the RMP [42], or changes in the magnitude of K^+ conductances, very prominent in the RMS neuroblasts [13]. The exact mechanisms explaining the significant depolarization of RMP observed in the RMS neuroblasts after the blockade of NKCC1 is beyond the scope of the present study. Nevertheless, the alteration of RMP after the blockade of Cl^- cotransporters observed for the first time in migratory neuroblasts in the present study and in other cell types in previous reports (see above) suggests a relevant role for NKCC1 in the control of resting excitability in developing neuronal cells.

Our results show that the blockade of GABA_A receptors accelerates the migration speed of the neuroblasts in the RMS. These observations are in agreement with previous reports showing that tonically released GABA acts on GABA_A receptors to diminish the velocity of migration in RMS neuroblasts [12,43,44], as well as in the cortex [45]. In our study, the speed of migration was accelerated by GABA_A pharmacological blockade in both neuroblasts transduced with the control sequence and those treated with bumetanide or transduced with shNKCC1. This is in agreement with the depolarizing driving force for GABA measured in all groups of cells (Figure 6Aiii,Biii), as well as with the hyperpolarization of RMP values observed after treating the cells with gabazine. The statistical analysis of the effects of blockers of GABA_A receptors and NKCC1 shows a lack of interaction between these two mechanisms. Thus, our results strongly support the notion that the motogenic action of NKCC1 does not depend on the activity of GABA_A receptors. It is very likely that the effects of NKCC1 blockade on neuroblast migration are accounted for by the sustained depolarization of the RMP observed in cells with impaired NKCC1 function. In support of this hypothesis, a similar study by Bolteus and Bordey [12] shows that depolarization induced by KCl treatment diminishes the speed of RMS neuroblast migration. In that study, the diminution of speed induced by depolarization was only partially reverted, but not abolished, by the pharmacological blockade of GABA_A receptors with 100 μ M bicuculline. Thus, our results, together with previous findings, show that the migration of neuroblasts is very sensitive to manipulations affecting the resting membrane potential, therefore suggesting that the relationship between the membrane potential and the speed of migration might be more important than previously thought.

The precise way in which external and internal processes regulate neuroblast migration in the RMS remains unclear. Modifying bulbar neuronal activity has little effect on tangential migration. For instance, anosmia [46], naris occlusion [47], or surgical disruption of the RMS [48] failed to alter neuroblast migration. In contrast, migration in the RMS is highly sensitive to brain hyperexcitability, associated with status epilepticus [49], or to brain injury, which accelerates the speed of migration [49,50]. Further studies are needed to understand how these external signals can change neuroblast excitability and adjust their rate of integration into the OB circuits or injured brain areas. Considering the important role of NKCC1 in the integration of external signaling and resting excitability, further investigations should determine whether pathological brain activity can impact the function of this Cl^- cotransporter, particularly since previous studies have shown that NKCC1 function is sensitive to activity-dependent modulation [51,52]. In this regard, a recent study showed that the pharmacological blockade of NKCC1 diminishes the proliferation of glial precursors induced by ischemia in the cortex [53].

NKCC1 is inactivated during the transition from migrating RMS neuroblasts to GCs

In the present study we used organotypic slice cultures to compare the role of NKCC1 on the excitability of neuroblasts and in maturing newborn neurons. The use of this model seems particularly appropriate for the present study since both the neuroblast migration and GC integration occur in the organotypic control, in a similar way to those reported from acute models [12,31]. First, the speed of migration in our system is remarkably similar to previous values measured in acute slices [12]. Moreover, the labeled precursors evolve in the organotypic culture, first as a contingent of neuroblasts migrating in the RMS to increasingly populate the OB. Once in the GCL, the cells change their morphology and show functional maturation (Figure 1; Additional file 3), including the development of synaptic activity. A similar system has been previously used to study the control of differentiation in dopaminergic cells [54].

Our results show that the early maturation of OB interneurons, at least when considering GCs, is paralleled by changes in both GABAergic signaling and NKCC1 function. After reaching the bulb, the neural precursors acquire a mature morphology and establish synaptic interactions with pre-existing neurons, some of which are GABAergic [55]. Our results show that GABA action evolves from a non-synaptic, temporally and spatially diffuse tonic influence in the RMS to include synaptic contacts on GCs during the first weeks of development. The tonic influence of GABA was

preserved in GCs during the entire duration of our study, regardless of the development of synaptic contacts. These results are in agreement with a previous report showing that, in early postnatal tissue, GABA produced a depolarizing influence, capable of shunting the firing activity on GCs [56]. Correspondingly, we detected a small tonic current and a slightly positive driving force for Cl^- in GCs (Figures 5 and 6A iii). Slightly positive driving forces for GABA are frequently observed in immature interneurons [57-59] and are believed to favor neuronal development [14].

The manipulations affecting the activity of NKCC1 showed that the strong depolarizing action of GABA in RMS neuroblasts largely depends on this Cl^- transport system. Meanwhile, in GCs, GABA exerted a much weaker depolarizing driving force that was not affected by the manipulations blocking the activity of NKCC1. These results are confirmed by the immunohistochemical data showing a much higher proportion of neuroblasts expressing NKCC1 compared to GCs in slices injected with the control sequence, and a significant drop of NKCC1⁺ neuroblasts but not GCs in cells transduced with shNKCC1 (Additional file 4). In accordance with the functional and anatomical evidence of low expression of NKCC1 on GCs, we failed to find any effect of the manipulation of this Cl^- transporter on the development of GC dendrite projections (data not shown). Thus, it seems unlikely that GC synaptic connectivity in the OB is shaped by NKCC1, in contrast to the dentate gyrus [21].

Wang and colleagues [56] showed that, after the second postnatal week, GABA hyperpolarizes the GC, held at -60 mV, at the time that KCC2 mRNA appears. We observed a strong hyperpolarization of E_{GABA} when GFP⁺ cells reached the GCL. However, GABA action did not become hyperpolarizing during our study, possibly owing to delays in this process due to *in vitro* conditions, or because our recorded cells were young and did not yet express KCC2 (Additional file 7B). Several reports have shown that KCC2 mRNA expression is abundant in mature neurons, whilst negligible expression was detected in neuronal progenitors [17,60]. It is conceivable that, at later stages of maturation, KCC2 is involved in the synaptic connectivity of GCs, as it is in other systems [40,61]. Nonetheless, our results indicate that the first significant decrease in $[\text{Cl}^-]_i$ in GCs, and therefore the simultaneous hyperpolarization of E_{GABA} , results from a functional down-regulation of NKCC1.

Altogether our data suggest that variations in NKCC1 activity can efficiently control various steps of OB postnatal neurogenesis. Accordingly, we show that an important change in the functional expression of this cotransporter accompanies the developmental transition from neuroblasts to developing GCs. NKCC1 is highly

active in the RMS precursor cells and its functional expression drops drastically in the newborn GCs, greatly affecting the action of GABAergic signaling on this early period of functional integration. Concerning the neuroblasts in the RMS, our results show, for the first time, an important role of NKCC1 in the control of neuroblast excitability. Our electrical determinations indicate that NKCC1 controls at least two different aspects of neuroblast activity. The first concerns the control of $[Cl^-]_i$ and therefore the integration of the most prominent neurotransmitter signaling system in the RMS, GABA_A-receptor-mediated activity [12,13,31,43,62,63]. The second aspect concerns the regulation of intrinsic excitability and depends on the control of the RMP. Thus, our study suggests that NKCC1 integrates relevant external signaling and the intrinsic neuronal excitability to finely tune the migration speed of RMS neuronal precursors *en route* to the OB.

Materials and methods

Culture preparation

Postnatal day 7 C57BL/6J mouse pups (Janvier, Le Genest Saint Isle, France) were decapitated and the brains removed and placed into ice-cold artificial cerebrospinal fluid (ACSF) containing (in mM) 124 NaCl, 3 KCl, 1.3 MgSO₄, 26 NaHCO₃, 1.25 NaH₂PO₄, 10 glucose and 2 CaCl₂, saturated with 95% O₂/5% CO₂. Sagittal forebrain slices, 300 μm thick, containing the migratory pathway from the SVZ-RMS and OB were prepared with a vibratome (Leica VT1200). All experimental procedures were performed in accordance with the Charter of Fundamental Rights of the European Union (2000/C 364/01), the European Communities Council Directive of 24 November 1986 (86/609/EEC), and European Union guidelines. They were reviewed and approved by our Institutional Animal Welfare Committee.

Each brain slice was transferred to a Millicell-CM Culture Plate Insert (Millipore PICM ORG 50; Billerica, MA, USA), the excess ACSF was removed and the insert was placed in a 35-mm Petri dish containing 1 ml of medium, as described by Stoppini and colleagues [64]. Culture medium consisted of 46% MEM (31095-029, Invitrogen, Paisley, UK), 25% HBSS (24020-091, Invitrogen), 25% horse serum, 20 nM HEPES, 6 mg/ml D-glucose and gentamycin (Invitrogen, Paisley, UK). The slices were incubated at 37°C in a humid atmosphere of 5% CO₂. The medium was changed three times per week.

Lentiviral vectors

We used lentiviral vectors created using the PTRIPDU3 method to express GFP in neural precursors and neuroblasts [65]. The first vector was constructed such that GFP expression was under the control of the PGK

promoter. For the second vector, a previously published shRNA sequence for NKCC1, shNKCC1-1 (5'ACA-CACTTGTCCTGGGATT3'), was introduced under the control of the U6 promoter [21]. This vector was similar to the first, also containing a sequence to induce GFP expression, but under the control of the CMV promoter. As a control, we constructed a third vector in which the shRNA sequence was replaced with a non-target scrambled sequence (5'GCCAGATTTCTCAGGTGATAA3'). Twenty-four hours after culturing the slices, 270 pg of p24 from one of the constructions was injected in the SVZ, in a volume of 18 nl.

Electrophysiology

GFP⁺ RMS neuroblasts and olfactory bulb GCs, at 3 to 11 dpi, were recorded using the patch-clamp technique in different configurations as detailed below. For data acquisition and electrical control an EPC-10 amplifier (Heka Elektronik, Lambrecht/Pfalz, Germany) was used. Before the recordings, a piece of Millicell membrane containing the slice was cut out and transferred to a chamber. During the recording session the slices were continuously perfused at a rate of 1.5 ml/minute with extracellular solution containing (in mM) 124 NaCl, 3 KCl, 2.4 CaCl₂, 1 MgCl₂, 25 NaHCO₃, 1.25 NaH₂PO₄, and 10 glucose. The extracellular solution was heated to 35°C at the entrance of the chamber (inline heater TC-324B, Warner Instruments, Inc; Hamden, CT, USA). Glass pipettes with a resistance of 6 to 7 MΩ were used for the recordings.

The reversal potential of GABA_A-receptor-mediated currents was determined using the perforated patch-clamp configuration with gramicidin - a cation-specific channel - in order to obtain voltage control of the cell while leaving $[Cl^-]_i$ intact [66]. For perforated-patch recordings one of the following patching solutions was used in each experiment: (1) 150 mM KCl, 10 mM Hepes; or (2) 150 mM KGlucuronate, 10 mM Hepes, both at pH 7.5. A stock of 1 mg/ml of gramicidin was freshly prepared on the day of the experiment and a final concentration of 2 μg/ml was added to the patching solution. The pipette tip was dipped in gramicidin-free patching solution before backfilling it with the gramicidin-containing solution. A giga-seal was established with the cell membrane and the recording session began when a stable access resistance between 50 and 70 MΩ was obtained.

To determine the reversal potential of the currents elicited by the GABA_A receptor agonist muscimol, a 1 s voltage-ramp from -80 to +20 mV and back to -80 mV was applied to cells recorded in the perforated configuration. Muscimol (10 μM) was pressure ejected (10 to 20 psi) near the cell membrane during the ramp. At least three ramps were applied and averaged before and during

muscimol application. The averaged currents were subtracted and the resulting trace was used to measure the reversal potential of the muscimol-triggered current. Reversal potentials were determined by linear fitting of the current trace using Igor (Wavemetrics, Inc; Lake Oswego, OR, USA), whereas $[Cl^-]_i$ was estimated from the Nernst equation. Repeated agonist application can influence the apparent Cl^- reversal potential (E_{GABA}), particularly in immature neurons in which Cl^- regulation mechanisms are less effective [67]. Therefore, we ensured that the interval between agonist applications was sufficiently long (20 s) to avoid shifts in E_{GABA} .

To determine the cells' RMP we used the method described by Verheugen *et al.* [38]. Briefly, we kept the cells in the cell-attached mode at -60 mV and applied a voltage-ramp from -100 to +200 mV to activate a voltage-dependent K^+ conductance. Patching solutions 1 or 2, described above, were used in the absence of gramicidin for these determinations. Both solutions have a concentration of 150 mM K^+ ; therefore, the reversal potential of the K^+ conductance was assumed to be equal to the RMP. The difference between average RMP values and the reversal potential of the muscimol-induced current values was used to determine the driving force for GABA in each condition and cell type.

Tonic currents were recorded in the whole cell mode at -60 mV holding potential. For these recordings solution 1 was supplemented with 2 mM ATPMg, 0.2 mM GTPNa and 0.2 mM EGTA. We recorded the spontaneous activity of the cells held at -60 mV for at least 2 minutes in normal ACSF before applying 10 μ M gabazine to the bath, and then recorded spontaneous activity for 5 minutes after the application of the drug. To measure the tonic $GABA_A$ current, we assessed the average holding current necessary to maintain the voltage at -60 mV, during 10 s in control ACSF, starting 40 s before application of gabazine, and 30 s after its application in the bath. Bicuculline blocks SK-type K^+ channels [68] and may affect the measurement of tonic current. Thus, gabazine is the drug of choice for studying tonic GABA conductance. For one group of cells, $GABA_A$ -dependent tonic currents were determined in the cells recorded first in the perforated-patch configuration and subsequently transformed into whole-cell mode by gentle suction within the recording pipette ($n = 6$ neuroblasts and 3 GCs). A second group was recorded in whole cell mode in the absence of gramicidin on the intracellular pipette ($n = 4$ neuroblasts and 3 GCs). No significant differences were observed between the values obtained with either method, neither for the RMS neuroblasts ($P = 0.57$) nor for the GCs ($P = 0.51$); thus, the results were pooled.

Only the cells maintaining a stable access resistance (less than 20% variation during the experiment) were analyzed. The series resistance values were between

10 and 25 M Ω and no compensation was applied. The spontaneous currents were filtered at 2.9 KHz with a low-pass Bessel filter and were sampled at 8.33 KHz. To analyze the passive properties of cells, the currents generated by a 20 ms, 20 mV hyperpolarizing pulse were filtered at 5 kHz and sampled at 10 kHz. All values were corrected for the measured liquid junction potential (-2 and -8 mV for solutions 1 and 2, respectively).

Drugs

All drugs were purchased from Sigma-Aldrich and used at the following concentrations: 10 μ M bumetanide was used to block NKCC1 activity. In all the experiments using bumetanide the slices were pre-incubated for 1 h in 10 μ M bumetanide and the same concentration was present during our electrophysiological and imaging recordings (Sigma-Aldrich, Saint-Quentin Fallavier, France). 10 μ M gabazine and 10 μ M muscimol were used as antagonist and agonist of $GABA_A$ receptors, respectively, and 10 nM alpha-latrotoxin was used to trigger neurotransmitter release.

Immunohistochemistry

Organotypic brain slices were fixed in 4% paraformaldehyde before being cryoprotected in 12.5% sucrose, 5% glycerol for 2 h at 4°C and incubated overnight in 25% sucrose, 10% glycerol. Slices were rapidly frozen and thawed three times over liquid N_2 vapor, washed in PBS before a second fixation in paraformaldehyde for 15 minutes, and incubated for 2 h in 10%, NGS 0.1% Triton blocking solution at room temperature. For immunolabeling of chloride cotransporters, slices previously injected with one of the lentiviral vectors were then incubated at 1 or 6 dpi in the primary antibody for NKCC1 (1:200; T4, Developmental Studies Hybridoma Bank) or at 6 dpi in the antibody for KCC2 (1:200; 07-432, Upstate, Billerica, MA, USA) mixed with the anti-GFP (1:1,000; 06-896, Upstate). A different set of organotypic slices at 6 div were incubated in the following primary antibodies: neuronal nuclei marker NeuN (1:200; MAB377, Chemicon, Billerica, MA, USA; $n = 3$), DCX (1:1,000; Ab18723, Abcam, Cambridge, UK; $n = 3$), GABA (1:1,000; A2052, Sigma; $n = 3$) or TH (1:4,000; 22941, ImmunoStar, Hudson, Wisconsin, USA; $n = 3$). All primary antibodies were diluted in blocking solution and incubated for 48 h at 4°C, followed by washes in PBS. Then, slices were incubated for 2 h in an alexa-conjugated secondary antibody (Alexa Fluor 568 (1:750) for the chloride transporters and Alexa Fluor 488 (1:800) for the rest of the proteins; both from Molecular Probes, Paisley, UK) at room temperature and washed in PBS. NeuN, DCX, GABA and TH stained cultures were then incubated in DAPI for 15 minutes at room temperature and washed in PBS. All the slices were then

mounted in ProLong Gold antifade (P36930, Molecular Probes). For analysis, images of 0.5 to 1 μm thick were acquired either in an upright microscope (Zeiss), equipped with a confocal spinning disk unit (Andor, Belfast, UK) or in an inverted microscope (Axioscope FS, Zeiss), equipped with a structured illumination device (Apotome, Zeiss).

To determine the number of GFP⁺ cells in the field that were also positive for NKCC1, we used the Intensity Correlation Analysis package of ImageJ [69]. The region analyzed was first defined by the limits of the GFP signal of a cell in the z-axis. Signal was considered a cell when the morphology corresponded to a round or elliptical cell body of 6 to 12 μm with at least one projection. We considered as double labeled those cells showing a positive intensity correlation quotient for both GFP and alexa 568 signals. The number of labeled cells per field (313 \times 313 μm) was determined in three independent images for each region of interest (RMS and GCL), and the results were normalized to the total number of GFP⁺ cells analyzed in each image and then averaged (Figure 2; Additional file 4).

Time-lapse videomicroscopy

The Petri dish containing the cultured slice was transferred to a closed chamber with the temperature control adjusted to maintain the medium at 36°C (DH40i, Warner Instruments, Inc.). During the entire time-lapse session, the tissue was oxygenated using a mixture of 95% O₂/5% CO₂. To prevent evaporation of the culture medium, the internal environment of the chamber was continuously humidified (Humidifying module warmer HMW-1, Warner Instruments, Inc.). Mosaic images of the entire RMS were re-constructed from individual frames (x-, y- and z-axis), acquired every 20 minutes during a period of 3 h. We used an inverted fluorescence microscope (Axioscope FS, Zeiss) equipped with a 20 \times long-distance objective (LD plan Neofluar, Zeiss). For analysis, a maximal projection of the different planes taken along the z-axis was obtained for each time-lapse. Individual RMS neuroblasts could be identified and their trajectories were tracked using Metamorph (Molecular Devices). Two migration variables were analyzed, the speed and displacement along the x-axis, corresponding to the tangential displacement in the RMS, provided that the rostro-caudal axis of the RMS was positioned parallel to the x-axis (Figure 3B,C). Delta x (Δx), the total displacement of a cell along the rostro-caudal axis of the RMS during an experimental session, was used as an index of the direction of migration in the RMS-OB pathway.

Statistical analysis

All results are expressed as averages \pm standard error of the mean. Experiments involving the comparison of a

control and an experimental group were analyzed using the Student's *t*-test. For multiple comparisons either one- or two-way ANOVA were used with the Tukey method as a post-hoc test; the use of each is detailed in the Results section. For multiple comparisons of the immunohistochemical data Kruskal-Wallis and Mann-Whitney methods were used; in these cases significance was evaluated based on the Bonferroni adjustment of *P*-values. Differences were considered significant for values of *P* < 0.05. Distributions were compared using the Kolmogorov-Smirnov test in the R program [70].

Additional material

Additional file 1: Figure S1. Immunohistochemical staining for doublecortin (DCX) and neuronal nuclei marker (NeuN) in 6-div organotypic cultures. **(A)** Low magnification image (10 \times) from a sagittal organotypic slice stained with DAPI (blue) depicting the cellular nuclei in the rostral migratory stream (RMS), as delimited by dashed lines. **(B)** DCX (green) and DAPI labeling in the RMS elbow area. Inset: approximate location of the area shown in (C). **(C)** High magnification detail of the RMS double stained for DCX and DAPI. **(D)** Low magnification image showing the distribution of NeuN-expressing cells (green) in the granule cell layer (GCL) and absence of positive cells in the RMS in a sagittal organotypic slice of the OB. Outlined by dashed lines, the borders of the OB and the accessory OB are depicted for reference. **(E)** High magnification detail of the GCL region and OB RMS showing NeuN-expressing cells in the former and the cellular nuclei stained with DAPI. **(F)** Higher magnification detail of the OB RMS and GCL double stained for DAPI and NeuN. Scale bars: 250 μm for (A,E), 200 μm for (B,F), 20 μm for (C), and 500 μm for (D).

Additional file 2: Figure S2. Immunohistochemical staining for GABA and tyrosine hydroxylase (TH) in 6-div organotypic cultures. **(A,B)** Photomicrograph showing GABAergic cells (green) and cellular nuclei co-stained with DAPI (blue) in the RMS (A), and in the GCL regions (B). **(C)** Photomicrograph showing the presence of TH-positive cells (green) surrounding a glomerulus in the OB, in an organotypic slice co-stained with DAPI. Inset: approximate location of the region magnified in (Ci). **(Ci)** High magnification photomicrograph showing a TH-positive cell co-stained with DAPI in an OB glomerulus. **(D)** Photomicrograph showing TH-expressing cells in the OB of an organotypic slice. Inset: approximate location of the region depicted in (Di). **(Di)** High magnification photomicrograph of a TH-expressing cell double labeled with DAPI in the OB. Scale bars: 50 μm for (A), 20 μm for (B), 100 μm in (C,D), 20 μm in (Ci,Di).

Additional file 3: Figure S3. Functional maturation of recorded GFP⁺ cells. **(A)** Scatter plot and linear regression analysis showing a lack of correlation between input membrane resistance (IR) and days post-injection (dpi) for neuroblasts migrating in the RMS. **(B)** A significant correlation was found in maturing GCs. Each point in the plot represents the values of one recorded GFP⁺ cell.

Additional file 4: Figure S4. Immunohistochemical staining for NKCC1 in control and shNKCC1 neuroblasts and GCs. **(A)** Upper panel: photomicrograph showing the immunolabeling for NKCC1 (red) and GFP⁺ (green) on migrating neuroblasts in the RMS transduced with the control non-target sequence. Lower panel: product of the differences from the mean (PDM) analysis for the double staining, as described for Figure 2. **(Ai)** Upper panel: photomicrograph showing the immunolabeling for NKCC1 (red) on GFP⁺ (green) neuroblasts located in the RMS and transduced with the shNKCC1 sequence. Lower panel: PDM analysis for the upper panel image. **(B)** Upper panel: photomicrograph showing the immunolabeling for NKCC1 (red) and GFP (green) on GCs transduced with the control non-target sequence. Lower panel: PDM analysis for the double labeling. **(Bi)** Upper panel: photomicrograph showing the immunolabeling for NKCC1 (red) in GFP⁺ (green)

neuroblasts located in the GCL and transduced with the shNKCC1 sequence. Lower panel: PDM analysis for the upper panel image. For (A, Ai,B,Bi) scale bars are 10 μ m. Photographs in (A,Ai) were taken from cultures fixed at 6 dpi and in (B,Bi) at 7 dpi. **(C)** Normalized average (\pm standard error of the mean) number of neuroblasts and GCs showing correlated NKCC1 and GFP signals in cells transduced with the control and shNKCC1 sequences.

Additional file 5: Video 1. Video showing the migration of neuroblasts transduced with the non-target control sequence in the RMS. The total duration of the session is 3 hours.

Additional file 6: Video 2. Video showing the migration of neuroblasts transduced with the shNKCC1 sequence in the RMS. The total duration of the session is 3 hours.

Additional file 7: Figure S5. Immunohistochemical staining for KCC2 in the organotypic slices 6 days after transduction with the shNKCC1- and GFP-encoding sequences. **(A)** Upper panel: photomicrograph of the RMS showing KCC2 immunostaining (red) in GFP⁺ migrating neuroblasts (green). Lower panel: product of the differences from the mean (PDM) analysis for the immunostaining in (A), as explained in Figure 2. **(B)** Upper panel: photomicrograph showing KCC2 immunostaining (red) of GFP⁺ maturing interneurons in the GCL. Lower panel: PDM analysis for the immunostaining in (B). Scale bars: 10 μ m in (A) and 20 μ m in (B).

Additional file 8: Figure S6. The reversal potential of GABA_A-mediated responses in neuroblasts and GCs did not evolve in the culture. **(A,B)** Scatter plot and linear regression analysis showing the lack of correlation between the reversal potential for GABA and days post-injection (dpi) in neuroblasts migrating in the RMS (A) and in GCs (B). Each point in the plot represents the values of one recorded GFP⁺ cell.

Additional file 9: Figure S7. Example traces showing the measurements of voltage-dependent K⁺ current reversal in neuroblasts recorded in the cell attached configuration. Upper panel: three independent traces (gray) and their average (black) recorded in response to the voltage ramp depicted in the lower panel. **(A,B)** The blue dashed lines represent the linear fit applied to the average current trace and the black dashed lines indicate the corresponding voltage value for the reversal of K⁺ voltage-dependent currents in a control cell (A) and in a shNKCC1 cell (B). Scale bars: 100 pA and 2 ms for (A) and 50 pA and 2 ms for (B).

Abbreviations

ACSF: artificial cerebrospinal fluid; DCX: doublecortin; div: days *in vitro*; dpi: days post-injection; GABA: gamma-aminobutyric acid; GC: granule cell; GCL: granule cell layer; GFP: green fluorescent protein; IR: input membrane resistance; KCC2: K⁺-Cl⁻; NKCC1: Na⁺-K⁺-2Cl⁻; OB: olfactory bulb; PBS: phosphate-buffered saline; PGC: periglomerular cell; RMP: resting membrane potential; RMS: rostral migratory stream; shRNA: short hairpin RNA; SVZ: subventricular zone; TH: tyrosine hydroxylase.

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Authors' contributions

The experimental work of this study was achieved at the laboratory for Perception and Memory, at the Pasteur Institute in Paris. S-MG and PM-L conceived and designed the experiments. K-M produced the viral vectors,

organotypic cultures, immunohistochemical staining and corrected the English in the manuscript. S-MG performed the electrophysiological and migration experiments as well as the data analysis and interpretation. S-MG and PM-L wrote the manuscript.

Competing interests

The authors declare that they have no competing interests.

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