Title: Naked mole-rat brain neurons have reduced ASIC-mediated currents and are resistant to acid-induced cell death

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Abstract

Background: Regulation of brain pH is a critical homeostatic process. Changes in brain pH modulate various ion channels and receptors and thus neuronal excitability. Tissue acidosis, resulting from hypoxia or hypercapnia, can activate acid-sensing ion channels (ASICs), a family of primarily Na^+ permeable ion channels, although ASIC1a homomeric channels are also permeable to Ca^{2+}. During brain acidosis, ASIC activation produces neuronal death through a process called acidotoxicity, which acts alongside glutamate-mediated excitotoxicity. Naked mole-rats (NMRs, *Heterocephalus glaber*) are long-lived, fossorial, eusocial rodents that display remarkable behavioral/cellular hypoxia and hypercapnia resistance. In the central nervous system, ASIC subunit expression is similar between mouse and NMR with the exception of much lower expression of ASIC4 throughout the NMR brain. However, ASIC function and neuronal sensitivity to sustained acidosis has not been examined in the NMR brain.

Results: Whole-cell patch-clamp electrophysiology of cultured NMR and mouse cortical and hippocampal neurons demonstrated that NMR neurons have smaller voltage-gated Na^+ channel currents and more hyperpolarized resting membrane potentials. We further demonstrate that acid-mediated currents in NMR neurons are of smaller magnitude than in mouse, and that all currents in both species are fully blocked by the ASIC antagonist benzamil. Using the ASIC1a blocker Psalmotoxin-1 (PcTx1), we find that ASIC1a plays a major role in ASIC-mediated currents in mouse neurons, whereas in NMR neurons, only a few responses were blocked by 10 nM PcTx1. However, when nmrASIC1a was heterologously expressed in Chinese hamster ovary cells, neither 10 nor 100 nM PcTx1 inhibited acid-evoked currents. We also show that nmrASIC1a channel is Ca^{2+} permeable. Finally, we demonstrate that NMR neurons show greater resistance to acid-induced cell death than mouse neurons.

Conclusions: NMR neurons are less excitable than mouse neurons and show significant cellular resistance to acidotoxicity, likely resulting from smaller ASIC-mediated currents, which may be neuroprotective against acid-induced neuronal death.

Key words

ASIC, acid-induced currents, acidotoxicity, naked mole-rat, Psalmotoxin-1, hippocampus

Abbreviations

ASIC, acid-sensing ion channel; CHO, Chinese Hamster Ovary; CNS, central nervous system; DRG, dorsal root ganglion, NaV, voltage-gated Na^+ channel; NMR, naked mole-rat; PcTx1, Psalmotoxin-1; TTX, tetrodotoxin

Background

Acid-sensing channels (ASICs) are ion channels of the ENaC/Deg superfamily and most subunits are activated by extracellular protons [1,2]. Six different ASIC subunits are encoded by 4 ASIC genes (ASIC1a, ASIC1b, ASIC2a, ASIC2b, ASIC3 and ASIC4), which assemble as homo- or heterotrimers [3]; neither ASIC2b nor ASIC4 form proton-sensitive homotrimers. ASICs are primarily permeable to Na^+, although the ASIC1a homomeric channels are also Ca^{2+} permeable [2]. In the central nervous system (CNS), neurons have been shown to primarily express mainly ASIC1a homomers and heteromers of ASIC1a/2a and ASIC1a/2b [4–8], where they have been demonstrated to have key roles in synaptic plasticity [9–11] and fear conditioning [12–14], as well as being major players in neuronal death resulting from brain ischemia [5,15–17], and neurodegenerative diseases [18–20].

Regulation of brain pH is a highly complex and important process [21]. Brain tissue acidosis can result either from an increase in tissue partial pressure of carbon dioxide (PCO_2)

during hypercapnia, or from the accumulation of the byproducts of anaerobic metabolism, such as lactate and protons, during hypoxia [22]. During periods of tissue acidosis, activation of ASICs by extracellular acidification is worsened by the release of allosteric modulators such as lactate [23], spermine [16] and arachidonic acid [24,25]. In addition to the activation of the Ca^{2+} permeable ASIC1a channel [15,16], a drop in pH also modulates the activity of numerous others ion channels, including voltage-gated ion channels [26–29] and glutamate receptors [30,31], therefore leading to disturbance in ion homeostasis, excitotoxicity and ultimately neuronal death [5,15,16].

Naked mole-rats (NMRs, *Heterocephalus glaber*) are subterranean rodents belonging to the Bathyergidae African mole-rat family found in East Africa [32]. Unusually for a mammal, NMRs are eusocial [33,34]. However, NMRs also display a range of remarkable physiological peculiarities, which is beginning to make a significant impact on biomedical research [35]. The unusual physiology of the NMR includes: extreme longevity with no increased risk of death with ageing [36], an apparent absence of age-related neurodegenerative disorders [37,38], resistance to cancer [39–41], insensitivity to certain noxious and irritant stimuli [42–45] and hypoxia/hypercapnia resistance resulting altered NMDA receptor function and an ability to utilize fructose as an energy source [46–49]. It is striking that NMRs are resistant to many pathological conditions known to involve ASICs. Recordings of ASIC-mediated currents in dorsal root ganglion (DRG) sensory neurons demonstrated an increased frequency and magnitude of ASIC responses in NMR neurons compared to mouse neurons [43], with APETx2, an inhibitor of ASIC3-containing ASICs, demonstrating a key role for ASIC3, even though nmrASIC3 itself is non-functional [50]. Although we have previously mapped out ASIC expression in different NMR brain regions and found similar expression between mouse and NMR with the exception of much lower ASIC4 levels throughout the NMR brain [51], no one has yet studied the function of ASICs in NMR brain neurons.

In this study, we investigated acid-induced currents in mouse and NMR neurons using whole-cell patch clamp recording of cultured neonatal hippocampal and cortical neurons. We find that NMR neurons have ASIC-mediated currents of significantly smaller peak current amplitude than those recorded from mouse neurons. Using the selective ASIC1a blocker PcTx1, ASIC1a homomers were observed to play a major role in ASIC-mediated currents in mouse neurons while in NMR neurons, only few responses were blocked. When nmrASIC1a was heterologously expressed in CHO cells, we demonstrated that it is Ca^{2+} permeable, but that PcTx1 fails to produce inhibition. Lastly, we demonstrated that NMR neurons are resistant to acid-induced cell death. Overall, these results suggest that the reduced sensitivity of NMR neurons to extracellular acidosis may be neuroprotective.

Methods

Animals

All experiments were conducted in accordance with the United Kingdom Animal (Scientific Procedures) Act 1986 Amendment Regulations 2012 under a Project License (70/7705) granted to E. St. J. S. by the Home Office; the University of Cambridge Animal Welfare Ethical Review Body also approved procedures. Breeding couples of 1 male and 2 female C57/bl6 mice were conventionally housed with nesting material and a red plastic shelter; the holding room was temperature-controlled (21 °C) and mice were on a normal 12-hour light/dark cycle with food and water available *ad libitum*. Naked mole-rats were bred in house and maintained in a custom-made caging system with conventional mouse/rat cages connected by different lengths of tunnel. Bedding and nesting material were provided along with running wheels and chew blocks. The room was warmed to 28 °C and humidified, with a heat cable to provide extra warmth running under 2–3 cages, and red lighting (08:00–16:00) was used.

Neuronal cultures

P0-P2 mice and P0-P5 naked mole-rats were used to prepare cortical and hippocampal neuronal cultures. Multiple pups (2-4) were used to prepare a single culture. Following decapitation, heads were immediately placed in dishes containing ice-cold Hank's Balanced Salt Solution (HBSS) solution (20 mM HEPES, 30 mM glucose in HBSS, Life Technologies). Brains were removed, transferred to a new dish and the two hippocampi and cortices were isolated. Tissues were subsequently incubated in an enzymatic digestion solution: 2 mg/mL papain (Worthington Biochemical Corporation) in Hibernate-Ca^{2+} solution (Brain Bits), activated by 0.5 mM Glutamax (Life Technologies) at 37 °C for 30 min in a 5 % CO_2 incubator. The digestion solution was then replaced by HBSS solution supplemented with DNAse I (250 Kunitz units/mL, Sigma Aldrich) and tissues were slowly triturated (5-7 times) using a P1000 pipette. Neuronal suspensions were filtered through a 100 µm nylon cell strainer (Corning) to remove non-dissociated pieces of tissues before centrifugation for 5 mins at 1100 rpm at room temperature. Supernatants were discarded, and the pellets carefully resuspended in HBSS solution. After further centrifugation for 5 min 1100 rpm at room temperature, pellets were resuspended in MEM/HS solution: 10 % heat-inactivated horse serum (Life Technologies), 2 mM glucose, 0.0025 % Glutamax, and 0.2 mg/mL primocin (InVivogen). Hippocampal and cortical neurons were plated on 35 mm plastic dishes (Fisher Scientific), previously coated with 100 mg/mL poly-L-lysine (Sigma-Aldrich), rinsed with water and dried, at a density of 300 000 cells/mL (2 mL/dish). After a 4-hour incubation in a 37 °C / 5 % CO_2 incubator, the MEM/HS solution was removed and the dishes were flooded with Neurobasal/B27 solution (1X B27 Supplement, 0.0025 % Glutamax, and 0.2 mg/mL primocin). Naked mole-rat neurons were kept at 33 °C in 5 % CO_2 incubator, whereas mouse neurons were kept at 37 °C in a 5 % CO_2 incubator; this is due to NMRs being cold-blooded and NMR cells do not withstand 37 °C for long periods of time [52]. Half of the medium was exchanged for fresh medium every 2-3 days until the cultures were used for experiments.

Chinese Hamster Ovary Cell Culture and transfection

Chinese hamster ovary (CHO) cells (Sigma-Aldrich) were maintained using standard procedures, as previously described [50,53]. Twenty-four hours before transfecting cells, 35-mm dishes (Fisher Scientific) were coated with 100 mg/mL poly-L-lysine (Sigma-Aldrich); cells from a 70–80 % confluent flask were trypsinized and resuspended in 5 mL CHO medium; and a volume was taken to seed cells at a 1:10 dilution, 2 mL/dish. For transfections, an enhanced green fluorescent protein (eGFP) expression vector was used to enable identification of transfected cells, and DNA was transfected using 0.5 µg nmrASIC1a (or nmrASIC1b) DNA and 0.01 µg eGFP DNA; the transfection reagent Lipofectamine LTX (Life Technologies) was used according to the manufacturer's protocol.

Electrophysiology

CHO cells. Whole-cell patch-clamp recordings from CHO cells were performed at room temperature 24-hours after transfection. For ion substitution experiments, the intracellular solution contained (in mM): 140 KCl, 2 $MgCl_2$, 5 EGTA, 10 HEPES in MilliQ water; pH was adjusted by adding KOH, and the osmolality was adjusted to 310–315 mOsm with sucrose. The extracellular solutions contained (in mM): 140 NaCl (or LiCl, or N-Methyl-D-glucamine-chloride, NMDGCl), 1.8 $CaCl_2$ (or 5 for Ca^{2+} substitution), 2 $MgCl_2$, 10 HEPES in MilliQ water; osmolality was adjusted to 300–310 mOsm with sucrose, and pH was adjusted to pH 7.4 or pH 6 by adding NaOH. For experiments with the ASIC1 antagonist Psalmotoxin-1 (PcTx1), the extracellular solution contained (in mM): 40 NaCl, 100 NMDG, 4 KCl, 2 $CaCl_2$, 1 $MgCl_2$, 10 HEPES, 4 glucose in MilliQ water; pH was adjusted by adding NaOH, and the osmolality was adjusted to 300–310 mOsm with sucrose. The intracellular solution contained (in mM): 110 KCl, 10 NaCl, 1 $MgCl_2$, 1 EGTA, 10 HEPES, 2 Na_2ATP, 0.5 Na_2GTP, adjusted to pH 7.3 with KOH and to 300–310 mOsm with sucrose. Patch pipettes were pulled (Model P-97, Flaming/Brown puller; Sutter Instruments) from borosilicate glass capillaries (Hilgenberg GmbH) and had a resistance of 3–6 MΩ. Data were acquired using an EPC10 amplifier and Patchmaster software (HEKA). Whole-cell currents were recorded at 20 kHz, pipette and membrane capacitance were compensated using Patchmaster

macros, and series resistance was compensated by >60 %. For ion substitution experiments, the following protocol was used: 5 seconds pH 7.4; 5 seconds pH 6; and 5 seconds pH 7.4, the holding potential ranging from -100 mV to 80 mV. To minimize tachyphylaxis when repetitively stimulating nmrASIC1a [43,54], cells were recorded once for a particular potential. The responses of 4 to 9 cells to pH 6 at each potential were averaged and used to plot the current/potential curves. Permeability ratios were calculated as described in [2]. For recordings examining the effect of PcTx1 (Smartox) on nmrASIC1a currents, the following protocol was used: 5 seconds pH 7.4; 1 second pH 6.5; 30 seconds 10 nM PcTx1; 1 second pH 6.5; 90 seconds wash with pH 7.4; 1 second pH 6.5; 30 seconds 100 nM PcTx1 and 1 second pH 6.5; 90 seconds wash with pH 7.4.

Neuronal cultures. Hippocampal and cortical neurons from mouse and NMR were used for whole-cell patch-clamp recordings at 9-12 days *in* vitro (DIV9-12). Recordings were performed at room temperature using the following solutions: extracellular (in mM): 140 NaCl, 4 KCl, 2 $CaCl_2$, 1 $MgCl_2$, 4 glucose, 10 HEPES, adjusted to pH 7.4 with NaOH and 300–310 mOsm with sucrose; intracellular (in mM): 110 KCl, 10 NaCl, 1 $MgCl_2$, 1 EGTA, 10 HEPES, 2 Na_2ATP, 0.5 Na_2GTP, adjusted to pH 7.3 with KOH and to 310–315 mOsm with sucrose. Acidic extracellular solutions were made using MES (pH 5.0). Patch pipettes were pulled (Model P-97, Flaming/Brown puller; Sutter Instruments,) from borosilicate glass capillaries (Hilgenberg GmbH) and had a resistance of 6-10 MΩ. Data were acquired using an EPC10 amplifier and Patchmaster software (HEKA). Whole-cell currents were recorded at 20 kHz, pipette and membrane capacitance were compensated using Patchmaster macros, and series resistance was compensated by >60 %. Cell capacitances and resting membrane potentials were measured just after cell opening in whole-cell configuration. To study macroscopic voltage-gated currents, a standard voltage-step protocol was used whereby cells were held at -120 mV for 200 msecs before stepping to the test potential (-80 mV - +65 mV in 5 mV increments) for 50 msecs, returning to the holding potential (-60 mV) for 200 msecs between sweeps. In some experiments, tetrodotoxin (300 nM, Alomone Labs) was

perfused for 30 seconds before repeating the voltage-step protocol. To measure neuronal acid-sensitivity, cells were exposed to the following protocol: 5 seconds of pH 7.4; 5 seconds of pH 5; and 5 seconds of pH 7.4. ASIC antagonists (100 μM Benzamil, Sigma, or 10 nM PcTx1) were perfused during 30 seconds before applying another 5 second pulse of pH 5. After 90 second wash time with pH 7.4 solution, a 5 second pulse of pH 5 was applied to check for reversal of any block observed. Current amplitude was measured in Fitmaster (HEKA) by taking the maximum peak response and subtracting the mean baseline amplitude in the preceding 50 msec (voltage-gated currents) or ~2.5 sec (ASIC currents); current amplitude was normalized for cell size by dividing by cell capacitance. Using Igor Pro, for each individual cell that underwent the voltage-step protocol, the following equation was fitted to the normalized inward currents:

$$i(x) = \Gamma \cdot x \cdot \frac{1 - e^{-\frac{x - Erev}{25mV}}}{1 - e^{-\frac{x}{25mV}}} \cdot \frac{1}{(1 - e^{-\frac{x - VHalf}{slope}})^3}$$

where Erev is the reversal potential; Vhalf the half-activating potential; Γ a constant and x the command potential. Similarly, a Boltzmann equation was fitted to the normalized outward currents:

$$\frac{I}{Imax} = \frac{1}{1 + e^{\frac{(VHalf - Vm)}{slope}}}$$

where Vm is the membrane voltage and VHalf the voltage at half-maximal activation. To determine the inactivation time of the ASIC-mediated currents, a single exponential was fitted. Data are expressed as mean ± standard error of the mean (SEM). Using Prism (GraphPad), paired t-tests were used to compare the effects of ASIC antagonists on proton-gated currents within both mouse and NMR neuron datasets; unpaired t-tests were used to compare parameters, such as neuronal resting membrane potential and capacitance and ASIC-mediated current amplitude, between mouse and NMR neuron datasets.

Acid-induced cell death assays

Mouse and NMR neurons were used to measure acid-induced cell death at DIV9-12. The pH 7.4 and pH 5 extracellular solutions used were the same as those described above for electrophysiology experiments. Neuronal cultures from both mouse and NMR were rinsed twice with 37 °C solution (pH 7.4 or pH 5) and then incubated with pH 7.4 or pH 5 solution for 2 hours at 37 °C. Cultures from both conditions were then rinsed with warm pH 7.4 solution and incubated during 30 min with pH 7.4 solution containing 1.5 mM propidium iodide (PI, Sigma Aldrich) to stain necrotic cells and Hoechst 33342 (dilution 1/2500, Sigma) to label all nuclei. Labelled cultures were imaged using an epifluorescence microscope (Olympus) equipped with a 20X objective (Olympus) and a QImaging camera. To determine the percentage of dead necrotic PI-positive cells, we used the software Fiji to count the total number of cells per field of view by counting nuclei on the Hoechst images, and subsequently counting the number of PI-positive nuclei on the PI images. One to three dishes per condition were used, and three different images per dish were taken. Data were collected from three different cultures and each culture was prepared from multiple animals. A one-way ANOVA test (Prism, GraphPad) corrected for multiple comparisons (Tukey test) was used to compare the percentage of cell death in each field of view at pH 7.4 and pH 5 in both species. Data are expressed as mean ± standard error of the mean (SEM).

Results

Basic electrophysiological properties and voltage-gated Na^+ channel activity differ between NMR and mouse neurons

Electrophysiological recordings from NMR neurons have been performed in both DRG sensory neurons [43,50] and CNS neurons [46,47,55]. However, neuronal activity from NMR CNS has only been recorded in brain slices in the form of field excitatory postsynaptic potentials and the basic electrophysiological properties of NMR neurons in hippocampal and cortical cultures have not yet been described.

We first compared the capacitance and resting membrane potential of NMR and mouse neurons from both cortical and hippocampal neuronal cultures (Fig. 1). The capacitance of NMR neurons was significantly smaller than in mouse neurons in both cortical and hippocampal cultures (cortex: 17.27 ± 1.02 pF versus 28.72 ± 2.32 pF for NMR (n = 30) and mouse (n = 24) neurons, respectively; hippocampus: 17.41 ± 1.03 pF versus 38.13 ± 3.27 pF for NMR (n = 18) and mouse (n = 26) neurons, respectively; unpaired two-sided t-tests, **** $p < 0.0001$, Fig. 1a). Resting membrane potentials were measured as soon as the whole-cell configuration was established and NMR neurons were significantly more hyperpolarized than mouse neurons, in both cortical and hippocampal cultures (cortex: -57.03 ± 2.64 mV versus -44.05 ± 2.84 mV for NMR (n = 30) and mouse (n = 21) neurons, respectively; hippocampus: -55.11 ± 4.79 mV versus -43.85 ± 2.59 mV for NMR (n = 18) and mouse (n = 26) neurons, respectively; unpaired two-sided t-tests; ** $p < 0.01$; * $p < 0.05$, Fig. 1b).

We then investigated macroscopic voltage-gated currents in NMR and mouse neurons, using a voltage-step whereby cells were held at -120 mV for 200 msecs before stepping to the test potential (-80 mV to +65 mV in 5 mV increments) for 50 msecs, and returning to the holding potential (-60 mV) for 200 msecs between sweeps (Fig. 1c). Both NMR and mouse neurons showed inward and outward currents (Fig. 1c-e). We applied 300 nM tetrodotoxin (TTX) for 30 seconds after a first standard protocol, to investigate the contribution of TTX-sensitive voltage-gated Na^+ channels (NaVs) to the macroscopic voltage-gated inward currents recorded in NMR neurons (Fig. 1c, right panel). In both cortical and hippocampal NMR neurons, the voltage-gated inward currents were fully blocked by 300 nM TTX (n = 8 and n = 2 for cortical and hippocampal neurons, respectively). The fact that no inward current remained after application of 300 nM TTX indicates that only NaVs were activated with our voltage-step protocol and that there was no measurable contribution of voltage-gated Ca^{2+} channels to the inward currents recorded. Moreover, these results indicate that cortical and hippocampal NMR neurons only express TTX-sensitive NaVs.

Strikingly, NMR neurons had a significantly smaller inward peak current density in both cortical and hippocampal cultures (cortex: 28.79 ± 3.83 pA/pF versus 202.40 ± 21.35 pA/pF for NMR (n = 19) and mouse (n = 17) neurons, respectively; hippocampus: 39.00 ± 6.39 pA/pF versus 196.70 ± 28.28 pA/pF for NMR (n = 13) and mouse (n = 16) neurons, respectively; unpaired two-sided t-tests, **** $p < 0.0001$, Fig. 1d.i-ii). Additionally, the voltage of half-activation (Vhalf) and the peak inward current amplitude potential (peak Vm) were more depolarized in NMR neurons compared to mouse neurons (cortex: Vhalf: -36.12 ± 2.80 mV versus -45.42 ± 1.99 mV and peak Vm: -12.43 ± 3.02 mV versus -31.27 ± 2.61 mV for NMR (n = 19) and mouse (n = 17) neurons, respectively; hippocampus: Vhalf: -37.10 ± 2.34 mV versus -45.71 ± 1.30 mV and peak Vm: -17.62 ± 4.00 mV versus -30.68 ± 3.09 mV for NMR (n = 13) and mouse (n = 16) neurons, respectively; unpaired two-sided t-tests, **** $p < 0.0001$, ** $p < 0.01$; * $p < 0.05$, Fig. 1d.iii-iv). These results suggest that NMR neurons may be less excitable compared to mouse neurons, with more hyperpolarized resting membrane potentials and smaller voltage-gated inward currents that are activated at more depolarized potentials, i.e. a greater depolarizing stimulus is required to activate NMR voltage-gated inward currents that produce much smaller currents.

By contrast, current-voltage curves for voltage-gated outward currents were similar between NMR and mouse neurons from both cortical and hippocampal neurons (Fig. 1e.i) and Vhalfs were not significantly different (cortex: 7.13 ± 1.91 mV versus 10.77 ± 2.18 mV for NMR (n = 24) and mouse (n = 17) neurons, respectively; hippocampus: 5.15 ± 2.50 mV versus 11.59 ± 2.26 mV for NMR (n = 14) and mouse (n = 16) neurons, respectively; unpaired two-sided t-tests, $p = 0.221$ and $p = 0.094$ for cortex and hippocampus respectively, Fig. 1e.ii).

Acid-induced currents are mediated by ASICs in both NMR and mouse neurons

The expression profile of ASIC subunits is similar in mouse and NMR brains, with the exception of lower levels of ASIC4 throughout the NMR brain [51], results suggesting that

functional ASIC-mediated currents should be present in NMR as others have shown in mouse [4,5,7,8,15].

A 5 second pulse of pH 5 was applied to NMR and mouse neurons from both hippocampal and cortical neurons and rapidly activating and inactivating acid-induced responses were recorded in every cell of both species (Fig. 2a). However, the peak current density of acid-mediated responses recorded in NMR neurons was significantly smaller than in mouse neurons, in both hippocampal and cortical neurons (cortex: 15.41 ± 1.82 pA/pF versus 85.66 ± 10.90 pA/pF for NMR (n = 31) and mouse (n = 24) neurons, respectively; hippocampus: 20.54 ± 2.86 pA/pF versus 100.90 ± 17.68 pA/pF for NMR (n = 22) and mouse (n = 26) neurons, respectively; unpaired two-sided t-tests, **** $p < 0.0001$, *** $p < 0.001$, Fig. 2b). By contrast, the inactivation time constant of the acid-mediated currents was similar between NMR and mouse neurons (cortex: 0.35 ± 0.012 s versus 0.44 ± 0.056 s for NMR (n = 27) and mouse (n = 18) neurons, respectively, unpaired two-sided t-test, $p=0.0516$; hippocampus: 0.47 ± 0.06 s versus 0.35 ± 0.04 s for NMR (n = 13) and mouse (n = 19) neurons, respectively; unpaired two-sided t-test, $p = 0.0861$, Fig. 2c).

The transient nature of the acid-mediated inward currents in both NMR and mouse neurons is characteristic of ASIC-mediated currents [3] and to confirm the involvement of ASICs we utilized the ASIC antagonist benzamil [56]. After a first application of pH 5 for 5 seconds, we applied 100 μM benzamil for 30 seconds before a second pH 5 pulse, then followed by a wash period of 90 seconds (Fig. 2d, 2g). In cortical and hippocampal mouse neurons, all the acid-induced currents were reversibly blocked by 100 μM benzamil (cortex: pH 5: 133.60 ± 19.01 pA/pF; benzamil: 16.77 ± 5.28 pA/pF; wash: 133.80 ± 19.88 pA/pF (n = 9); hippocampus: pH 5: 151.00 ± 33.64 pA/pF; benzamil: 7.84 ± 2.67 pA/pF; wash: 89.26 ± 20.48 pA/pF (n = 10); one-way paired ANOVA test, Tukey's multiple comparison test, *** $p < 0.001$, ** $p < 0.01$, Fig. 2e-f). This is in agreement with previous studies indicating that acid-evoked currents in mouse hippocampal and cortical neurons are mediated by ASICs [4,8,15].

Similarly, in cortical and hippocampal NMR neurons, acid-induced currents were reversibly inhibited by 100 μM benzamil (cortex: pH 5: 9.66 ± 1.09 pA/pF; benzamil: 2.78 ± 0.49 pA/pF; wash: 9.13 ± 0.89 pA/pF (n = 7); hippocampus: pH 5: 11.13 ± 1.88 pA/pF; benzamil: 3.50 ± 0.53 pA/pF; wash: 8.15 ± 1.30 pA/pF (n = 6); one-way paired ANOVA test, Tukey's multiple comparison test, *** $p < 0.001$, ** $p < 0.01$, Fig. 2h-i). This is the first demonstration of functional ASIC-mediated currents in CNS NMR neurons.

ASIC1a plays a greater role in mouse neuronal acid responses than in NMR neuronal responses

In CNS mouse neurons, ASIC1a plays a major role in ASIC-mediated currents as it has been reported that neurons express mainly ASIC1a homomers, ASIC1a/2a and ASIC1a/2b heteromers [4–8]. Hence, we investigated the contribution of ASIC1a to the ASIC-mediated currents in NMR neurons.

After a first application of pH 5 for 5 seconds, we applied 10 nM PcTx1, which inhibits homomeric ASIC1a [57] and ASIC1a/2a heteromeric channels [58], for 30 seconds before a second pH 5 pulse, followed by a wash period for 90 seconds and a third pH stimulation (Fig. 3). Currents were considered blocked by the drug if the peak current density was reduced by more than 50%. In mouse neurons, two populations of responses were observed (Fig. 3a): in some neurons, the ASIC-mediated current was blocked by 10 nM PcTx1 (Fig. 3a.i), but in others, no inhibition was observed (Fig. 3a.ii). In cortical mouse neurons, 5 neurons out of 11 (45%) were blocked by 10 nM PcTx1 (pH 5: 91.90 ± 17.97 pA/pF; PcTx1: 52.29 ± 17.09 pA/pF; wash: 71.01 ± 14.14 pA/pF (n = 11), one-way ANOVA test, Tukey's multiple comparison test, ** $p < 0.01$, Fig. 3b.i-ii), and more than half of the mouse hippocampal neurons (54%, 7 out of 13 neurons) were reversibly blocked by 10 nM PcTx1 (pH 5: 88.67 ± 18.69 pA/pF; PcTx1: 19.65 ± 6.05 pA/pF; wash: 49.39 ± 10.06 pA/pF (n = 13), one-way ANOVA test, Tukey's multiple comparison test, * $p < 0.05$) (Fig. 3c.i-ii). This confirms previous studies where ASIC1a homomers were shown to be involved in ASIC-mediated currents in mouse hippocampal neurons [4].

Two populations of responses were also observed in NMR hippocampal and cortical neurons, where some neurons were fully inhibited by 10 nM PcTx1 (Fig. 3d.i) and some were not (Fig. 3d.ii). However, compared to mouse neurons, the proportion of fully blocked neurons was notably less in NMR neurons (Fig. 3e-f). In cortical NMR neurons, only 1 out of 13 neurons (0.07%) was reversibly inhibited by 10 nM PcTx1 (Figure 3e.i-ii). Although the mean peak current density after application of 10 nM PcTx1 was reduced compared to the pH 5 pulse before the drug application (pH 5: 16.01 ± 2.80 pA/pF; PcTx1: 12.96 ± 2.22 pA/pF; wash: 13.74 ± 1.91 pA/pF; n = 13; one-way ANOVA test, Tukey's multiple comparison test, ** $p < 0.01$) no recovery in current density was observed after the wash period, but rather a further decrease was seen. The decrease observed with repetitive stimulation is likely due to tachyphylaxis, a known phenomenon for ASIC1a [43,54] and which may also contribute to the decrease in current observed between the initial pH 5.0 stimulation and effect of PcTx1 on the second pH 5.0 stimulation. Similarly, only 2 hippocampal NMR neurons were reversibly inhibited by 10 nM PcTx1 out of 12 (16%) (Fig. 3f.i-ii). Tachyphylaxis is also likely to occur in hippocampal NMR neurons, because only the responses before and after the drug application were significantly different from each other with no recovery being observed after the wash (pH 5: 16.01 ± 2.80 pA/pF; PcTx1: 12.96 ± 2.22 pA/pF; wash: 13.74 ± 1.91 pA/pF; n = 13; one-way ANOVA test, Tukey's multiple comparison test, ** $p < 0.01$). These results are somewhat contradictory, the known ASIC1a antagonist PcTx1 has relatively little impact on most currents recorded in NMR neurons and yet tachyphylaxis, a key characteristic of nmrASIC1a homomers [43], appears to occur in most neurons. To date however, the effect of PcTx1 on nmrASIC1a has not been reported and so it is possible that the lack of effect observed with PcTx1 is due to its comparative inability to inhibit nmrASIC1a. However, 2 NMR hippocampal neurons are fully, reversibly blocked by 10 nM PcTx1 suggesting that PcTx1 may inhibit better different ASIC combinations in NMR neurons.

PcTx1 does not inhibit homomeric nmrASIC1a

According to the published amino acid sequences, the binding site for PcTx1 on ASIC1a is conserved between mouse and NMR [59,60] (Uniprot numbers nmrASIC1a: G9DXC2; mouse ASIC1a: Q6NXK8), which suggests that PcTx1 should inhibit nmrASIC1a. The confirmation of published literature that PcTx1 reversibly inhibits a significant proportion of acid-mediated currents in mouse hippocampus and cortical neurons (Fig. 3b-c) demonstrates that the protocol used to measure the effect of PcTx1 is appropriate, but when expressed in Chinese Hamster Ovary (CHO) cells we observed no inhibition of nmrASIC1a by PcTx1. We used a 1 second pulse of pH 6.5 to minimize tachyphylaxis, which is both pH and stimulation-time dependent [43], followed by application of 10 nM PcTx1 for 30 seconds, but observed no reversible inhibition of nmrASIC1a following a 90 second wash (Fig. 4a-b). Even at 100 nM PcTx1 failed to inhibit ASIC1a (Fig. 4a-b) (pH 6.5: 429.80 ± 139.10 pA/pF; 10 nM PcTx1: 407.10 ± 109.20 pA/pF; wash: 401.00 ± 116.70 pA/pF; 100 nM PcTx1: 569.10 ± 324.60 pA/pF; wash: 176.8 ± 55.48 pA/pF; n = 7; one-way paired ANOVA test, Tukey's multiple comparison test, p = 0.2905). Therefore, as we were not able to demonstrate any inhibition of nmrASIC1a by PcTx1 in a heterologous expression system, it is difficult to make any firm conclusions about the relative lack of PcTx1 activity observed on acid-mediated currents in NMR neurons and thus the contribution of ASIC1a to the responses observed (Fig. 3).

nmrASIC1a is Ca^{2+} permeable whereas nmrASIC1b is not

Although we are unable to make any firm conclusions on the presence of functional nmrASIC1a homomers in NMR CNS neurons, we have previously shown that ASIC1a is expressed throughout the NMR brain as in mouse [51]. In addition, we have shown that nmrASIC1a displays similar proton-sensitivity with regard to activation and steady-state inactivation as mASIC1a [43]. What makes ASIC1a a key player in causing acidotoxic neuronal death during ischaemia is its Ca^{2+}-permeability [15], which is unique amongst ion channels formed by different ASIC subunit configurations [1,2]. To test if nmrASIC1a is also Ca^{2+}-permeable, we

expressed nmrASIC1a or nmrASIC1b in CHO cells and recorded responses to 5 second pulses at pH 6 while holding the cells at different voltages ranging from -100 mV to 80 mV. Extracellular solutions contained either Na^+, Li^+ or Ca^{2+} as the main permeant cations (Fig. 5). At more hyperpolarized potentials, Ca^{2+} mediated currents through nmrASIC1a were recorded (Fig. 5a-b) (Ca^{2+} reversal potential: -73.94 mV; n = 5-6 cells for each voltage). Additionally, and as expected, nmrASIC1a was highly permeable to Na^+ and Li^+ [2] (pNa^+/pLi^+ = 1.87, pNa^+/pK^+ = 6.40, pNa^+/pCa^{2+} = 16.21; Na^+ reversal potential: 46.90 mV; n = 4-6 cells for each voltage; Li^+ reversal potential: 31.04 mV; n = 5-6 cells for each voltage) (Fig. 5a-b). By contrast, nmrASIC1b was not permeable to Ca^{2+}, but permeable to other cations (Fig. 5c-d) (pNa^+/pLi^+ = 0.62, pNa^+/pK^+ = 6.76; Na^+ reversal potential: 48.27 mV; n = 6-9 cells for each voltage; Li^+ reversal potential: 60.13 mV; n = 5-7 cells for each voltage). Thus, this confirms that nmrASIC1a is Ca^{2+} permeable and once activated could contribute to acidotoxicity, whereas nmrASIC1b is only permeable to other cations such as Na^+ or Li^+.

NMR neurons are resistant to acid-induced cell death

ASICs are involved in acid-induced cell death, so-called acidotoxicity, which can occur during periods of ischemia [5,15,16]. In particular, ASIC1a-containing channels play a pivotal role in Ca^{2+}-mediated cell death, because protons and lactate produced during acidosis can synergistically activate these Ca^{2+}-permeable channels leading to Ca^{2+} entry and consequently cell death [15,16]. nmrASIC1a is Ca^{2+} permeable, but because NMR neurons exhibit significantly smaller ASIC currents (Fig. 2), we hypothesized that this could lead to a less toxic entry of Ca^{2+} in NMR neurons and therefore be neuroprotective when neurons are in an acidic environment. We exposed neuronal cultures from mouse and NMR cortices to a pH 7.4 or a pH 5 solution for 2 hours at 37 °C. Nuclei were stained by Hoechst 33342 and necrotic (dead) neurons were labeled using propidium iodide (PI) (Fig. 6). Percentages of dead neurons were calculated by counting the number of necrotic PI-positive cells over the total number of Hoechst 33342-positive neurons

in the field of view, and a one-way ANOVA test corrected for multiple comparisons was used to compare neuronal death at pH 5 between both species (Fig. 6c). At pH 7.4, percentages of dead cells were comparable between mouse and NMR cortical cultures (mouse - pH 7.4: 20.31 ± 3.50 %, n = 18 fields of view; NMR – pH 7.4: 13.31 ± 1.53 %, n = 27 fields of view; from 3 independent experiments), suggesting no differences in neuronal death under basal conditions between NMR and mouse cultures. When incubated for 2 hours in pH 5 solution, mouse cortical neurons exhibited a significantly increased percentage of cell death compared to all other conditions (mouse – pH 5: 60.55 ± 4.87 %, n = 18 fields of view; NMR – pH 5: 22.29 ± 2.30 %, n = 28 fields of view, from 3 independent experiments, $p < 0.0001$, ****). By contrast, when NMR cortical neurons were exposed to pH 5 for 2 hours, no difference in the level of cell death was observed compared to incubation at pH 7.4. Similar results were obtained using mouse and NMR hippocampal neurons (data not shown). This indicates that NMR neurons are resistant to acid-induced cell death, possibly due to their reduced ASIC currents compared to mouse neurons.

Discussion

In this study, we recorded for the first time from cultured NMR brain neurons and described their basic electrophysiological properties (Fig. 1). NMR neuronal capacitance was smaller than in mouse neurons, suggesting that NMR neurons have a smaller dendritic and/or axonal arborization. This may be specific to culturing NMR neurons as in NMR brains, neurons appear to successfully grow neurites and present a large arborization [55,61]. Additionally, we showed that the resting membrane potential of NMR neurons is more hyperpolarized than in mouse neurons. However, previous data from others found the resting membrane potential of NMR hippocampal neurons to be further hyperpolarized (NMR cortical neurons: -57.03 ± 2.64 mV, NMR hippocampal neurons: -55.11 ± 4.79 mV, Fig. 1; NMR hippocampal pyramidal neurons (4 months old): -70.3 ± 6.1 mV, NMR hippocampal dentate granule cells (4 months old): -75.1 ±

3.6 mV, [55]). One explanation for the difference observed is likely the experimental conditions, here we cultured neurons, whereas Penz and colleagues made recordings in brain slices. Secondly, we isolated neurons from neonatal animals, whereas slice recordings were made from animals aged at least 4-months.

A standard voltage-step protocol that has been already successfully used in NMR sensory neurons to predominantly isolate NaV activity [43] was used to measure macroscopic voltage-gated current activity in mouse and NMR neurons. Inward currents recorded in NMR brain neurons were similar to currents recorded in NMR sensory neurons (NMR cortex: Vhalf: -36.12 ± 2.80 mV; NMR hippocampus: Vhalf: -37.10 ± 2.34 mV; NMR DRG neurons: Vhalf: from -34.2 ± 0.1 mV to -44.6 ± 0.5 mV [43]). However, NMR inward currents significantly differed from mouse currents: the peak current density was significantly smaller and the Vhalf and peak Vm values were more depolarized, suggesting that NMR NaVs are less excitable than in mouse neurons and that they convey less currents when activated. These differences in NMR NaV activity likely result from differences in amino acid sequence of NaV subunits and/or differential expression of accessory subunits and warrant further investigation. We also found that addition of 300 nM TTX completely abolished all voltage-gated inward currents demonstrating an absence of TTX resistant NaV subunits in the NMR brain. It should also be noted that we have made recordings under normoxic, standard laboratory conditions, whereas NMRs live in a hypoxic and hypercapnic environment [62,63], which may influence channel activity *in vivo*.

ASIC activity in brain neurons is now accepted as a key factor in numerous physiological and pathological conditions [64]. However, nothing is known about acid-induced responses in NMR brain neurons, which is of considerable interest considering the behavioral hypoxia/hypercapnia resistance and lack of acid-induced nocifensive behavior display by NMR that are likely adaptations to adapting to a safe, but relatively hypoxic and hypercapnic habitat [35]. Recently, we described ASIC subunit expression in the NMR CNS [65], which is similar to that in the mouse CNS, with the exception of lowered ASIC4 levels in the NMR brain. However,

evidence for functional ASIC activity is lacking and yet is of particular interest in light of our recent finding that nmrASIC3 forms non-functional homomers [50]. Here, we find that NMR brain neurons ASIC-mediated currents, in both hippocampus and cortex, as demonstrated by acid-induced responses being fully, reversibly blocked by 100 μM benzamil (Fig. 2). However, the peak current density of NMR ASIC-mediated responses was significantly reduced compared to responses recorded in mouse neurons. The reasons for such reduction in ASIC currents in NMR neurons is not known and additional research is needed to determine what underpins this different, e.g. are regulators of ASIC plasma membrane trafficking different in NMR?

With regard to the ASIC currents themselves, they have similar decay kinetics in both NMR and mouse neurons (Fig. 2C), which suggests that a similar mixture of ASIC subunits are expressed, as our previous mRNA based analysis suggested [65]. Previous studies have shown ASIC1a to be a key component of ASIC-mediated responses in rodent hippocampal and cortical neurons, as demonstrated by their sensitivity to the ASIC1a homomer and ASIC1a/2a heteromer inhibitor PcTx1 [4,5,58]. In agreement with these studies, we observed that ~50% of mouse hippocampal and cortical neurons were inhibited by 10 nM PcTx1 (Fig. 3). However, in NMR neurons PcTx1 did not prove so effective, with only a few neurons (< 20%) being inhibited PcTx1 (Fig. 3), a surprising result in view of the known ASIC1a/ASIC2a expression in NMR brain [65] and similarity in current kinetics between mouse and NMR neurons. However, when tested in a heterologous system, we observed that neither 10 nM nor 100 nM PcTx1 produced any significant, reversible inhibition of nmrASIC1a. Considering that mASIC1a and nmrASIC1a have 98% sequence homology and the amino acid sequence of the PcTx1 binding site is very similar between these channels [58–60], this is a surprising result and makes it difficult to interpret the functional contribution of ASIC1a to acid-evoked currents in NMR neurons. It is possible that PcTx1 inhibition of nmrASIC1a homomers has differential pH-dependence [66] and additional research should be carried to understand the structural interaction between nmrASIC1a and

PcTx1. Unlike the differential PcTx1 sensitivity observed, we confirm here that as in other species [2], nmrASIC1a is Ca^{2+} permeable (Fig. 5) and thus has the potential to contribute to acidotoxicity.

Due to nmrASIC1a, like mASIC1a, displaying Ca^{2+} permeability (Fig. 5) and a similar proton sensitivity [43], it is possible that nmrASIC1a is involved in acidotoxicity as has been described for mASIC1a in mouse neurons [15,16]. However, incubation with a pH 5.0 solution showed that unlike mouse neurons, NMR neurons do not display any significant acid-induced cell death (Fig. 6). This is the first demonstration of resistance to acid-induced neuronal death in NMR neurons. In rodent neurons, some factors shown to be protective against acid-induced neuronal injury, including lower temperature [67], pharmacological blockade or genetic deletion of ASIC activity [15] or ASIC trafficking [68]. Considering the similar prevalence of ASIC currents and the Ca^{2+}-permeability of nmrASIC1a, we suggest that the reduced ASIC-mediated current amplitude observed in NMR neurons may be an additional neuroprotective mechanism in NMR brains, alongside the previously described increased hypoxia-inducible transcription factor (HIF1-α) expression [69] or more efficient *in vivo* CO_2 buffering [47], which cannot account for these *in vitro* results. One possible mechanism for the decreased amplitude observed is reduced ASIC plasma membrane trafficking which is known to be modulated by extracellular acidic environment [68]. However, it is also possible that the reduced acid-induced cell death observed is not ASIC-dependent. Acid-induced cell death is a complex mechanism and dependent on several molecular players. ASIC activation appears to play a major role in neuronal injury [64], but many other ion channels are also modulated by a drop in extracellular pH, such as NaVs [27,28] and glutamate receptors [31], and here we show that NMR neurons also have smaller NaV-mediated currents, which may also add a layer of neuroprotection.

Conclusions

In this work, we describe for the first time the basic electrophysiological properties of NMR neurons in culture and showed that the resting membrane potential of NMR neurons is more hyperpolarized than that of mouse neurons, suggesting that NMR neurons are less excitable than mouse neurons. We then demonstrated that acid-induced currents in NMR neurons are, as in mouse, ASIC-mediated, but the role of ASIC1a remains unclear due to differences in ASIC1a pharmacology across species. The key result is that acid-induced cell death is virtually absent in NMR neurons, with reduced ASIC amplitude likely contributing to this observation, and thus this is a further adaptation enabling NMR to live in a subterranean, hypercapnic/hypoxic environment.

Declarations

Ethics: All experiments were conducted in accordance with the United Kingdom Animal (Scientific Procedures) Act 1986 Amendment Regulations 2012 under a Project License (70/7705) granted to E. St. J. S. by the Home Office; the University of Cambridge Animal Welfare Ethical Review Body also approved procedures.

Availability of data and material: The datasets supporting the conclusions of this article are available in the University of Cambridge Apollo repository: https://doi.org/10.17863/CAM.18704

Competing interests: The authors have no competing interests.

Funding: This work was supported by an Isaac Newton Trust Research Grant from the University of Cambridge. ZH was funded by a EMBO Long-Term Fellowship (ALTF1565-2015).

Authors' contributions: ZH and ESS designed the study. ZH performed all the experiments and analysed the data. ZH and ESS wrote the manuscript.

Acknowledgements: Thanks to all members of ESS lab for their useful discussions, and in particular to the assistant staff in the Department of Pharmacology and Animal Facility for their technical help and animal husbandry.

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Figure Legends

Fig. 1. Electrophysiological properties of NMR neurons differ from mouse neurons. NMR cortical and hippocampal neurons have significantly smaller capacitance (**a**) and more hyperpolarized resting membrane potential (**b**) than mouse neurons. **c.** Example of whole-cell patch-clamp recordings made from a mouse (left, black) and a NMR (middle, blue) hippocampal neuron in response to the voltage step protocol and the inhibition of response in NMR neurons by 300 nM TTX (right, blue). Cortical and hippocampal NMR neurons have significantly smaller voltage-gated inward currents (**a, b**), a more depolarized Vhalf (**c**) and a more depolarized peak activation membrane potential Vm (**d**) compared to mouse neurons. **e.** Voltage-gated outward currents in NMR cortical and hippocampal show no significant difference to those in mouse neurons. * $p <$

0.5; ** $p < 0.01$; **** $p < 0.0001$, unpaired t-tests within each structure. Numbers in brackets indicate the number of recorded cells.

Fig. 2. ASICs mediate acid-induced currents in NMR and mouse CNS neurons. **a.** Both mouse (black trace, left) and NMR (blue trace, right) neurons respond to a pH 5 solution with a transient inward current. **b.** Acid-induced currents are of significantly smaller amplitude in NMR neurons compared to mouse neurons. **c.** Inactivation time constants of the acid-induced responses are similar between NMR and mouse neurons. **d.** Example trace of an acid-induced current elicited by a pH 5 solution and the effect of 100 μM benzamil in a mouse cortical neuron. **e-f.** Acid-induced currents in both cortical and hippocampal mouse neurons were reversibly blocked by 100 μM benzamil (n = 9 and n = 10 cortical and hippocampal neurons, respectively). **g.** Example trace of an acid-induced current evoked by a pH 5 solution in a cortical NMR neuron showing inhibition by 100 μM benzamil. **h-i.** In both cortical (n = 7) and hippocampal (n = 6) NMR neurons, acid-induced currents were reversibly blocked by 100 μM benzamil. ** $p < 0.01$; **** $p < 0.0001$, one-way ANOVA paired tests, Tukey's multiple comparison tests. Numbers into brackets indicates the number of recorded cells.

Fig. 3. Sensitivity of acid-induced currents to the ASIC1a blocker PcTx1. **a.** Two populations of acid-induced currents were recorded in mouse neurons: (**a.i**) those blocked by 10 nM PcTx1 and those not inhibited by 10 nM PcTx1 (**a.i**); currents were considered blocked if their peak current density was reduced by more than 50%. **b-c.** In both cortical and hippocampal mouse neurons, application of 10 nM PcTx1 reduced acid-induced current amplitudes (**b.i** – **c.i**) in most of the recorded neurons (**b.ii** – **c.ii**; 5/11 and 7/13 for cortical and hippocampal neurons, respectively). **d.** Similarly, in NMR cortical and hippocampal neurons, some acid-induced currents were blocked by 10 nM PcTx1 (**d.i**) while others were not (**d.ii**). **e-f.** In both cortical and hippocampal NMR neurons, 10 nM PcTx1 inhibited less than a quarter of the currents recorded (1/13 and 2/12 for

cortical and hippocampal neurons, respectively). * $p < 0.5$; ** $p < 0.01$, one-way paired ANOVA tests, Tukey's multiple comparison tests.

Fig. 4. PcTx1 does not inhibit nmrASIC1a. **a.** nmrASIC1a was heterologously expressed in CHO cells and currents were evoked by a 1 second pulse of pH 6.5 solution with the effect of 10 nM and 100 nM PcTx1 being recorded. **b.** No significant change in amplitude was observed when PcTx1 was applied. n = 7 cells, one-way paired ANOVA test, $p = 0.2905$.

Fig. 5. nmrASIC1a is Ca^{2+} permeable, whereas nmrASIC1b is not. Whole-cell patch clamp recordings from CHO cells heterologously expressing nmrASIC1a (**a-b**) or nmrASIC1b (**c-d**) using different cation concentrations in the extracellular solution while perfusing a pH 6 solution at different voltages ranging from -100 mV to 80 mV. nmrASIC1a is permeable to Ca^{2+}, Na^{+} and Li^{+} permeability (**a-b**), whereas nmrASIC1b is only permeable to Na^{+} and Li^{+} (**c-d**). Four to nine cells were recorded for each voltage and for each extracellular solution composition.

Fig. 6. NMR cortical neurons show resistance to acid-induced cell death. Mouse (**a**) and NMR (**b**) cortical neurons were incubated for 2 hours with either a pH 7.4 or pH 5 solution and then stained with Hoetsch 33342 to label nuclei and PI to label necrotic dead cells. **c.** Percentages of PI-positive neurons (i.e. dead) at pH 7.4 were similar between mouse and NMR neurons. The percentage of mouse dead neurons was higher when cells were incubated in a pH 5 solution, compared to pH 7.4 condition and compared to the percentage of cell death obtained with NMR neurons at pH 5. NMR cortical neurons did not exhibit any significant difference with regard to cell death when exposed to pH 5.0 compared to pH 7.4. **** $p< 0.0001$, one-way ANOVA test, Tukey's multiple comparison test. Percentages of each field of view were used to compare acidotoxicity between species. The number of fields of view analyzed for each condition are the following: n =

18 for mouse neurons (for both pH 7.4 and pH 5), n = 27 and 28 for NMR neurons at pH 7.4 and at pH 5, respectively; obtained from 3 independent cultures for both species.

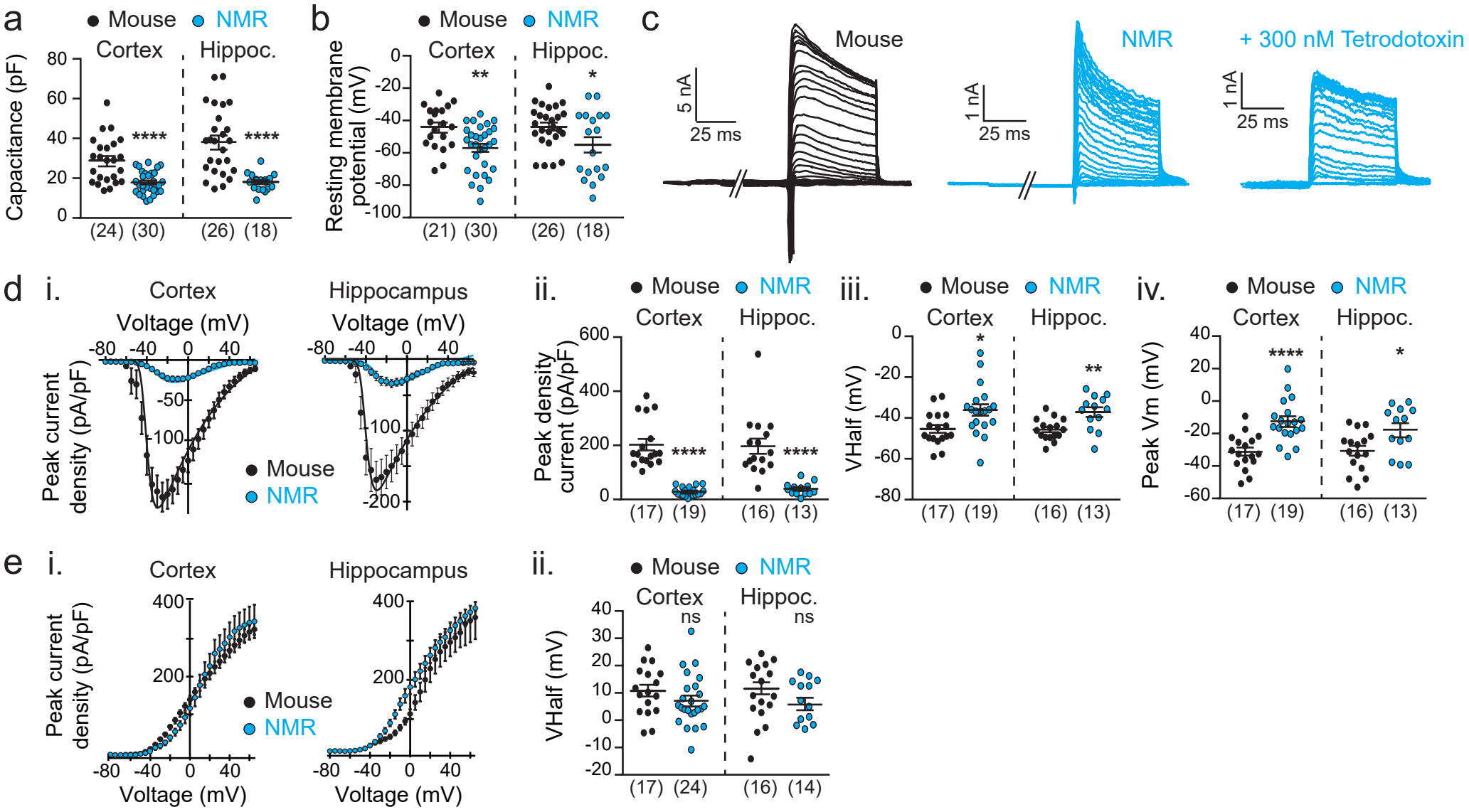
a
Mouse
NMR
Cortex
Hippoc.
Capacitance (pF)
80
60
40
20
0
****
****
(24) (30)
(26) (18)
b
Mouse
NMR
Cortex
Hippoc.
Resting membrane potential (mV)
0
-20
-40
-60
-80
-100
**
*
(21) (30)
(26) (18)
c
Mouse
5 nA
25 ms
NMR
1 nA
25 ms
+ 300 nM Tetrodotoxin
1 nA
25 ms
d
i.
Cortex
Hippocampus
Voltage (mV)
-80 -40 0 40
Peak current density (pA/pF)
-50
-100
-200
Mouse
NMR
ii.
Mouse
NMR
Cortex
Hippoc.
Peak density current (pA/pF)
600
400
200
0
****
****
(17) (19)
(16) (13)
iii.
Mouse
NMR
Cortex
Hippoc.
VHalf (mV)
0
-20
-40
-60
-80
*
**
(17) (19)
(16) (13)
iv.
Mouse
NMR
Cortex
Hippoc.
Peak Vm (mV)
40
20
0
-20
-40
-60
****
*
(17) (19)
(16) (13)
e
i.
Cortex
Hippocampus
Peak current density (pA/pF)
400
200
Mouse
NMR
-80 -40 0 40
Voltage (mV)
ii.
Mouse
NMR
Cortex
Hippoc.
VHalf (mV)
40
30
20
10
0
-10
-20
ns
ns
(17) (24)
(16) (14)

FIG. 1

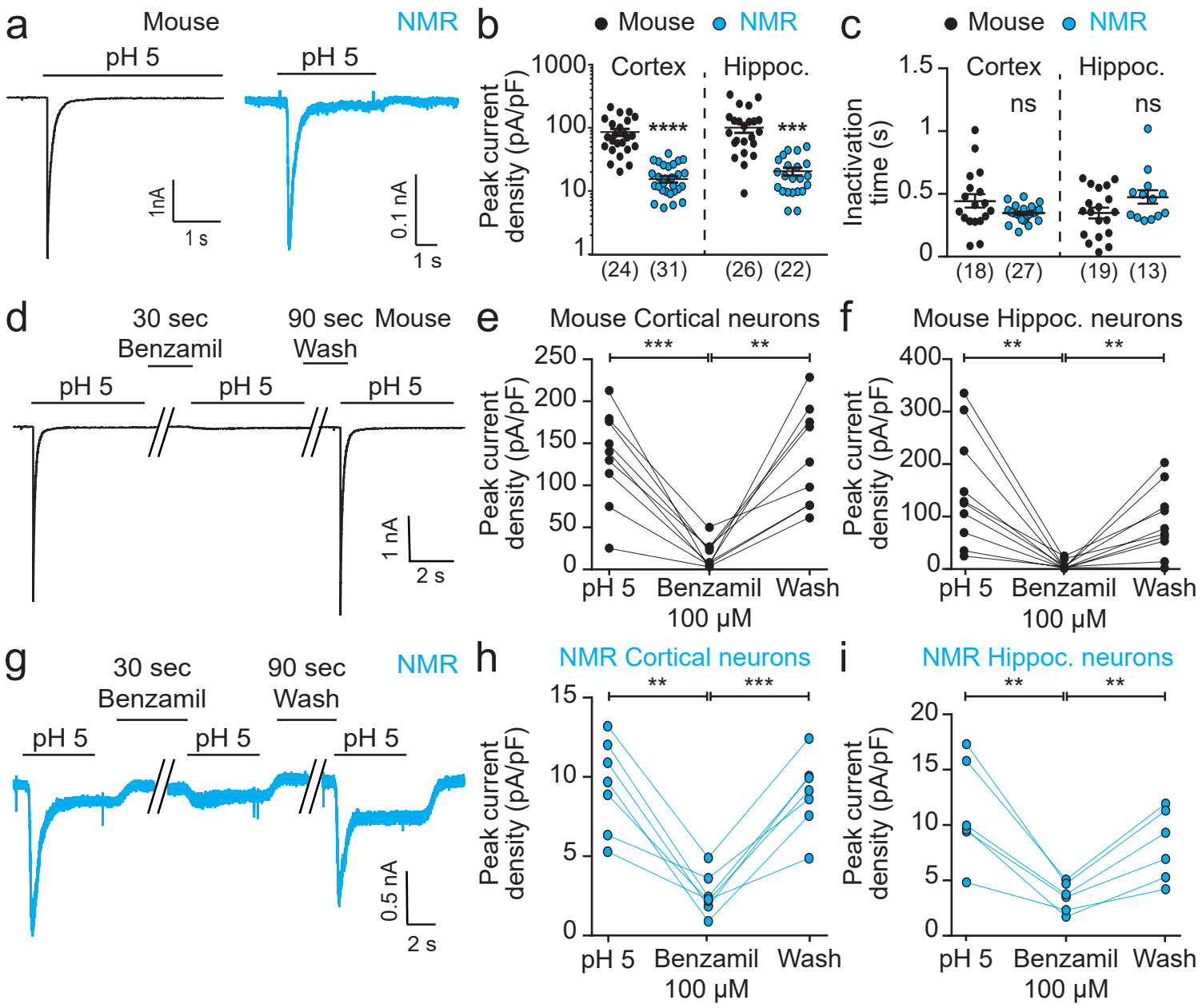
a
Mouse
NMR
pH 5
pH 5
1nA
1 s
0.1 nA
1 s
b
Mouse
NMR
Cortex
Hippoc.
Peak current density (pA/pF)
1000
100
10
1
****
***
(24) (31)
(26) (22)
c
Mouse
NMR
Cortex
Hippoc.
Inactivation time (s)
1.5
1
0.5
0
ns
ns
(18) (27)
(19) (13)
d
30 sec Benzamil
90 sec Wash
Mouse
pH 5
pH 5
pH 5
1 nA
2 s
e
Mouse Cortical neurons
Peak current density (pA/pF)
250
200
150
100
50
0
***
**
pH 5
Benzamil 100 μM
Wash
f
Mouse Hippoc. neurons
Peak current density (pA/pF)
400
300
200
100
0
**
**
pH 5
Benzamil 100 μM
Wash
g
30 sec Benzamil
90 sec Wash
NMR
pH 5
pH 5
pH 5
0.5 nA
2 s
h
NMR Cortical neurons
Peak current density (pA/pF)
15
10
5
0
**
***
pH 5
Benzamil 100 μM
Wash
i
NMR Hippoc. neurons
Peak current density (pA/pF)
20
15
10
5
0
**
**
pH 5
Benzamil 100 μM
Wash

FIG. 2

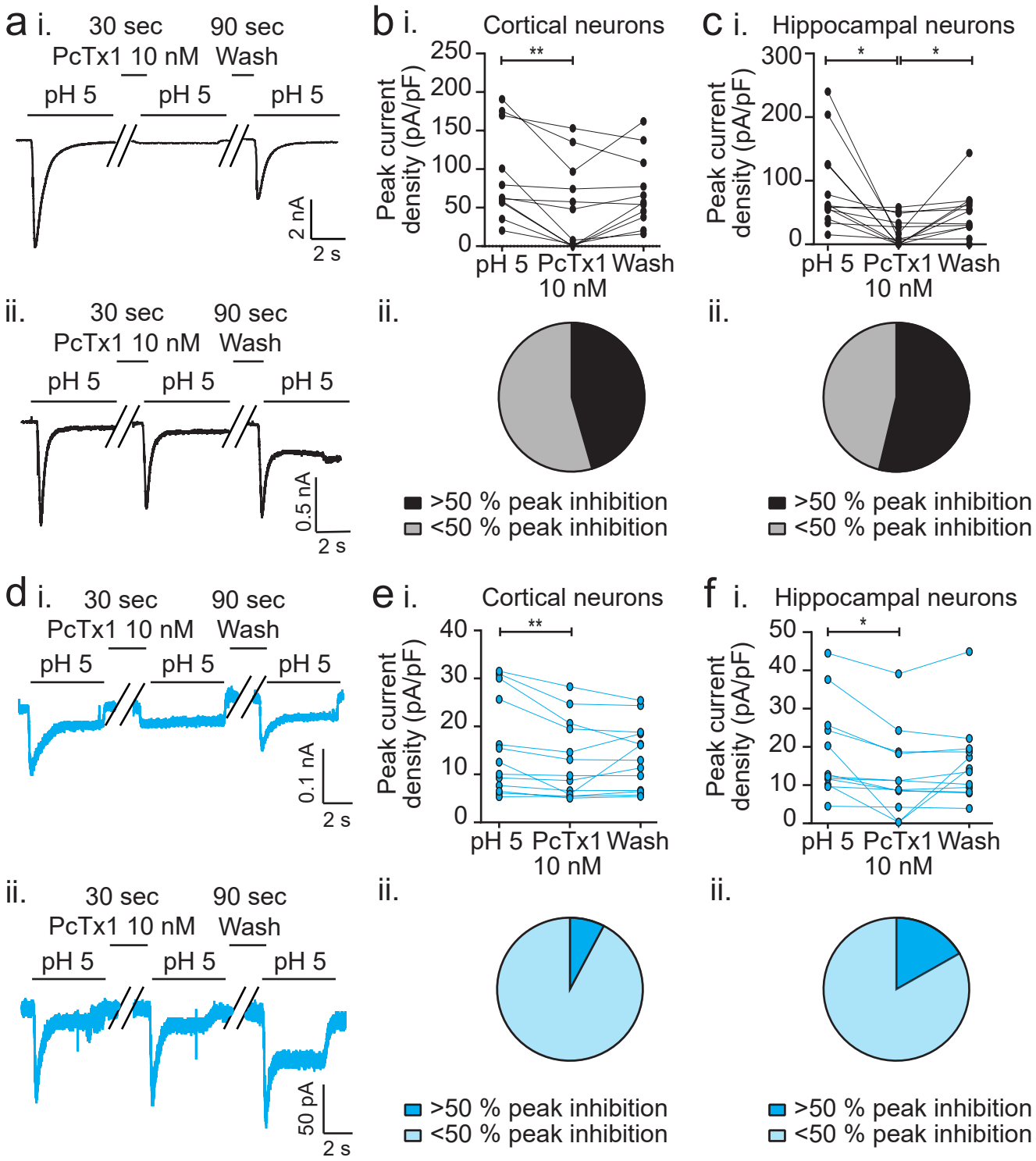
a i.
30 sec
90 sec
PcTx1 10 nM
Wash
pH 5
pH 5
pH 5
2 nA
2 s
ii.
30 sec
90 sec
PcTx1 10 nM
Wash
pH 5
pH 5
pH 5
0.5 nA
2 s
b i.
Cortical neurons
Peak current density (pA/pF)
250
200
150
100
50
0
**
pH 5
PcTx1 10 nM
Wash
ii.
>50 % peak inhibition
<50 % peak inhibition
c i.
Hippocampal neurons
Peak current density (pA/pF)
300
200
100
0
*
*
pH 5
PcTx1 10 nM
Wash
ii.
>50 % peak inhibition
<50 % peak inhibition
d i.
30 sec
90 sec
PcTx1 10 nM
Wash
pH 5
pH 5
pH 5
0.1 nA
2 s
ii.
30 sec
90 sec
PcTx1 10 nM
Wash
pH 5
pH 5
pH 5
50 pA
2 s
e i.
Cortical neurons
Peak current density (pA/pF)
40
30
20
10
0
**
pH 5
PcTx1 10 nM
Wash
ii.
>50 % peak inhibition
<50 % peak inhibition
f i.
Hippocampal neurons
Peak current density (pA/pF)
50
40
30
20
10
0
*
pH 5
PcTx1 10 nM
Wash
ii.
>50 % peak inhibition
<50 % peak inhibition

FIG. 3

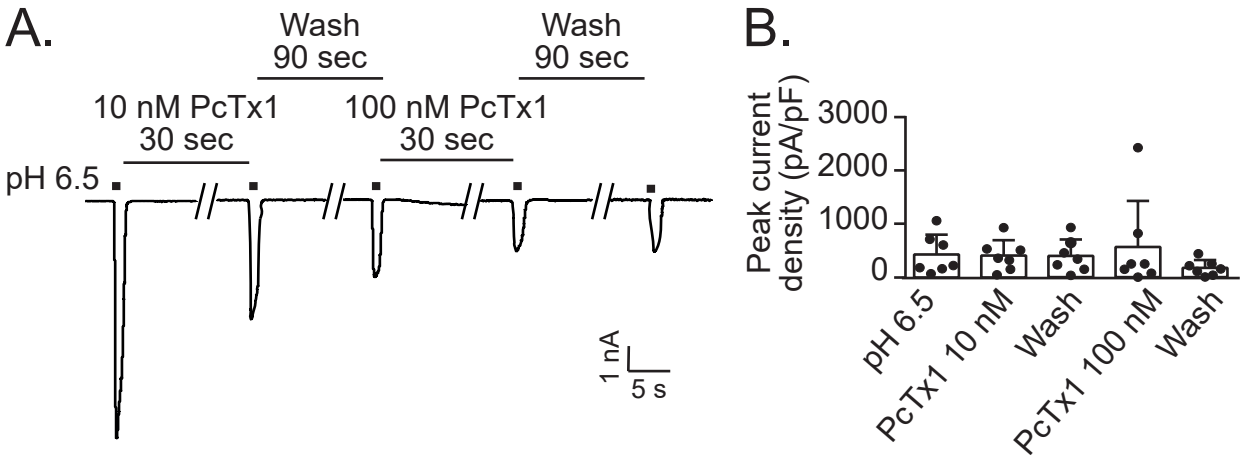
A.
Wash
90 sec
Wash
90 sec
10 nM PcTx1
30 sec
100 nM PcTx1
30 sec
pH 6.5
1 nA
5 s
B.
Peak current density (pA/pF)
3000
2000
1000
0
pH 6.5
PcTx1 10 nM
Wash
PcTx1 100 nM
Wash

FIGURE 4

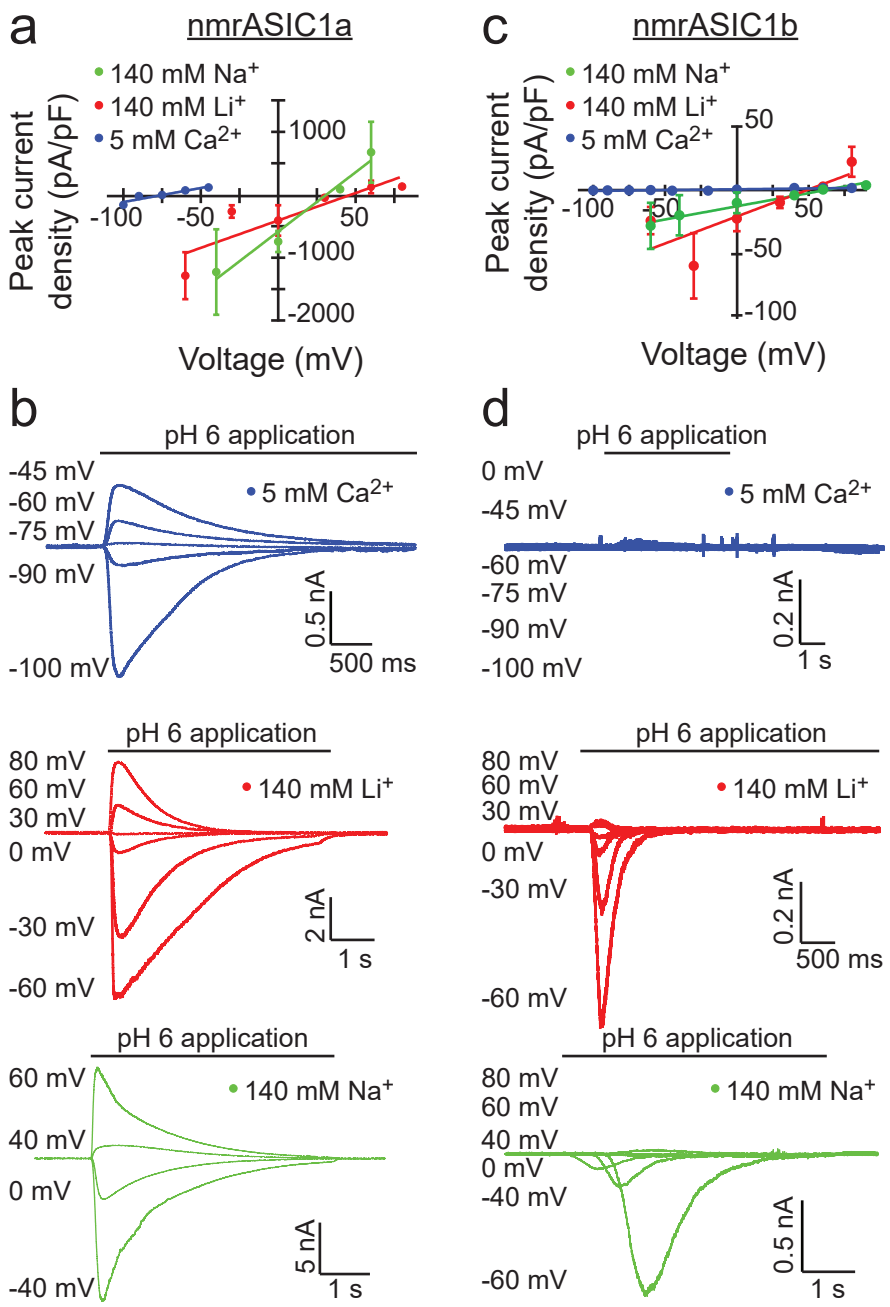
a
nmrASIC1a
140 mM Na+
140 mM Li+
5 mM Ca2+
Peak current density (pA/pF)
1000
-1000
-2000
-100
-50
50
Voltage (mV)
c
nmrASIC1b
140 mM Na+
140 mM Li+
5 mM Ca2+
Peak current density (pA/pF)
50
-50
-100
-100
-50
50
Voltage (mV)
b
pH 6 application
-45 mV
-60 mV
-75 mV
-90 mV
-100 mV
5 mM Ca2+
0.5 nA
500 ms
pH 6 application
80 mV
60 mV
30 mV
0 mV
-30 mV
-60 mV
140 mM Li+
2 nA
1 s
pH 6 application
60 mV
40 mV
0 mV
-40 mV
140 mM Na+
5 nA
1 s
d
pH 6 application
0 mV
-45 mV
-60 mV
-75 mV
-90 mV
-100 mV
5 mM Ca2+
0.2 nA
1 s
pH 6 application
80 mV
60 mV
30 mV
0 mV
-30 mV
-60 mV
140 mM Li+
0.2 nA
500 ms
pH 6 application
80 mV
60 mV
40 mV
0 mV
-40 mV
-60 mV
140 mM Na+
0.5 nA
1 s

FIG. 5

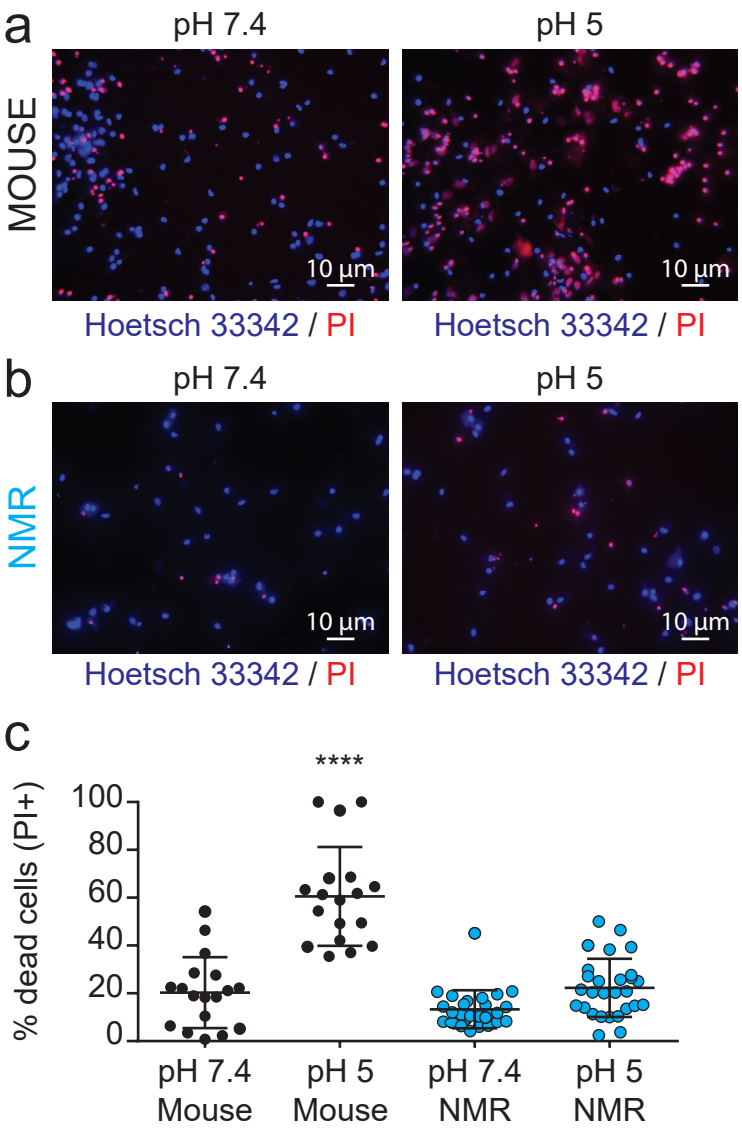
a
pH 7.4
pH 5
MOUSE
10 µm
10 µm
Hoetsch 33342 / PI
Hoetsch 33342 / PI
b
pH 7.4
pH 5
NMR
10 µm
10 µm
Hoetsch 33342 / PI
Hoetsch 33342 / PI
c
% dead cells (PI+)
100
80
60
40
20
0
****
pH 7.4 Mouse
pH 5 Mouse
pH 7.4 NMR
pH 5 NMR

FIG. 6