Swelling activated Cl\textsuperscript{-} channels in microglia
Biophysics, pharmacology and role in glutamate release

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Abbreviations: DCPIB, 3-\{aminosulfonyl\}-5-\{butylamino\}-4-phenoxybenzoic acid (bumetanide), 4-\{2-butyl-6,7-dichloro-2-cyclopentylinden-1-on-5-yl\}oxybutyric acid; FFA, fluoranamic acid; G\textsubscript{Cl\textsubscript{swell}}, instantaneous slope conductance at reversal potential; glibenclamide, 5-chloro-N-\{4-[N-(cyclohexylcarbamoyl)sulfamoyl]phenethyl\}-2-methoxybenzamide; GDP\textsubscript{S}, guanosine 5'-O-\{beta-thio\}diphosphate; GTP\textsubscript{S}, guanosine 5'-O-\{gamma-thio\}triphosphate; IAA-94, indanyloxacetic acid 94; I\textsubscript{Cl\textsubscript{swell}}, volume-activated Cl\textsuperscript{-} current; NPPB, 5'-\{5-nitro-2-\{3-phenylpropylamino\}benzoic acid\}; riluzole, 2-amino-6-\{trifluoromethoxy\}-benzothiazole; tamoxifen, (Z)-1-(\{p-dimethylamino ethoxyphenyl\})-1,2-diphenyl-1-butene

Microglia have a swelling-activated Cl\textsuperscript{-} current (which we call I\textsubscript{Cl\textsubscript{swell}}), and while some of its biophysical properties and functional roles have been elucidated, its molecular identity is unknown. To relate this current to cell functions and determine whether it is regulated by mechanisms other than cell swelling, it is important to establish both biophysical and pharmacological fingerprints. Here, we used rat microglia and a cell line derived from them (MLS-9) to study biophysical, regulatory and pharmacological properties of I\textsubscript{Cl\textsubscript{swell}}. The whole-cell current was activated in response to a hypo-osmotic bath solution, but not by voltage, and was time-independent during long voltage steps. The halide selectivity sequence was I>Br>Cl (Eisenman sequence I) and importantly, the excitatory amino acid, glutamate was permeant. Current activation required internal ATP, and was not affected by the guanine nucleotides, GTP\textsubscript{S} or GDP\textsubscript{S}, or physiological levels of internal Mg\textsuperscript{2+}. The same current was activated by a low intracellular ionic strength solution without an osmotic gradient. I\textsubscript{Cl\textsubscript{swell}} was reversibly inhibited by known Cl\textsuperscript{-} channel blockers (NPPB, fluoranamic acid, glibenclamide, DCPIB), and by the glutamate release inhibitor, riluzole. Cell swelling evoked glutamate release from primary microglia and MLS-9 cells, and this was inhibited by the blockers (above), and by IAA-94, but not by tamoxifen or the Na\textsuperscript{+}/K\textsuperscript{+}/Cl\textsuperscript{-} symport inhibitor, bumetanide. Together, these results confirm the similarity of I\textsubscript{Cl\textsubscript{swell}} in the two cell types, and point to a role for this channel in inflammation-mediated glutamate release in the CNS.

Introduction

After acute CNS injury, microglia respond by retracting their long processes, and changing morphology and volume as they migrate to the site of damage and become phagocytic. We identified a Cl\textsuperscript{-} current in rat microglia that is activated by cell swelling.\textsuperscript{1} The regulatory volume decrease (RVD) that follows cell swelling requires Cl\textsuperscript{-} and K\textsuperscript{+} efflux and accompanying water loss. By constructing an extensive pharmacological fingerprint of the Cl\textsuperscript{-} current (which we call I\textsubscript{Cl\textsubscript{swell}}), we showed that in rat microglia it contributes to both homeostatic volume regulation and RVD,\textsuperscript{2} and to phagocytosis\textsuperscript{3} and microglial proliferation.\textsuperscript{1} Multiple functional roles are anticipated because the Cl\textsuperscript{-} channel regulates the membrane potential of microglia.\textsuperscript{3} Several Cl\textsuperscript{-} channels have been described in mammalian cells, and while some have been cloned, the molecular identity of swelling-activated Cl\textsuperscript{-} currents is unknown (reviewed in refs. 4 and 5). There is biophysical and pharmacological evidence that I\textsubscript{Cl\textsubscript{swell}} in immune cells differs from the more common volume-regulated anion channel (‘VRAC’).\textsuperscript{2} Therefore, it is important to distinguish I\textsubscript{Cl\textsubscript{swell}} from other Cl\textsuperscript{-} currents when assessing its roles in microglia, and studying its regulation by factors other than cell swelling. Identifying the gene underlying the I\textsubscript{Cl\textsubscript{swell}} in microglia has been severely hampered by the difficulty of transfected primary microglia.\textsuperscript{2} We previously derived a cell line (called MLS-9) from rat microglia, and found that it can readily be transfected.\textsuperscript{4} If this cell line expresses the same I\textsubscript{Cl\textsubscript{swell}}\textsuperscript{b} it will be useful for future studies aimed at identifying the channel gene, and identifying its regulatory interactions with other molecules. Therefore, we first addressed whether MLS-9 cells express an I\textsubscript{Cl\textsubscript{swell}} with the same properties as the current in primary microglia.
Rese... If these anion channels release excitatory amino acids, microglia are ideally positioned to affect other CNS cells, because they form an essentially continuous network in the healthy CNS, with long processes abutting neurons and glial cells. Furthermore, after injury, microglia migrate to the site of injury, as shown in our recent studies of K+ channels in rat models of optic nerve transection, ischemic stroke and intracerebral hemorrhage.8,10 In addition, glutamate is involved in signaling between neurons, astrocytes and microglia; thus, potential outcomes of activating IClswell include trans-activating NMDA receptors to evoke excitotoxic neuronal injury;11 activating microglia through their metabotropic glutamate receptors, leading to production of other neurotoxic mediators;12,13 and mediating glia-to-neuron signaling under inflammatory conditions.14

Here, we first show that the biophysical and pharmacological properties of IClswell in MLS-9 cells are as previously determined for primary rat microglia. We then assessed current regulation by intracellular nucleotides and ionic strength, and constructed a pharmacological profile that we used to determine the contribution of this channel to glutamate release. Together, our results suggest that IClswell is a potential target for reducing glutamatergic signaling between microglia and other CNS cells and for inhibiting neurotoxicity after acute CNS injuries.

Results

Cell swelling activates a chloride current (IClswell) in microglia. Throughout this study, whole-cell recordings were used with solutions and voltage protocols designed to isolate anion currents from the several cation currents in microglia (e.g., Kv1.3, KCa2.3, KCa3.1). For electrophysiological recordings (except when analyzing anion selectivity), the bulky cation, NMDG+, replaced Na+ and K+ in the bath (Solution 4; see Methods for all solutions) and potassium channels were inactivated using a holding potential of 0 mV.

When voltage ramps were applied to microglia bathed in iso-osmotic solution (containing 144 mM Cl-), the whole-cell current was negligible (Fig. 1A and C). After acute CNS injury (e.g., stroke, trauma), the excitatory amino acid, glutamate, causes neuronal injury; thus, a crucial finding was that IClswell channels in microglia are significantly permeable to glutamate (and aspartate).1,2 If these anion channels release excitatory amino acids, microglia are ideally positioned to affect other CNS cells, because they form an essentially continuous network in the healthy CNS, with long processes abutting neurons and glial cells. Furthermore, after injury, microglia migrate to the site of injury, as shown in our recent studies of K+ channels in rat models of optic nerve transection, ischemic stroke and intracerebral hemorrhage.8,10 In addition, glutamate is involved in signaling between neurons, astrocytes and microglia; thus, potential outcomes of activating IClswell include trans-activating NMDA receptors to evoke excitotoxic neuronal injury;11 activating microglia through their metabotropic glutamate receptors, leading to production of other neurotoxic mediators;12,13 and mediating glia-to-neuron signaling under inflammatory conditions.14

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Results

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When voltage ramps were applied to microglia bathed in iso-osmotic solution (containing 144 mM Cl-), the whole-cell current was negligible (Fig. 1A and C). After hypo-osmotic solution 5 (~60% of normal osmolarity) was perfused into the bath, the cell swelled within minutes (Fig. 1A) and an outward-rectifying current activated, increased with time (vertical arrow) and reached a quasi-stationary plateau. (B) When re-exposed to iso-osmotic bath (Solution 4), the cell shrank (inset) and the current declined to its pre-swelling level. (C) The instantaneous slope conductance was calculated at the experimentally measured reversal potential (Erev) by fitting each current trace with a mono-exponential function and taking the derivative at Erev. The instantaneous slope conductance was then normalized to the cell size (capacitance in pF), and plotted as a function of time after establishing the whole-cell recording. Current activation and deactivation are shown during two cycles of swelling and shrinking (same cell as in A and B).

Figure 1. Cell swelling activates a large Cl- current. A representative whole-cell recording of currents in an MLS-9 cell in response to voltage ramps from -100 to +80 mV, from a holding potential of -10 mV. (A) The first five sweeps show a very small current in iso-osmotic NMDG+ bath solution (Solution 4; 177 mOsm/kgH2O; see Methods). When the bath was perfused with the hypo-osmotic Solution 5, the cell swelled (inset) and an outward-rectifying current activated, increased with time (vertical arrow) and reached a quasi-stationary plateau. (B) When re-exposed to iso-osmotic bath (Solution 4), the cell shrank (inset) and the current declined to its pre-swelling level. (C) The instantaneous slope conductance was calculated at the experimentally measured reversal potential (Erev) by fitting each current trace with a mono-exponential function and taking the derivative at Erev. The instantaneous slope conductance was then normalized to the cell size (capacitance in pF), and plotted as a function of time after establishing the whole-cell recording. Current activation and deactivation are shown during two cycles of swelling and shrinking (same cell as in A and B).
potential for Cl with 54 mM Cl in the pipette and 74 mM Cl\(^{-}\) in the bath. Ion substitution experiments (see below) confirmed the anion selectivity of the swelling-activated current. When the original iso-osmotic bath was restored (Fig. 1B), the cell shrank back to its original size and the current decreased over several minutes to a very low level. By plotting the instantaneous slope conductance calculated at \(E_{\text{rev}}\) (Fig. 1C), the time course of current activation and its swelling sensitivity is clearly illustrated during repeated cycles of swelling and shrinking. A less hypo-osmotic solution (75–80% of normal) more slowly activated a more variable current (not shown). For the remainder of the study, the current was activated using the 60% hypo-osmotic bath solution (Solution 5).

The swelling-activated Cl current in primary rat microglia and the MLS-9 cell line showed neither time-dependent activation nor inactivation during 2-sec long voltage clamp steps (Fig. 2A and C). Current-versus-voltage (I–V) relations were constructed from step currents and superimposed on ramp currents from the same cells (Fig. 2B and D). As above, \(E_{\text{rev}}\) (about -8 mV after junction potential correction) was very close to the calculated Cl Nernst potential, and there was significant outward rectification despite the similarity in internal and external Cl concentrations. Our subsequent use of the ramp protocol to monitor the conductance was supported by the similarity of I–V relations generated by ramp and step protocols.

The relative permeability of the channels to different anions (Table 1) was determined by first activating the current with a hypo-osmotic bath containing 120 mM NaCl, and then exchanging the bath with a hypo-osmotic solution containing 120 mM of the Na salt of the test anion (I, Br, glutamate). The reversal potential was measured with each bath solution, and the change (\(\Delta E_{\text{rev}}\)) calculated; from which the permeability ratio (\(P_{x}/P_{\text{Cl}}\)) was calculated using the modified Goldman-Hodgkin and Katz (GHK) voltage equation (Eqn. 1).

\[
P_{x/\text{Cl}} = \frac{[\text{Cl}]_{x\text{ before}} \exp(\Delta E_{\text{rev}} \cdot Z_{x}/RT)}{[\text{Cl}]_{x\text{ after}}}[A_{x\text{ after}}]/[A_{\text{Cl before}}]
\]

(Eqn. 1)

where \([\text{Cl}]_{x\text{ before}}\) is the initial 74 mM Cl\(^{-}\) concentration, \([\text{Cl}]_{x\text{ after}}\) is the 4 mM Cl\(^{-}\) concentration remaining after changing the external anion, \([A_{x}\text{ after}]\) is the concentration of the test anion after the solution change, and \(z\) is the valence. For each anion, several cells were tested and the permeability ratios were calculated and averaged. Importantly, for these experiments, a 3 M KCl agar bridge was used to prevent junction potential changes when the anion species was changed in the bath. With the 120 mM NaCl bath solution, the expected Nernst potential for Cl was -20 mV. The average reversal potentials, and calculated relative anion permeabilities (Table 1) indicate a permeability sequence of I > Br \(\geq\) Cl > glutamate, where > denotes a significant difference (p < 0.05) and \(\geq\) indicates a trend that did not reach statistical significance. The sequence among the halides corresponds with Eisenman sequence I, and is similar to swelling activated anion currents in many other cell types (see Discussion). The relative permeability for F could not be determined because NaF substitution activated a large unidentified current (not shown).

Spontaneous activation of the Cl current by low intracellular ionic strength. With a low-ionic strength pipette solution and essentially iso-osmotic bath the Cl current spontaneously activated and reached a peak by ~5 min (Fig. 3A). The cells

| Table 1. Anion selectivity of \(I_{\text{Cl}\text{swell}}\) in MLS-9 cells |
|-----------------|---------------|----------------|------------------|
| Anion           | \(E_{\text{rev}}\) (mV) ± SE | \(P_{x}/P_{\text{Cl}}\) ± SE | Number of cells |
| I               | -26.2 ± 0.4   | 1.15 ± 0.03    | 5               |
| Br              | -23.5 ± 0.4   | 1.03 ± 0.04    | 5               |
| Cl              | -22.9 ± 0.6   | 1.00 ± 0.00    | 5               |
| Glutamate       | 16.6 ± 1.3    | 0.12 ± 0.01†   | 7               |

\(I^{+}\) permeability is higher than Br (p < 0.05). Glu permeability is lower than any of the halides (p < 0.01).
did not obviously swell under these conditions, but to eliminate this possibility, a group of recordings was made with a hypo-osmotic, low ionic strength pipette solution. Not only did the current develop spontaneously with both pipette solutions, but also the time course and maximal conductance were similar. With the hypo-osmotic pipette solution, $G_{Cl}$ reached a peak conductance of $0.68 \pm 0.06 \text{nS/pF}$ ($n = 5$) with a half-maximal activation time ($t_{1/2}$) of $149 \pm 11$ sec. With the iso-osmotic pipette solution, peak $G_{Cl}$ was $0.67 \pm 0.07 \text{nS/pF}$ ($n = 5$) and $t_{1/2}$ was $160 \pm 12$ sec.

With low ionic-strength pipette solution (Fig. 3B and solid curve), the current spontaneously activated and at the indicated $t_{1/2}$ (160 sec), $G_{Cl}$ reached $0.35 \pm 0.03 \text{nS/pF}$ (Fig. 3C and closed circles; $n = 5$). With high ionic-strength pipette solution (Fig. 3B and dashed curve), the current did not spontaneously activate; $G_{Cl}$ was only $0.06 \pm 0.02 \text{nS/pF}$ at 160 sec (Fig. 3C and open circles; $n = 12$; $p < 0.05$). Nevertheless, a swelling-induced conductance developed in hypo-osmotic bath Solution 5 (Fig. 3B and seen in 11/11 cells).

**Figure 3.** Spontaneous activation of the Cl$^-$ current by low intracellular ionic strength. The instantaneous slope conductance was calculated at $E_e$ and normalized to the cell capacitance (as in Fig. 1C), and plotted as a function of time after establishing whole-cell recordings. (A) The Cl$^-$ current spontaneously activated (open circles; $n = 5$) when whole-cell recordings were established with a low ionic strength pipette solution (Solution 2; ionic strength, 76 mM; see Methods) that was nominally iso-osmotic with the bath (Solution 4; ~300 mOsm/kgH$_2$O). Similar current activation was seen (closed circles; $n = 5$) when the osmolarity of low ionic strength pipette solution was reduced to ~260 mOsm/kgH$_2$O by omitting sucrose. (B) The Cl$^-$ conductance rapidly and spontaneously developed with low ionic strength pipette solution (solid curve; $t_{1/2}$ = time required for half-maximal activation). Spontaneous activation was prevented when the ionic strength of the pipette solution was 146 mM (dashed curve; 80 KAsp/40 KCl) but the current was activated by cell swelling when the bath was perfused with hypo-osmotic Solution 5. (C) Expanded time scale to show rapid activation of the Cl$^-$ conductance with low ionic strength pipette solution (closed circles; $n = 5$), and lack of activation with high ionic strength pipette solution (open circles; *p* < 0.05, $n = 12$).

**Activation of the Cl$^-$ current requires intracellular ATP.** The role of adenine nucleotides in regulating the anion channels was investigated by excluding ATP from the normal pipette solution (Solution 1). After establishing each whole-cell recording, we waited 5–10 min to allow endogenous ATP to diffuse out of the cell, and then perfused in a hypo-osmotic bath (Solution 5). The peak $G_{Cl}$ in response to cell swelling (Fig. 4A) was $0.06 \pm 0.01 \text{nS/pF}$ without intracellular ATP versus $1.47 \pm 0.06 \text{nS/pF}$ with 2 mM ATP ($n = 3$ cells each). Because omitting ATP increases free intracellular $\text{Mg}^{2+}$ from 0.08 to 0.6 mM, we next tested the effect of reducing $\text{Mg}^{2+}$ in the absence of ATP (Fig. 4B). The swelling-induced conductance was not significantly different when $\text{Mg}^{2+}$ was reduced without ATP; peak $G_{Cl}$ was $0.26 \pm 0.08 \text{nS/pF}$ ($\text{Mg}^{2+}$ 0.08 mM) compared with $0.10 \pm 0.02 \text{nS/pF}$ ($\text{Mg}^{2+}$ 0.6 mM).

To determine whether guanine nucleotides can substitute for ATP, GTP$\gamma$S was added to the ATP-free patch pipette solution (Fig. 4C). GTP$\gamma$S did not support activation of the swelling-induced Cl$^-$ conductance; peak $G_{Cl}$ was $0.25 \pm 0.04 \text{nS/pF}$ (200 $\mu$M GTP$\gamma$S) versus $0.09 \pm 0.03 \text{nS/pF}$ without ATP. Not surprisingly then, the presence of GDP$\beta$S with ATP in the pipette (Fig. 4D) did not alter the peak swelling-induced $G_{Cl}$, which was $0.77 \pm 0.16 \text{nS/pF}$ ($n = 6$) compared with $0.83 \pm 0.12 \text{nS/pF}$ ($n = 7$) without GDP$\beta$S. Finally, we asked whether intracellular GTP$\gamma$S affects the spontaneous current activation seen with a low ionic-strength pipette solution containing ATP. As expected from results in Figure 3A, the current spontaneously activated (Fig. 4E), and reached a peak conductance of $0.66 \pm 0.07 \text{nS/pF}$ ($n = 5$). Surprisingly, GTP$\gamma$S increased the conductance to $1.05 \pm 0.10 \text{nS/pF}$ ($n = 8$; $p < 0.05$).

**Pharmacological profile of the volume-sensitive Cl$^-$ channel.** After inducing the swelling-activated current with hypo-osmotic Solution 5 and waiting for the conductance to reach a quasi-stable plateau, we perfused in the same hypo-osmotic solution containing an inhibitor. Figure 5 shows representative Cl$^-$ currents in response to voltage ramps before and after adding NPPB, glibenclamide, riluzole or DCPIB; and conductance versus time graphs, which show blocker reversibility. Table 2 summarizes the percent block with each drug, calculated by comparing
that of Cl− (Table 1). We previously reported similar glutamate permeability for the channel in primary microglia.1,2 Thus, we asked whether this channel might contribute to glutamate release from swollen MLS-9 and primary microglial cells. Extracellular glutamate accumulation was measured in the medium after swelling the cells in hypo-osmotic medium, with or without each of the blockers used in Table 2. For comparison, we also used bumetanide, a Na+/K+/Cl− co-transport inhibitor.

A functional role for the Cl− channel in glutamate release. We found that the swelling-activated anion channel in MLS-9 cells is permeable to glutamate with a relative permeability about 12% of that of Cl− (Table 1). We previously reported similar glutamate permeability for the channel in primary microglia.1,2 Thus, we asked whether this channel might contribute to glutamate release from swollen MLS-9 and primary microglial cells. Extracellular glutamate accumulation was measured in the medium after swelling the cells in hypo-osmotic medium, with or without each of the blockers used in Table 2. For comparison, we also used bumetanide, a Na+/K+/Cl− co-transport inhibitor.

When the normal iso-osmotic bath (Solution 3) was changed to hypo-osmotic Solution 6, glutamate release increased by 280%
Neither bumetanide nor a very low concentration of tamoxifen (1 μM) affected glutamate release. In primary rat microglia, hypo-osmotic Solution 5 induced a 210% increase in glutamate release, and this was inhibited by NPPB, IAA-94 and flufenamic acid, but not by the Na+/K+/Cl- symport inhibitor, bumetanide.

Discussion

The first goal of this study was to create a biophysical fingerprint of I_{Clswell} in MLS-9 cells, and compare it with the rat microglia from which they were derived, and which we previously characterized.1,2 For all properties assessed, currents in the two cells were indistinguishable, including time- and voltage independent gating, mild outward rectification in symmetrical Cl- solutions, and a broad anion permeability, with I- > Br- > Cl- >> glutamate. The same I_{Clswell} was activated by a hypo-osmotic bath, or by reducing the internal ionic strength, a treatment that activates I_{Clswell} in some cells without swelling (reviewed in ref. 15). A stretch-induced Cl- current in microglia16 has similar outward rectification, voltage-independent activation and lack of inactivation. Although ‘volume-regulated anion channels’ (VRAC) are present in many cell types (recently reviewed in refs. 4, 5 and 17), differences in biophysical and pharmacological properties between classical VRAC and I_{Clswell} indicate more than one molecular entity.2 The microglial current is most similar to I_{Clswell} in lymphocytes.18,19 Properties that distinguish the immune cell current from other cell types include lack of inactivation at strongly depolarized potentials, even in the presence of divalent cations; lack of block by external SITS, DIDS or ATP; and a very low single-channel conductance (1–3 pS).1,2,16 Other similarities between I_{Clswell} in

Figure 5. Pharmacological profile of the volume-sensitive Cl- channel. Representative experiments illustrate reversible inhibition of the Cl- current by 125 μM NPPB (A), 500 μM glibenclamide (B), 300 μM riluzole, (C) and 20 μM DCPIB (D). For each compound, the upper part shows currents in response to voltage ramps before swelling (traces marked ‘1’), during the plateau phase at the peak of swelling-induced activation (marked ‘2’), and after adding the inhibitor (marked ‘3’). Each lower graph shows instantaneous slope conductance, calculated at E_{rev} (as in Fig. 1C) versus time after establishing whole-cell recordings.

(to 2.80 ± 0.22 a.u.; p < 0.01; Fig. 6A), and this was inhibited by NPPB and flufenamic acid at concentrations that block the swelling-induced Cl- current in MLS-9 cells (Table 2), and in primary microglia.1,2 Specifically, at 20 μM, DCPIB displayed similar inhibition to NPPB; both reduced glutamate release to the baseline level. Glutamate release was inhibited by the Cl channel blocker, IAA-94, at a concentration that reduces the swelling-induced current in primary microglia,1,2 and by riluzole at a concentration that reduced the Cl- conductance in MLS-9 cells (Table 2).
lymphocytes and microglia are the selectivity sequence, activation by low internal ionic strength, and the pharmacological profile (see below). $I_{\text{Clwell}}$ differs from VRAC (also called 'volume-sensitive organic anion channel', VSOAC) in other cells, which have a larger single-channel conductance (20–70 pS), are blocked by external SITS, DIDS and ATP, and exhibit voltage-sensitive inactivation, with a variable time course that might depend on extracellular Mg$^{2+}$ (reviewed in refs. 4, 5 and 17). Properties of the currents that show similarities across cell types are lack of voltage- and time-dependent activation, broad permeability to anions (following Isenman sequence I), and mild outward rectification in symmetrical Cl$^{-}$ solutions.

Our observation of spontaneous activation of $I_{\text{Clwell}}$ in MLS-9 cells with a low ionic strength pipette solution is consistent with our earlier studies of human T lymphocytes, and primary rat microglia. Activation apparently did not require swelling in the present study; the osmolarity of the low ionic-strength pipette solution was reduced to avoid swelling. Moreover, Cl$^{-}$ was present in the pipette solution, but we cannot rule out residual ATP inside the cell.

Anion channel blockers are not perfectly selective; hence, we used several blockers from different chemical classes to construct a pharmacological toolbox for comparing the current in MLS-9 cells, primary microglia and other cell types, and for assessing its functional role in glutamate release. $I_{\text{Clwell}}$ in MLS-9 cells was reversibly blocked by NPPB and by flufenamic acid at concentrations (<200 μM) that block the current in primary rat microglia,24 and by 500 μM glibenclamide, which is an inhibitor of VRAC and of CFTR Cl$^{-}$ channels, but also inhibits ATP-sensitive K$^{+}$ channels.27 Of note, $I_{\text{Clwell}}$ in microglia was reversibly blocked by riluzole, which is considered a neuroprotective drug and is in clinical trials for amyotrophic lateral sclerosis (ALS) and spinal cord injury. These CNS disorders involve inflammation, and while riluzole reduces inflammation

Table 2. Cl$^{-}$ current reduction by pharmacological compounds

| Compound name | Number of cells | Concentration (μM) | Inhibition (%) |
|---------------|----------------|--------------------|---------------|
| NPPB          | 3              | 125                | 86.9 ± 2.2    |
| Flufenamic acid| 3              | 100                | 53.1 ± 7.7    |
| Glibenclamide | 3              | 200                | 78.4 ± 5.4    |
| DCPiB         | 4              | 20                 | 82.8 ± 3.0    |
| Riluzole      | 6              | 100                | 22.4 ± 1.9    |
|               | 10             | 300                | 74.8 ± 0.9    |
|               | 3              | 400                | 88.3 ± 2.5    |

![Figure 6](image-url)

Figure 6. A functional role for the Cl$^{-}$ channel in glutamate release. MLS-9 cells (A) or primary rat microglia (B) were exposed to hypo-osmotic Solution 5 (205 mOsm/kg H$_{2}$O) for 1 hr with or without a pharmacological compound at the indicated concentration. The concentration of extracellular glutamate was quantified with the Amplex® Red Kit, after background subtraction, and removing contaminating endogenous H$_{2}$O$_{2}$ (see Methods). Glutamate release was normalized to the control group and reported in arbitrary units (a.u.), with the number of individual experiments indicated on each bar. Comparisons of control versus drug treatments are indicated as *p < 0.05, **p < 0.01. Drug treatments are compared with hypo-osmotic solution: †p < 0.05, ††p < 0.01.
in vivo, this has been interpreted to result from reduced neuron death. The ability of riluzole to inhibit neuronal voltage-gated Na⁺ channels and glutamate receptors is usually cited as the mechanism by which it reduces glutamate release in vivo, and from nerve terminals in vitro (reviewed in ref. 28). However, riluzole has pleiotropic effects on ion channels; it also blocks VRAC in human glioma cells,22 and activates small-conductance Ca²⁺-activated K⁺ channels in neurons.23 It is important to consider whether the neuroprotective and anti-inflammatory actions of riluzole in vivo also involve inhibition of microglial activation.

Based on pharmacological parallels between I_{Clswell} and cell functions, several roles for this channel have been proposed in microglia and other cells. Because of its activation by cell swelling, the most obvious role is in the regulatory volume decrease, and we demonstrated such a role for I_{Clswell} in primary rat microglia.2 However, we also found that in rat microglia I_{Clswell} contributes to the membrane potential,3 their proliferation,1 and their ability to phagocytose bacteria.2 In murine microglia, membrane stretch slowly induced a Cl⁻ current, and cell morphology was altered by several drugs, including SITS and DIDS.16,31 The underlying mechanism likely differs from I_{Clswell} in rat microglia, which is not blocked by SITS or DIDS at normal membrane potentials.1,2 It is important to note that SITS and DIDS are potent inhibitors of Cl⁻/HCO₃⁻ exchangers, which confounds their use in cell function studies. An anion channel (which the authors called VRAC) contributed to zymosan-induced glutamate release from microglia and was enhanced by respiratory burst activity,32 but the authors concluded that it was a different entity from the I_{Clswell} we described here and previously.1,2 In other cell types (reviewed in ref. 17), pharmacological studies support roles for I_{Clswell} in proliferation,9,33,34 migration,35,36 and apoptosis.37

The channels underlying I_{Clswell} in rat microglia and MLS-9 cells have a substantial permeability to the excitatory amino acids, aspartate (p_{aspartate}/p_{Cl⁻} = 0.22) and glutamate (p_{glutamate}/p_{Cl⁻} = 0.12 to 0.18; present study). We found that glutamate release from rat microglia and MLS-9 cells was evoked by cell swelling, and was inhibited by several drugs that block I_{Clswell}: NPPB, IAA-94, flufenamic acid, DCPIB and riluzole. Bumetanide, an inhibitor of the Na⁺/K⁺/Cl⁻ symporter, did not reduce glutamate release, nor did tamoxifen (tested on MLS-9 cells only), which is an estrogen response modifier that also inhibits some volume-regulated anion channels (reviewed in ref. 4 and 34). VRAC is one of several pathways involved in glutamate release from glial cells.11,17,38 Release of excitatory amino acids from microglia39 is consistent with a growing body of evidence linking inflammation to excitotoxic damage of neurons. We13 and others12 have shown that rat microglia can be activated through metabotropic glutamate receptors, and then become capable of killing neurons.

Because microglia are closely associated with neurons and synapses, and can efficiently synthesize and release glutamate, they are poised to contribute to neuron injury. Our results, together with the literature, suggest that I_{Clswell} can contribute to the increases in extracellular glutamate observed after acute CNS injuries (e.g., stroke, trauma), and their therapeutic potential should be considered. If, as our present and earlier data suggest, the gene underlying I_{Clswell} in immune cells differs from other cell types, this channel might be a more specific target for reducing inflammation.

**Materials and Methods**

**Cell cultures.** Primary microglia were isolated from brains of 1- to 2-day-old Sprague Dawley rats, as before.2,13,40 Whole brain tissue (without meninges) was mashed through a stainless steel sieve, suspended in minimal essential medium (MEM) and centrifuged (10 min, 1,000 g). The cell pellet was re-suspended and seeded into flasks with MEM with 10% fetal bovine serum (FBS) and 100 μM gentamycin. Cell culture reagents were purchased from Invitrogen (Burlington, ON, Canada) unless otherwise indicated. After culturing for two days, cellular debris, non-adherent cells and supernatant were removed and fresh medium was added to the flask. The mixed cultures were allowed to grow for 7–10 days and then shaken for 3 h on an orbital shaker at 8–10 Hz in a standard tissue culture incubator. Detached microglia in the supernatant were centrifuged (10 min, 1,000 g), re-suspended and plated at 3.5 x 10⁴ cells per 15 mm diameter cover slip for electrophysiology or in 12-well tissue culture plates (BD Falcon, Mississauga, ON Canada) at 10⁵ cells/well for glutamate measurements. The highly purified cultures were 99%–100% microglia, as judged by labeling with isopelectin B4, tomato lectin (Sigma, St. Louis, MO) and OX-42 antibody (Serotec, Raleigh, NC). Before experiments, microglia were cultured for another 1–5 days in reduced serum (2% FBS) to establish a more resting state, indicated by low expression of several inflammatory molecules and membrane receptors.46

We derived the MLS-9 cell line by treating pure cultures of rat microglia with colony-stimulating factor-1, and have used it for electrophysiology studies of K⁺ channels,4 and pharmacological studies of retroviral drug transport.42,43 MLS-9 cells were thawed and cultured for several days in MEM with 10% FBS and 100 μM gentamycin. For experiments, cells were harvested with phosphate buffered saline (PBS) containing 0.25% trypsin and 0.1% EDTA, washed with MEM, centrifuged (10 min, 1,000 g) and re-suspended, and then plated in the culture medium at 4.5 x 10⁴ cells/coverslip for electrophysiology, or at 10⁵ cells/well in 12-well plates for glutamate measurements.

**Patch-clamp recordings.** Whole-cell recordings were made with an Axopatch 200A amplifier (Molecular Devices, Sunnyvale, CA), digitized with a DigiDATA 1322A board, filtered at 5 kHz and sampled at 10 kHz. Pipettes (4–7 MΩ resistance) were pulled from thin wall borosilicate glass (WPI, Sarasota, FL) on a Narishige puller (Narishige Scientific, Setagaya-Ku, Tokyo). Data were acquired and analyzed with pCLAMP software (ver 9; Molecular Devices). Junction potentials, which were reduced by using agar bridges made with bath solution, were calculated with the utility in pCLAMP, confirmed using a 3 M KCl electrode,44 and corrected before data analysis. To display changes throughout the course of an experiment, most summarized results are shown as the instantaneous slope conductance at the reversal potential (G_{Clswell}), which was obtained as follows. The whole cell current was fit with a mono-exponential function (using the Clampfit utility), and exported to Origin software (ver 7.0; OriginLab, Northampton,
MA), where the derivative at the reversal potential was calculated, normalized to the cell membrane capacitance, and plotted as a function of time after obtaining the whole-cell configuration.

**Recording solutions.** Unless otherwise indicated, chemicals were from Sigma-Aldrich (Oakville, ON, Canada). The standard pipette solution (“Solution 1”) contained (in mM): 50 NMDG-Cl, 70 NMDG-aspartate, 1 CaCl₂, 1 MgCl₂, 10 HEPES, 2 MgATP (pH 7.2; 280 mOsm/kgH₂O), with internal Ca²⁺ buffered to -20 nM using 10 EGTA. For experiments using a low-ionic strength pipette solution (“Solution 2”), the NMDG-aspartate was replaced with an iso-osmolar concentration of sucrose. For ionic strength calculations

\[
I = \frac{1}{2} \sum C_{i} z_{i}^{2}
\]

where \( z \) and \( C \) represent the valence and concentration of each ion in the solution. Before recording, the cells on coverslips were rinsed in standard bath solution, and then mounted in a perfusion chamber (Model RC-25, Warner Instruments, Hamden, CT). All bath solutions were pH 7.4. When required, bath solutions were rapidly exchanged using a gravity perfusion system flowing at 1.5–2 ml/min. Whole-cell recordings were established with iso-osmotic bath solutions (300–310 mOsm/kgH₂O), as follows. The standard bath solution (“Solution 3”) contained (in mM) 125 NaCl, 5 KCl, 1 CaCl₂, 1 MgCl₂, 10 HEPES and 5 D-glucose. Then, to minimize cation currents, Na⁺ and K⁺ were replaced with the bulky cation, NMDG⁺; such that the bath contained 140 NMDG-Cl, 1 CaCl₂, 1 MgCl₂, 10 HEPES and 5 D-glucose (“Solution 4”). To activate the Cl⁻ current, two hypo-osmotic bath solutions were made by diluting Solution 4 with a solution containing only 1 CaCl₂, 1 MgCl₂, 10 HEPES and 5 D-glucose (28 mOsm/kgH₂O). The resulting “Solution 5” (177 mOsm/kgH₂O) was used for electrophysiology, and “Solution 6” (205 mOsm/kgH₂O) was used for studying swelling-induced glutamate release. For anion selectivity studies only, the hypo-osmotic bath solutions contained 120 mM of the Na⁺ salt of the test anion (Cl⁻, I⁻, Br⁻, glutamate). Osmolarities were measured with a freezing point depression osmometer (Model 3MO, Advanced Instruments, Norwood, MA).

**Channel blockers and nucleotides.** Unless otherwise indicated, chemicals were from Sigma-Aldrich (Oakville, ON Canada). For experiments testing guanine nucleotides, immediately before use 200 μM GTPγS or GDPβS was added to the pipette solution from a frozen 20 mM nucleotide stock solution. Block of the swelling-activated current was investigated by adding known or putative Cl⁻ channel blockers of different chemical structures. Stock solutions (concentrations indicated) were stored at -20°C until used. The following compounds were dissolved in DMSO: the fenamates, NPPB (5-nitro-2-(3-phenylpropylamino) benzoic acid) (250 mM) and flufenamic acid (300 mM); glibenclamide (500 mM), a well known CFTR blocker; tamoxifen (10 mM); DCPIB (20 mM, Tocris Bioscience, MO USA), said to be a selective blocker of the volume-sensitive anion channel in cardiovascular tissues; and riluzole (300 mM), best known as a glutamate release inhibitor. Indanyllalkanoic acid (IAA-94, 300 mM) was dissolved in ethanol. Bumetanide (10 mM), an inhibitor of the Na⁺/K⁺/Cl⁻ symporter, was dissolved in double-distilled water.

For each blocker, the percent inhibition was calculated as

\[
\left(1 - \frac{G_{\text{rem}}}{G_{\text{Clmax}}} \right) \times 100
\]

where \( G_{\text{Clmax}} \) is the maximum conductance without a channel blocker, and \( G_{\text{rem}} \) is the remaining conductance in the presence of the blocker. \( G_{\text{Clmax}} \) was determined by fitting the plot of \( G_{\text{Clrev}} \) versus-time with a sigmoidal curve using the Boltzmann distribution, chosen because it consistently yielded low chi-squared values. Blocker data are presented as percent inhibition of \( G_{\text{Clrev}} \) unless otherwise stated.

**Measuring glutamate release.** Glutamate release from microglia was monitored with the two step Amplex® Red Glutamic Acid Assay Kit (Invitrogen, Burlington, ON), according to the manufacturer’s instructions. Step 1: L-glutamic acid is oxidized by glutamate oxidase to produce α-ketoglutarate, NH₃ and hydrogen peroxide (H₂O₂). H₂O₂ production is amplified by multiple cycles of the initial reaction in which L-Alanine and L-glutamate-pyruvate transaminase regenerate L-glutamic acid by transamination of α-ketoglutarate. Step 2: A highly fluorescent product, resorufin, is then generated in a 1:1 stoichiometry when H₂O₂ reacts with 10-acetyl-3,7-dihydroxy phenoxazine (Amplex® Red reagent) in a reaction catalyzed by horseradish peroxidase. Thus, resorufin fluorescence changes are proportional to the initial L-glutamic acid concentration in the cell supernatant. Because resorufin absorption and fluorescence emission maxima are about 571 nm and 585 nm, respectively, there is minimal contamination from cellular autofluorescence.

Microglia or MLS-9 cells growing in 12-well plates were washed with iso-osmotic bath solution, and 1 ml of a test solution was added. Iso-osmotic saline (Solution 3) was used for controls, and swelling experiments used hypo-osmotic saline (Solution 6), with or without a pharmacological compound. After 1 hr incubation at 37°C, 50 μl of each supernatant was transferred to a well in a 96-well plate. The resultant fluorescence emission was read at 590 nm on a Wallac 1420 VICTOR3™ plate reader (PerkinElmer, Woodbridge, ON, Canada). Microglia can produce H₂O₂, which would confound the assay; therefore, to assess this, the Step 2 reaction was conducted alone; i.e., 10-acetyl-3,7-dihydroxy phenoxazine was added to the cell supernatant. We found that this background H₂O₂ production, as well as the fluorescence from the medium, drugs and solvents, was very low; nevertheless, they were subtracted from the experimental measurements.

**Statistics.** Data from both electrophysiological and glutamate assays are expressed as mean ± SEM. Either Student’s t-tests (for single comparisons), or one-way ANOVA followed by Tukey’s test for multiple comparisons were conducted using Origin ver7.0 software (OriginLab, Northampton, MA). p < 0.05 was taken as statistically significant.

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