Relationships Between Calcium and pH in the Regulation of the Slow Afterhyperpolarization in Cultured Rat Hippocampal Neurons

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Submitted 2 December 2005; accepted in final form 19 July 2006

Kelly, Tony and John Church. Relationships between calcium and pH in the regulation of the slow afterhyperpolarization in cultured rat hippocampal neurons. J Neurophysiol 96: 2342–2353, 2006. First published August 2, 2006; doi:10.1152/jn.01269.2005. The Ca2+-dependent slow afterhyperpolarization (AHP) is an important determinant of neuronal excitability. Although it is established that modest changes in extracellular pH (pHo) modulate the slow AHP, the relative contributions of changes in the priming Ca2+ signal and intracellular pH (pHi) to this effect remain poorly defined. To gain a better understanding of the modulation of the slow AHP by changes in pHi, we performed simultaneous recordings of intracellular free calcium concentration ([Ca2+]), pHi, and the slow AHP in cultured rat hippocampal neurons coloaded with the Ca2+- and pH-sensitive fluorophores fura-2 and SNARF-5F, respectively, and whole cell patch-clamped using the perforated patch technique. Decreasing pHi from 7.2 to 6.5 lowered pHi, reduced the magnitude of depolarization-evoked [Ca2+], transients, and inhibited the subsequent slow AHP; opposite effects were observed when pHi was increased from 7.2 to 7.5. Although decreases and increases in pHi (at a constant pHo) reduced and augmented, respectively, the slow AHP in the absence of marked changes in preceding [Ca2+], transients, the inhibition of the slow AHP by decreases in pHi was correlated with low pHo-dependent reductions in [Ca2+], transients rather than the decreases in pHi that accompanied the decreases in pHi. In contrast, high pHi-induced increases in the slow AHP were correlated with the accompanying increases in pHi rather than high pHo-dependent increases in [Ca2+], transients. The results indicate that changes in pHi modulate the slow AHP in a manner that depends on the direction of the pHi change and substantiate a role for changes in pHi in modulating the slow AHP during changes in pHo.

INTRODUCTION

In many neurons of the CNS, trains or bursts of action potentials are followed by a prolonged slow afterhyperpolarization (AHP) that, in turn, is an important determinant of subsequent activity (reviewed by Storm 1990; Vogalis et al. 2003). Although the molecular correlate of the apamin-insensitive Ca2+-activated K+ current that underlies the slow AHP (sAHP) remains unknown, it is apparent that it can be modulated by a large number of neurotransmitters and second-messenger systems (reviewed by Vogalis et al. 2003; see also Stocker 2004). The slow AHP and sAHP are also sensitive to changes in extracellular pH (pHo); decreases and increases in pHi inhibit and augment, respectively, sAHP and the slow AHP, with consequent effects on neuronal excitability (Church 1999; Church and McLennan 1989; Kelly and Church 2004). Nevertheless, the relative contributions of changes in the priming Ca2+ signal and intracellular pH (pHi) to the modulation of sAHP and the slow AHP by changes in pHi remain unclear. Changes in pHi, acting directly or indirectly by changes in pHo (Church et al. 1998; Tombaugh and Somjen 1996, 1997), could modulate the Ca2+ influx through high-voltage–activated (HVA) Ca2+ channels constituting the primary source of Ca2+ for the activation of the slow AHP in hippocampal neurons (see Shah and Haylett 2000 and references therein). Alternatively, as shown previously for BK-type Ca2+-activated K+ channels that contribute to the fast AHP that follows a single action potential in hippocampal neurons (Church et al. 1998; see also Kume et al. 1990; Laurido et al. 1991), changes in pHi consequent on changes in pHo could affect directly the activities of the Ca2+-activated K+ channels that underlie the slow AHP.

To distinguish between these possibilities, in the present study we developed a technique in which whole cell perforated patch-clamp recordings of the slow AHP in cultured rat hippocampal neurons were obtained simultaneously with microspectrofluorimetric measurements of both intracellular free calcium concentration ([Ca2+]), and pH. The results indicate that the slow AHP is sensitive to changes in pHi and suggest that increases in pHi make a major contribution to high pHi-induced increases in the potential. In contrast, a pHi-dependent reduction in Ca2+ influx appears to be the major determinant of the inhibition of the slow AHP observed at low pHo.

METHODS

Cell preparation

All procedures conformed to guidelines established by the Canadian Council on Animal Care and were approved by The University of British Columbia Animal Care Committee.

Primary cultures of hippocampal neurons were prepared from 2- to 4-day-old postnatal Wistar rats (Animal Care Centre, University of British Columbia). Rat pups were anesthetized with 3% halothane in air and decapitated. Brains were removed rapidly and collected in ice-cold Leibovitz L-15 medium (Invitrogen Canada, Burlington, Canada) supplemented with 34 mM glucose (L-15/G). Hippocampi were removed, collected in ice-cold L-15/G, and then incubated for 15 min at 37°C in L-15/G medium containing 1 mg/ml papain (from papaya latex; Sigma-Aldrich Canada, Oakville, Canada) and 25 μg/ml DNAse (type II from bovine pancreas; Sigma-Aldrich Canada). Afterward, the L-15/G medium was discarded and replaced with Dulbecco’s modified Eagle medium F-12 (DMEM/F-12; Invitrogen Canada) supplemented with 29 mM NaHCO3 and 10% fetal bovine serum (FBS; Sigma-Aldrich Canada) (pH 7.4 at 37°C after equilibration with

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5% CO₂). Hippocampi were then mechanically dissociated using fire-polished Pasteur pipettes of decreasing tip diameters and the resulting cell suspension was plated at a density of 5–8 × 10⁵ neurons/cm² onto 15-mm-diameter glass coverslips coated with poly-d-lysine (100 μg/ml; Sigma-Aldrich Canada) and laminin (16.7 μg/ml; Sigma-Aldrich Canada). Neurons were allowed to adhere to substrate for 2 h before coverslips were transferred into 12 well plates containing DMEM/F-12 supplemented with 29 mM NaHCO₃ and substrate for 2 h before coverslips were transferred into 12 well plates containing DMEM/F-12 supplemented with 29 mM NaHCO₃ and 10% FBS. After 24 h, the growth medium was fully changed to Neurobasal Medium A (Invitrogen Canada) supplemented with B-27 Supplement (Invitrogen Canada), 0.5 mM glutamine (Invitrogen Canada), 50–100 U/ml penicillin (Sigma-Aldrich Canada), and 50–100 μg/ml streptomycin (Sigma-Aldrich Canada). Cultures were fed every 3–4 days by half-changing the existing medium with fresh supplemented Neurobasal Medium A. Gliial proliferation was inhibited 48 h after initial plating by adding 10 μM cytosine-β-D-arabinofuranoside hydrochloride (Sigma-Aldrich Canada). Each coverslip consisted primarily of hippocampal neurons with a maximum of 15% cells being glial. Neurons were used 7–14 days after plating.

**Solutions and chemicals**

The standard bath solution contained (in mM) NaCl 135, KCl 3, NaHCO₃ 21, MgCl₂ 1.5, CaCl₂ 4, d-glucose 10, and HEPES 5 (to increase extracellular buffering capacity and maintain a stable pHₗ); pH was 7.2 after equilibration with 5% CO₂-95% air at 30°C. In initial experiments, the bath solution contained 2 mM, rather than 4 mM, CaCl₂; although the effects of increases in pHₗ on the magnitudes of depolarization-evoked [Ca²⁺], transients and the subsequent slow AHPs were not significantly different at 2 versus 4 mM external Ca²⁺ (see RESULTS), the absolute magnitudes of depolarization-evoked [Ca²⁺] transients (50 ± 5 mV, n = 5) and the subsequent slow AHPs (1.7 ± 0.1 mV, n = 5) at 2 mM external Ca²⁺ under our experimental conditions (i.e., cultured neurons loaded with fura-2 and SNARF-5F) at pHₗ 7.2 were small, which effectively precluded the accurate assessment of the inhibitory effects of reductions in pHₗ on the parameters. Therefore we used the approach previously taken by others in hippocampal neurons (e.g., Knöpfel et al. 1990; Segal and Barker 1986) to increase the Ca²⁺ load by increasing external Ca²⁺ to 4 mM and conducted all subsequent experiments under these conditions. Low and high pH solutions contained 3 and 39 mM NaHCO₃ (pH 6.5 and 7.5, respectively, after equilibration with 5% CO₂-95% air); changes in [NaHCO₃] were balanced by equimolar changes in HCO₃⁻/CO₂-free medium, HEPES (10 mM) and NaCl isosmotically replaced NaHCO₃ and the solution was titrated with 10 M NaOH to pH 7.2 at 30°C. For Ca²⁺-free medium, CaCl₂ was omitted, [Mg²⁺] was increased to 5.5 mM, and 200 μM EGTA was added. D-(−)-2-Amino-5-phosphonopentanoic acid (d-AP5; 40 μM) and 6-cyano-7-nitroquinoxaline-2,3-dione (CNQX; 20 μM) were present in all bath solutions. Neurons were continuously superfused at a rate of 2 ml/min and all experiments were performed at 30°C.

Salts and experimental compounds, applied by superfusion, were obtained from Sigma-Aldrich Canada, with the exceptions of potassium methylsulfate (ICN Pharmaceuticals Canada, Montreal, Canada), the AM forms of fura-2 and seminaphthorhodafluor-5F 5-(and-6)-carboxylic acid (SNARF-5F; Molecular Probes, Eugene, OR), d-AP5 and CNQX (Tocris Bioscience, Ellisville, MO), and UCL 2027 (a generous gift from Drs. C. R. Ganelin and D. G. Haylett, Departments of Chemistry and Pharmacology, respectively, University College London, London, UK).

**Microspectrofluorimetry**

Concurrent measurements of [Ca²⁺], and pH were performed using the ratiometric indicators fura-2 and SNARF-5F, respectively. As detailed by Martínez-Zaguilán et al. (1996b), the excitation and emission characteristics of fura-2 and SNARF derivatives are sufficiently distinct to permit the accurate discrimination of [Ca²⁺], and pH-dependent signals from dual dye-loaded cells (see also Sheldon et al. 2004a). Neurons were incubated with 0.5–10 μM (see RESULTS) fura-2-AM for 30 min at 32°C in the presence of 0.04% Pluronic F-127; 10 μM SNARF-5F-AM was added during the final 10 min of incubation. After loading, coverslips were placed in standard medium for 20 min to ensure deesterification of the fluorophores and then mounted in a temperature-controlled perfusion chamber to form the base of the chamber.

Measurements of bulk cytosolic [Ca²⁺], and pH were performed at the somatic level using the dual-excitation and dual-emission ratio methods, respectively, using an imaging system (Atto Bioscience, Rockville, MD) in conjunction with an Axiovert 135 epifluorescence microscope (Carl Zeiss Canada, Don Mills, ON) equipped with two intensified charge-coupled device cameras (Atto Bioscience). A detailed description of the optical equipment used was previously presented (Sheldon et al. 2004a). In brief, as illustrated in Fig. 1, fura-2- and SNARF-5F-derived ratio pairs were collected continuously by alternating between the dual-excitation and dual-emission modes; each automated cycle took about 1.5 s to complete, including a <0.5-s delay between collecting fura-2- and SNARF-5F-derived ratio pairs, and was repeated every 2–15 s (typically 10 s) during the course of an experiment; during the recording of depolarization-evoked [Ca²⁺], transients, the acquisition of pH data was interrupted and fura-2-derived ratio pairs only were collected every 0.6–0.8 s.

A one-point calibration technique was used to convert background-corrected SNARF-5F–derived emission intensity ratio values (B₅₅₀/₆₄₀) into pH values, as described (Baxter and Church 1996; Sheldon et al. 2004a). In brief, neurons loaded with SNARF-5F and fura-2 were exposed at the end of an experiment to a pH 7.00 high-[K⁺] solution containing 10 μM nigericin and the resulting background-corrected ratio value at pH 7.00 in a given neuron was used as a normalization factor for the B₅₅₀/₆₄₀ ratio values obtained from that neuron during that experiment. SNARF-5F–derived B₅₅₀/₆₄₀ ratio values obtained during the one-point calibration procedure were not significantly different in patch-clamped (1.77 ± 0.04 ratio units; n = 6 neurons) versus neighboring intact (i.e., not patch-clamped) neurons on the same coverslip (1.69 ± 0.05 ratio units; n = 6 neuronal populations). The parameters required to convert experimentally derived B₅₅₀/₆₄₀ ratio values into pH values were determined in full calibration experiments in which neurons were exposed to 10 μM nigericin-containing high-[K⁺] media titrated to pH 5.5–8.5 in 0.5 pH unit increments (see Sheldon et al. 2004a); no differences in SNARF-5F calibration parameters were observed between neurons loaded with SNARF-5F alone or coloaded with SNARF-5F and fura-2 (data not shown; see also Sheldon et al. 2004a).

The parameters required to convert experimentally derived background-corrected fura-2–derived emission intensity ratio values (B₅₅₀/₆₄₀) into [Ca²⁺], values were determined in full in situ calibration experiments, as described in detail by Martínez-Zaguilán et al. (1991, 1996a,b). In brief, neurons loaded with fura-2 and SNARF-5F were exposed to K⁺-EGTA/Ca²⁺-EGTA buffers containing 10 μM 4-BrA23187 and 10 μM nigericin at three different pH values (pH 6.5–7.5 in 0.5 pH unit increments) with variable free Ca²⁺ concentrations (range 0–500 μM). Data from these experiments were used to obtain the calibration parameters that describe the effects of...
pH on the affinity of fura-2 for Ca\(^{2+}\) (see Martínez-Zaguilán et al. 1991, 1996a,b). Subsequently, the steady-state pH\(_i\), value measured immediately before an evoked [Ca\(^{2+}\)], transient was used to correct for the effect of pH on the \(K_d, R_{\text{min}},\) and \(R_{\text{max}}\) of fura-2, and [Ca\(^{2+}\)], was calculated from the equation

\[
[\text{Ca}^{2+}] = K_{d, \text{pH corr}} (R - R_{\text{min, pH corr}})/(R_{\text{max, pH corr}} - R)
\]

where \(K_{d, \text{pH corr}}\) is the pH-corrected \(K_d\) of fura-2 for Ca\(^{2+}\), \(R\) is the experimentally derived background-corrected fura-2 ratio value, and \(R_{\text{min, pH corr}}\) and \(R_{\text{max, pH corr}}\) are the minimum and maximum ratio values, respectively, corrected for pH (Martínez-Zaguilán et al. 1991, 1996a,b; see also Church et al. 1998).

To limit potential cross-contamination by nigericin, perfusion lines were replaced and the imaging chamber was decontaminated after each experiment by soaking first in ethanol, then in 20% Decon 75. To limit potential cross-contamination by nigericin, perfusion lines were replaced and the imaging chamber was decontaminated after each experiment by soaking first in ethanol, then in 20% Decon 75.

**Electrophysiology**

Because conventional whole cell recordings are associated with a rapid and marked rundown of the slow AHP in cultured hippocampal neurons, we used the amphotericin B perforated patch clamp technique described by Shah and Haylett (2000) to measure membrane potential \(V_m\) and the slow AHP in neurons loaded with fura-2 and SNARF-5F. Patch pipettes were pulled from 1.2 mm OD \(\times\) 0.9 mm ID borosilicate tubing (World Precision Instruments, Sarasota, FL). The first 100- to 200-\(\mu\)m section of the pipette tip was filled with a standard solution containing (in mM) KMMeSO\(_4\) 145, KCl 10, and HEPES 10 (titrated to pH 7.4 with 6 mM KOH) and then backfilled with the same solution but containing 1.2 mg/ml amphotericin B; final osmolality was about 290 mOsm/kg H\(_2\)O and open pipette resistance when filled was 2–5 M\(\Omega\). The reference bath electrode was a 3 M KCl, 4% agar bridge. After a seal \(> 1 \text{ G}\Omega\) was achieved, recordings were made (Axoclamp 2 or Axopatch 200B, Axon Instruments, Union City, CA) when access resistance was stable at \(< 50 \text{ M}\Omega\). Current and voltage waveforms were low-pass filtered at 3 kHz and digitized at 5–10 kHz using a Digidata 1322A controlled by pCLAMP software (v8, Axon Instruments).

During current-clamp recordings, a train of action potentials evoked by 4-ms suprathreshold depolarizing current pulses applied at 33 or 50 Hz (Master-8, A.M.P.I., Jerusalem, Israel) was used to generate a [Ca\(^{2+}\)], transient and the subsequent slow AHP; the number of action potentials in the train was held the same under control and test conditions. Under voltage-clamp conditions, a 80- to 200-ms depolarizing voltage step from the holding potential (−50 mV) to 0–20 mV was used to elicit a [Ca\(^{2+}\)], transient and the subsequent \(s_{\text{AHP}}\); leakage currents (estimated using 10-mV, 100-ms hyperpolarizing voltage steps from −50 mV) were subtracted off-line from all records of \(s_{\text{AHP}}\).

The criteria for detecting whether perforated patch recordings broke through included a change in the measured access resistance, an increase in the leakage current (which was estimated in all experiments), a sudden loss of fura-2 and SNARF-5F fluorescence from the patched cell, and, finally, a rapid rundown of the slow AHP itself (see Shah and Haylett 2000).
Data analysis

The change in pH_{2} evoked by a test maneuver was quantified as the difference between the steady-state pH_{2} value observed under the test condition with the mean of the steady-state pH_{2} values observed just before and, whenever possible, after full recovery from the test condition. In any given patch-clamped neuron, trains of 13 action potentials (see RESULTS) elicited [Ca^{2+}], transients of consistent amplitude, which were quantified as the difference between the B_{334}/B_{380} ratio value (or [Ca^{2+}]) measured immediately before the train of action potentials and the peak B_{334}/B_{380} ratio value (or peak [Ca^{2+}]) observed during the transient. In light of the relatively slow rates at which fura-2–derived ratio pairs were acquired, the computed means of two to five [Ca^{2+}], transients obtained before, during, and, whenever possible, after exposure to a test solution were used to quantify the effects of a test maneuver on the amplitudes of depolarization-evoked [Ca^{2+}], transients.

Experimentally induced changes in the slow AHP were quantified as the difference between the computed mean of the peak amplitudes of the slow AHPs evoked under a test condition with the mean of the slow AHP amplitudes evoked before and, whenever possible, after recovery from the test condition. If a distinct peak in the slow AHP was observed under control conditions, the amplitudes of the slow AHPs under test/wash conditions were measured at the same time interval after the end of the train of action potentials. If a distinct peak in the slow AHP was not evident under control conditions, slow AHP amplitude was measured 700 ms after the end of the train of action potentials (i.e., at a time point at which the slow AHP is usually near its peak and the medium AHP, if present, has decayed by >90%; see Kelly and Church 2004; Shah and Haylett 2000); measurements of the slow AHP under test/wash conditions were then made at the same time interval. Test measurements were conducted at the original control membrane potential by passing, when necessary, steady current through the recording electrode. Experimentally induced changes in S_{ahp} were quantified by comparing the peak amplitudes of the current obtained under control/wash and test conditions (see Kelly and Church 2004).

Data are presented as means ± SE, with the accompanying n value referring to either the number of patch-clamped neurons (each on a different coverslip) or, for neighboring intact (i.e., not patch-clamped) neurons on the same coverslip, the number of coverslips (i.e., neuronal populations) from which data were obtained. Data were analyzed in pCLAMP v.8 or Origin v.7 (OriginLab, Northampton, MA). Unless otherwise noted, statistical comparisons were performed using Student’s two-tailed t-test, paired or unpaired as appropriate. In all cases, statistical significance was assumed at the 5% level.

RESULTS

Fura-2 loading

Fura-2, a BAPTA derivative, chelates Ca^{2+} and thereby could modulate the Ca^{2+}-dependent slow AHP (e.g., Abel et al. 2004; Helmchen et al. 1996; Lancaster and Batchelor 2000; Lasser-Ross et al. 1997). Initially, therefore, we examined the effects of different concentrations of fura-2-AM in the loading medium on depolarization-evoked [Ca^{2+}], transients and the incidence of the slow AHP under our experimental conditions.

Under control pH_{2}, 7.2 HCO_{3}/CO_{2}-buffered conditions in the absence of fluorophores, resting V_{m} was ~60 ± 1 mV, input resistance (R_{m}) was 488 ± 11 MΩ, and a train of 13 action potentials (see following text) was followed by a slow AHP in 11/20 neurons (Table 1). A medium AHP was seen in only a minority (4/11) of cells that exhibited a slow AHP (see also Shah and Haylett 2000) and was not further analyzed. Voltage-clamp recordings performed on five neurons that exhibited a slow AHP under current-clamp conditions revealed that a 80- to 200-ms depolarizing step from ~50 to 0–20 mV elicited an S_{ahp} ≥40 pA in only 40% of these neurons (currents <40 pA at pH_{2} 7.2 were considered too small for reliable analysis under our experimental conditions). Although the overall incidence of S_{ahp} observed here is lower than that reported by Shah and Haylett (2000), who found that ~60% of their cultured hippocampal neurons exhibited a S_{ahp} >20 pA, it is consistent with the findings of Alger et al. (1994) in the same cell type. In light of these observations, the majority of subsequent experiments examined the slow AHP under current-clamp conditions.

In 12 of 16 neurons loaded with 2–10 μM fura-2-AM, a train of 13 action potentials elicited a transient increase in the fura-2–derived B_{334}/B_{380} ratio value; the remaining cells failed to exhibit an increase in [Ca^{2+}]. Of those cells that displayed a [Ca^{2+}], transient, only 25% exhibited a slow AHP, the peak amplitude of which was significantly smaller than that

<table>
<thead>
<tr>
<th>Loading Condition</th>
<th>n</th>
<th>Incidence, %</th>
<th>Peak amplitude, B_{334}/B_{380} ratio units</th>
<th>Slow AHP</th>
<th>Incidence, %</th>
<th>Peak amplitude, mV</th>
</tr>
</thead>
<tbody>
<tr>
<td>No fluorophore</td>
<td>20</td>
<td>n.a.</td>
<td>n.a.</td>
<td>55</td>
<td>5.3 ± 0.2</td>
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<tr>
<td>Fura-2-AM</td>
<td></td>
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<tr>
<td>2–10 μM</td>
<td>16</td>
<td>75</td>
<td>0.08 ± 0.01</td>
<td>25</td>
<td>1.5 ± 0.5</td>
<td></td>
</tr>
<tr>
<td>0.5–1 μM</td>
<td>14</td>
<td>100</td>
<td>0.18 ± 0.01b</td>
<td>64</td>
<td>5.1 ± 0.3c</td>
<td></td>
</tr>
<tr>
<td>Fura-2-AM (0.5–1 μM) + SNARF-5F-AM (10 μM)</td>
<td>72</td>
<td>100</td>
<td>0.19 ± 0.01b,c,d</td>
<td>58</td>
<td>4.7 ± 0.3b,c,d</td>
<td></td>
</tr>
</tbody>
</table>

Values are means ± SE; n = total number of perforated patch-clamped neurons examined under each experimental condition. In all cases, a train of 13 action potentials was used to evoke a [Ca^{2+}], transient and subsequent slow AHP. Concentrations of fluorophores indicated are those in the loading medium. Incidence of [Ca^{2+}], transient is the percentage of the total number of fura-2–loaded neurons examined that displayed a [Ca^{2+}], transient. Incidence of slow AHP is either the percentage of neurons not loaded with a fluorophore that displayed a slow AHP or the percentage of those fura-2–loaded cells that displayed a [Ca^{2+}], transient and a subsequent slow AHP. *P < 0.01 compared with the respective value obtained in the absence of fura-2. **P < 0.01 compared with the respective values obtained in cells loaded with 2–10 μM fura-2. *P > 0.7 compared with the respective value obtained in the absence of fluorophores. 4P > 0.7 compared with the respective values obtained in cells loaded with 0.5–1 μM fura-2 in the absence of SNARF-5F, n.a., not applicable.
observed in the absence of fura-2 (Table 1). In contrast, in each of 14 neurons loaded with 0.5–1 μM fura-2-AM, the same stimulus elicited an increase in \( B_{334}/B_{380} \) ratio values that was significantly larger than observed in neurons loaded with ≥2 μM fura-2-AM, and in 64% of these cells the \([Ca^{2+}]_i\) transient was followed by a slow AHP that was not significantly different from that observed in the absence of fura-2 (Table 1). Finally, in 72 cells loaded with ≥1 μM fura-2-AM, additional loading with 10–μM SNARF-5F-AM failed to significantly affect the amplitude of depolarization-evoked \([Ca^{2+}]_i\) transients or the incidence and amplitude of the subsequent slow AHP (Table 1), which remained stable for the duration of the recordings (typically 30–40 min). Consequently, subsequent experiments were performed in neurons coloaded with 10 μM SNARF-5F-AM and ≥1 μM fura-2-AM.

Simultaneous measurements of \( V_m \), \([Ca^{2+}]_i \), and \( pHi \)

In 42 neurons loaded with 10 μM SNARF-5F-AM and ≥1 μM fura-2-AM, resting \( V_m \) and \( R_h \) under control \( pHi \) 7.2 HCO\(_3\)/CO\(_2\)-buffered conditions were \(-59 ± 0.3 \) mV and \( 474 ± 18 \) MΩ, respectively (\( P > 0.6 \) in each case, compared with respective values obtained in the absence of fluorophores; see above). Resting \( pHi \) in perforated patch-clamped cells (\( pHi \), 7.19 ± 0.01; \( n = 42 \) neurons) was not significantly different (\( P > 0.3 \)) from that measured in neighboring intact (i.e., not patch-clamped) neurons on the same coverslips (\( pHi \), 7.23 ± 0.01; \( n = 42 \) coverslips) and was consistent with values previously reported in intact cultured fetal (Baxter and Church 1996) and acutely isolated adult (Beversee et al. 1996; Brett et al. 2002) rat hippocampal neurons in the presence of HCO\(_3\). Also as reported previously for intact hippocampal neurons under HCO\(_3\)/CO\(_2\)-buffered conditions (Baxter and Church 1996; Beversee et al. 1996; Brett et al. 2002; Smith et al. 1998), the distributions of resting \( pHi \) values in both perforated patch-clamped and intact neurons were unimodal, albeit the distribution was slightly skewed in patch-clamped cells (Fig. 2A). As illustrated in Fig. 2B, resting fura-2–derived \( B_{334}/B_{380} \) ratio values were significantly (\( P < 0.05 \)) higher in patch-clamped neurons (0.32 ± 0.01 ratio units; \( n = 42 \) neurons) than in neighboring intact neurons on the same coverslips (0.26 ± 0.01 ratio units; \( n = 42 \) coverslips), although both values were within the range observed previously in intact cultured hippocampal neurons with our recording system (e.g., Church et al. 1994, 1998).

Next, given the relative paucity of studies in which the slow AHP has been examined in neurons in primary culture (see Sah and Clements 1999; Shah and Haylett 2000), the amplitudes of depolarization-evoked \([Ca^{2+}]_i \), transients and the subsequent slow AHPs were significantly reduced under external Ca\(^{2+}\)-free conditions (not shown) or by the application of 200 μM Ca\(^{2+}\) (Fig. 3). In contrast, 10 μM isoproterenol and 5–10 μM UCL 2027 (a relatively selective inhibitor of the slow AHP in rat hippocampal neurons; Shah et al. 2001) significantly reduced the slow AHP but not the preceding \([Ca^{2+}]_i \) transient (Fig. 3).

Effects of a decrease in \( pHo \)

Under control \( pHo \) 7.2 HCO\(_3\)/CO\(_2\)-buffered conditions, resting \( pHi \) in intact and perforated patch-clamped neurons loaded with 10 μM SNARF-5F and ≥1 μM fura-2 was \( 7.30 ± 0.02 \) (\( n = 8 \) coverslips) and \( 7.31 ± 0.02 \) (\( n = 8 \) neurons; Fig. 5A), respectively. In the patch-clamped neurons, resting \([Ca^{2+}]_i \) was \( 70 ± 5 \) nM and a train of 13 action potentials elicited a \( 120 ± 8 \) nM increase in \([Ca^{2+}]_i \); the amplitude of the subsequent slow AHP was \( 4.4 ± 0.2 \) mV (Fig. 5A). Reducing \( pHo \) from 7.2 to 6.5 produced significant (\( P > 0.01 \) in both cases) reductions in \( pHi \) in both patch-clamped cells (to \( pHi \), 7.10 ± 0.03; Fig. 5A) and the neighboring intact cells on the same
The reductions in pH_i and the percentage reductions in the peak amplitudes of the [Ca^{2+}][i] transients and slow AHPs from the effects of exposure to pHo 6.5 were then compared with the respective values obtained at pHo 7.2 (Fig. 5A). Recovery of pH_i and the amplitudes of the [Ca^{2+}][i] transients and slow AHPs from the effects of exposure to pH 6.7 medium was slow and usually incomplete. In one cell in which a distinct s_{AHP} was discernable, the peak amplitude of s_{AHP} declined by 73%, from 95 pA at pHo 7.2 to 25 pA at pHo 6.5.

The reductions in pH_i and the percentage reductions in the peak amplitudes of the [Ca^{2+}][i] transients observed in patch-clamped cells on reducing pH_o from 7.2 to 6.5 were then plotted against the percentage reductions in the peak amplitudes of the slow AHPs measured in the same cells (Fig. 5B). The percentage reduction in the slow AHP at pH_i 6.5 was significantly correlated with the percentage reduction in the magnitude of the depolarization-evoked [Ca^{2+}][i] transient but not with the reduction in pH_i ($P < 0.01$ and $P = 0.17$, respectively). The lack of a significant correlation between the reductions in the slow AHP and pH_i was maintained when the reduction in pH_i was plotted as an increase in [H^+]_i or a percentage increase in [H^+]_i (not shown; see Fig. 8D).

Effects of an increase in pH_o

Under pH_o 7.2 HCO_3^-/CO_2-buffered conditions, resting pH_i in perforated patch-clamped and intact, unpatched neighboring neurons on the same coverslips was 7.22 ± 0.01 ($n = 16$ neurons; Fig. 6A) and 7.24 ± 0.02 ($n = 16$ coverslips), respectively ($P > 0.2$ in both cases, compared with the values observed in the preceding experimental series also at pH_o 7.2). In the patch-clamped neurons, resting [Ca^{2+}][i], was 80 ± 2 nM and a train of 13 action potentials elicited a 94 ± 3 nM increase in [Ca^{2+}][i], and a subsequent slow AHP, the peak amplitude of which was 3.3 ± 0.1 mV (Fig. 6A). Increasing pH_i from 7.2 to 7.5 significantly ($P < 0.01$ in both cases) increased pH_i in patch-clamped and intact neurons to 7.34 ± 0.01 (Fig. 6A) and 7.35 ± 0.02 pH units, respectively. In the patch-clamped neurons, high pH_o conditions failed to affect resting [Ca^{2+}][i]; (see also Church et al. 1998) but significantly ($P < 0.01$ in both cases) increased the peak amplitudes of the [Ca^{2+}][i] transient to 130 ± 4 nM (a 36 ± 1% increase) and the subsequent slow AHP to 5.3 ± 0.2 mV (a 69 ± 2% increase) (Fig. 6A). Similar results were obtained at 2 mM external Ca^{2+}, where increasing pH_o from 7.2 to 7.5 increased the amplitude of the [Ca^{2+}][i] transient evoked by 13 action potentials by 32 ± 2% ($n = 3$) and the amplitude of the subsequent slow AHP by 85 ± 4% ($P > 0.02$ in each case, compared with the increases observed at 4 mM [Ca^{2+}][i]). In addition, similar to the slow AHP and s_{AHP} observed during conventional whole cell recordings in rat CA1 neurons in slices (Kelly and Church 2004), the augmented
were reached when the increase in pH was plotted as a
change in the peak amplitudes of \([Ca^{2+}]_i\) transients obtained in the absence vs. presence of the indicated compound. **P \(< 0.01\) rather than the increase in the peak amplitude of the slow AHP observed on increasing pHo from 7.2 to 7.5 was 5.5 
from 7.2 to 7.5 in individual perforated patch-clamped neurons. Under control pHo 7.2 HCO3
in the absence and presence of apamin, respectively; (top) that were followed by slow AHPs (bottom). In this and subsequent figures, the [Ca2+], transients used to elicit the slow AHPs shown are indicated by +. Application of 5 \(\mu\)M UCL 2027 (A) or 10 \(\mu\)M isoproterenol (B) reduced the slow AHP without appreciably affecting the preceding [Ca2+], transient. Scale bars in B apply to the respective traces in A. C: summary of the mean \pm SE percentage changes in the peak amplitudes of [Ca2+], transients and the subsequent slow AHPs observed under the conditions indicated on the figure, compared with the peak amplitudes obtained under control conditions. n \(\geq 3\) in all cases. *P \(< 0.05\) and **P \(< 0.01\) for the difference between the corresponding measurement obtained in the absence vs. presence of the indicated compound.

The percentage increases in the peak amplitudes of [Ca2+], transients and the increases in pHo observed on raising pHo from 7.2 to 7.5 in individual perforated patch-clamped neurons were then plotted against the percentage increases in the peak amplitudes of the slow AHPs measured in the same cells (Fig. 6B). In contrast to results obtained on reducing pHo to 6.5, the high pHo-induced increase in the peak amplitude of the slow AHP was significantly correlated with the increase in pHo (P < 0.01) rather than the increase in the peak amplitude of the preceding [Ca2+], transient (P = 0.26). Similar conclusions were reached when the increase in pHo was plotted as a
decline in [H+]i or a percentage decrease in [H+]i (not shown; see Fig. 8D).

Effects of changing pHj at a constant pHo

Next, we examined whether changes in pHj at a constant pHo affect the magnitudes of depolarization-evoked [Ca2+], transients and the subsequent slow AHPs. Although externally applied weak acids (e.g., propionic acid, butyric acid) and weak bases (e.g., NH3, trimethylamine) are widely used to

\(\begin{align*}
\text{FIG. 4. Pharmacological characterization of depolarization-evoked [Ca2+], transients and subsequent slow AHPs.} A \text{ and B: simultaneous [Ca2+], and perforated patch-clamp recordings from 2 different cultured hippocampal neurons. Under control pHo 7.2 HCO3/CO2-buffered conditions, trains of 13 action potentials (applied at a) elicited [Ca2+], transients (top) that were followed by slow AHPs (bottom). In this and subsequent figures, the [Ca2+], transients used to elicit the slow AHPs shown are indicated by +. Application of 5 \(\mu\)M UCL 2027 (A) or 10 \(\mu\)M isoproterenol (B) reduced the slow AHP without appreciably affecting the preceding [Ca2+], transient. Scale bars in B apply to the respective traces in A. C: summary of the mean \pm SE percentage changes in the peak amplitudes of [Ca2+], transients and the subsequent slow AHPs observed under the conditions indicated on the figure, compared with the peak amplitudes obtained under control conditions. n \(\geq 3\) in all cases. *P \(< 0.05\) and **P \(< 0.01\) for the difference between the corresponding measurement obtained in the absence vs. presence of the indicated compound.

\(\begin{align*}
\text{FIG. 5. Effects of a decrease in pHj on pHo, [Ca2+], transients, and the slow AHP.} A: \text{simultaneous measurements of pHj (top), [Ca2+], (middle), and Vm (bottom) from a perforated patch-clamped hippocampal neuron under HCO3/CO2-buffered conditions. Decreasing pHj from 7.2 to 6.5 reduced pHo. Peak amplitudes of [Ca2+], transients (evoked by trains of 13 action potentials applied at a) and the subsequent slow AHPs were reduced at pHj 6.5 compared with pHj 7.2. Breaks in the pHj record represent 3-min pauses in the acquisition of pHj measurements to record the [Ca2+], transients shown in the middle panels. B: percentage decreases in the peak amplitudes of [Ca2+], transients (a) and decreases in pHj (b) observed in individual neurons on reducing pHj from 7.2 to 6.5, plotted against percentage decreases in the peak amplitudes of the slow AHPs observed in the same cells. Low pHj-induced decrease in the [Ca2+], transient, but not the low pHj-induced decrease in pHj, was significantly correlated with the inhibition of the slow AHP (P < 0.01 and P = 0.17, respectively, as determined by a t-test of the Pearson product-moment correlation coefficient for each data set). Solid line is a linear least-squares regression fit to the data points (a) relating the percentage decrease in the slow AHP to the percentage decrease in the [Ca2+], transient (r^2 = 0.97).
\end{align*}\)
mechanisms such that, in neurons with a low resting pH, in the absence of HCO\textsubscript{3}\textsuperscript{-}, Na\textsuperscript{+}-dependent Cl\textsuperscript{-}/HCO\textsubscript{3}\textsuperscript{-} exchange causes pH\textsubscript{i} to rise, whereas in neurons with a high resting pH, in the absence of HCO\textsubscript{3}\textsuperscript{-}, Na\textsuperscript{+}-independent Cl\textsuperscript{-}/HCO\textsubscript{3}\textsuperscript{-} exchange causes pH\textsubscript{i} to fall.

Consistent with the findings of Brett et al. (2002) in intact hippocampal neurons, switching from a HCO\textsubscript{3}\textsuperscript{-}-free to a HCO\textsubscript{3}\textsuperscript{-}-containing medium (pH\textsubscript{o} constant at 7.2) caused pH\textsubscript{i} to decrease in 11 of 13 perforated patch-clamped neurons with high (>7.14) initial pH\textsubscript{i} values in HCO\textsubscript{3}\textsuperscript{-}-free medium (of the remaining neurons, one showed no change and the other a small increase in pH\textsubscript{i}) and to increase in three of three neurons with low (≤7.14) initial pH\textsubscript{i} values in HCO\textsubscript{3}\textsuperscript{-}-free medium (Fig. 7). The addition of HCO\textsubscript{3}\textsuperscript{-} failed to significantly affect resting [Ca\textsuperscript{2+}], which was 86 ± 3 nM in the absence and 90 ± 3 nM in the presence of HCO\textsubscript{3}\textsuperscript{-} (P > 0.8; see also Ou-Yang et al. 1994a,b).

As illustrated in Fig. 8, A and B, changes in pH\textsubscript{i} at a constant pH\textsubscript{o} produced only minor changes in the peak amplitudes of depolarization-evoked [Ca\textsuperscript{2+}], transients (see also Church et al. 1998) but consistently altered the peak amplitudes of the subsequent slow AHPs. When the percentage changes in the peak amplitudes of [Ca\textsuperscript{2+}], transients and the changes in pH\textsubscript{i} observed in individual neurons on the addition of HCO\textsubscript{3} (pH\textsubscript{o} constant at 7.2) were plotted against the percentage changes in the peak amplitudes of the slow AHPs measured in the same cells (Fig. 8C), the percentage change in the slow AHP was significantly (P < 0.01) correlated with the change in pH\textsubscript{i} but not with the change in the peak amplitude of the preceding [Ca\textsuperscript{2+}], transient (P = 0.59). The same results were obtained

change pH\textsubscript{i} at a constant pH\textsubscript{o}, at above ambient temperature they produce only transient changes in pH\textsubscript{i} in hippocampal neurons (Bonnet et al. 2000; Church et al. 1998). In addition, NH\textsubscript{3} and trimethylamine directly inhibit the slow AHP and sI\textsubscript{sh}, in rat hippocampal neurons (Kelly and Church 2004, 2005). In light of these considerations, we used the transition from a nominally HCO\textsubscript{3}\textsuperscript{-}/CO\textsubscript{2}-free HEPES-buffered medium to a HCO\textsubscript{3}\textsuperscript{-}/CO\textsubscript{2}-buffered medium (pH\textsubscript{o} constant at 7.2) to change pH\textsubscript{i} at a constant pH\textsubscript{o}. As described by Brett et al. (2002) (see also Beverse et al. 1996; Smith et al. 1998), the addition of HCO\textsubscript{3}\textsuperscript{-} activates HCO\textsubscript{3}\textsuperscript{-}-dependent pH\textsubscript{i}-regulating

![Image](http://jn.physiology.org/)

FIG. 6. Effects of an increase in pH\textsubscript{o} on pH\textsubscript{i} [Ca\textsuperscript{2+}], transients, and the slow AHP. A: simultaneous measurements of pH\textsubscript{i} (top), [Ca\textsuperscript{2+}], (middle), and V\textsubscript{m} (bottom) from a perforated patch-clamped hippocampal neuron under HCO\textsubscript{3}\textsuperscript{-}/CO\textsubscript{2}-buffered conditions. Increasing pH\textsubscript{o} from 7.2 to 7.5 increased pH\textsubscript{i}. Peak amplitudes of [Ca\textsuperscript{2+}] transients (evoked by trains of 13 action potentials applied at A) and the subsequent slow AHPs were increased at pH\textsubscript{o} 7.5 compared with pH\textsubscript{o} 7.2. Breaks in the pH\textsubscript{i} record represent 2-min (at pH\textsubscript{o} 7.2) and 3-min (at pH\textsubscript{o} 7.5) pauses in the acquisition of pH\textsubscript{i} measurements to record the [Ca\textsuperscript{2+}] transients shown in the middle panels. B: percentage increases in the peak amplitudes of [Ca\textsuperscript{2+}] transients (circles) and increases in pH\textsubscript{i} (squares) observed in individual neurons on increasing pH\textsubscript{o} from 7.2 to 7.5, plotted against percentage increases in the peak amplitudes of the slow AHPs observed in the same cells. High pH\textsubscript{i}-induced augmentation of the slow AHP was significantly correlated with the increase in pH\textsubscript{i}, but not the increase in the [Ca\textsuperscript{2+}] transient (P < 0.01 and P = 0.26, respectively, as determined by a t-test of the Pearson product-moment correlation coefficient for each data set). Solid line is a linear least-squares regression fit to the data points (circles) relating the percentage increase in the slow AHP to the increase in pH\textsubscript{i} (r\textsuperscript{2} = 0.97).

FIG. 7. Changes in pH\textsubscript{i} evoked by the transition from a HCO\textsubscript{3}\textsuperscript{-}-free to a HCO\textsubscript{3}\textsuperscript{-}-containing medium at a constant pH\textsubscript{o}. Changes in steady-state pH\textsubscript{i} observed in patch-clamped (circles) and neighboring intact neurons on the same coverslips (squares) on the transition from a pH 7.2 HCO\textsubscript{3}\textsuperscript{-}-free medium to a pH 7.2 HCO\textsubscript{3}\textsuperscript{-}-containing medium are plotted against initial pH\textsubscript{i} values in HCO\textsubscript{3}\textsuperscript{-}-free medium. Separate linear least-squares regression fits to the data obtained from patch-clamped and intact neurons were not significantly different (P > 0.05) and all data points were best fit (r\textsuperscript{2} = 0.83) by a single linear least-squares regression fit (solid line) that had a negative slope and an x-intercept at pH\textsubscript{i} 7.14.

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when the change in pH was plotted as a change in [H$^+$]$_i$ (not shown) or as a percentage change in [H$^+$]$_i$ (Fig. 8D). A distinct $s_{\text{mp}}$ was discernable in one neuron in this experimental series; in this cell, pH$_i$ decreased by 0.10 pH units on the addition of HCO$_3^-$ (pH$_o$ constant at 7.2) and the peak amplitude of $s_{\text{mp}}$ declined by 25%, from 51 pA in the absence of HCO$_3^-$ to 38 pA in the presence of HCO$_3^-$.

**Discussion**

We developed a technique to measure [Ca$^{2+}$]$_i$, pH$_i$, and $V_m$ simultaneously in cultured hippocampal neurons and used it to substantiate and extend previous findings, made on the basis of electrophysiological recordings from CA1 pyramidal neurons in slices (Church 1992, 1999; Kelly and McLennan 1989; Kelly and Church 2004), that the slow AHP is modulated by changes in pH$_i$ at a constant pH$_o$ (Kelly and Church 2004), that the slow AHP is modulated by changes in pH$_i$ at a constant pH$_o$ (Church 1992, 1999; Church and McLennan 1989; Zaguilán et al. 1991) are relatively commonplace, concurrent measurements of all three parameters have for the most part been limited to larger cells (e.g., invertebrate neurons) impaled with ion-sensitive microelectrodes (ISMs) (but see Silver and Erecinska 1990). The technique described here provides a means to measure [Ca$^{2+}$]$_i$, pH$_i$, and $V_m$ simultaneously in small cells that are not readily amenable to stable impalements with ISMs and offers a means to better understand the relationships between cytosolic [Ca$^{2+}$] and [H$^+$] and the roles of both ions in the regulation of cellular excitability. The validity of the technique was attested to by the facts that resting pH$_i$ values, the distribution of resting pH$_i$ values, and the magnitudes of the changes in pH$_i$ evoked by changes in pH$_o$ or the addition of external HCO$_3^-$ in dual dye-loaded patch-clamped cells were in agreement with those obtained in previous studies where pH$_i$ alone was measured (e.g., Brett et al. 2002; Sheldon et al. 2004a; Smith et al. 1998). In addition, resting [Ca$^{2+}$]$_i$ values and the amplitudes of [Ca$^{2+}$]$_i$ transients evoked under control (pH$_o$ 7.2) conditions in perforated patch-clamped neurons were significantly correlated with the changes in pH$_i$, but not with the changes in the preceding depolarization-evoked [Ca$^{2+}$]$_i$ transients (P < 0.01 and P = 0.59, respectively, as determined by a t-test of the Pearson product-moment correlation coefficient for each data set). Solid line is a linear least-squares regression fit to the data points ($	extcircled{c}$) relating the percentage change in pH$_i$ to the peak amplitude of slow AHPs in the same cells. Changes in the slow AHPs were significantly correlated with the changes in pH$_i$, but not with the changes in the slow AHP to the change in pH$_i$ ($r^2 = 0.91$). D: like C, but with change in pH$_i$ plotted as percentage change in [H$^+$]. Solid line is a linear least-squares regression fit to the data points ($	extcircled{c}$) relating the percentage change in the slow AHP to the percentage change in [H$^+$] ($r^2 = 0.88$).

**Simultaneous measurements of $V_m$, [Ca$^{2+}$]$_i$, and pH$_i$**

Although simultaneous measurements of $V_m$ and [Ca$^{2+}$]$_i$ (e.g., Abel et al. 2004; Sah and Clements 1999), $V_m$ and pH$_i$ (e.g., Trapp et al. 1996; Willoughby and Schwiening 2002), and pH$_i$ and [Ca$^{2+}$]$_i$ (e.g., Austin et al. 2004; Sah and Clements 1999), the data points ($	extcircled{c}$) relating the percentage change in [Ca$^{2+}$]$_i$ transient to the percentage change in slow AHPs observed in the same cells. Changes in the slow AHPs were significantly correlated with the changes in pH$_i$, but not with the changes in the slow AHP to the change in pH$_i$ ($r^2 = 0.91$). D: like C, but with change in pH$_i$ plotted as percentage change in [H$^+$]. Solid line is a linear least-squares regression fit to the data points ($	extcircled{c}$) relating the percentage change in the slow AHP to the percentage change in [H$^+$] ($r^2 = 0.88$).
loaded with fura-2 and SNARF-5F corresponded well with those measured previously with fura-2 alone under similar stimulating (trains of action potentials, as opposed to depolarizing voltage steps) and recording (bulk cytosolic measurements at the soma) conditions (e.g., Abel et al. 2004; Church et al. 1994, 1998; Knöpfel et al. 1990; Lancaster and Batchelor 2000; Lee et al. 2005). Finally, not only were $V_{m}$ and $R_{m}$ in dual dye-loaded neurons patch-clamped using the perforated patch-clamp technique similar to values measured in unloaded cells but also the pharmacological and other characteristics of the slow AHP in dual dye-loaded neurons were comparable to those observed previously by ourselves (Kelly and Church 2004) and others (e.g., Shah and Haylett 2000; Shah et al. 2001) in the absence of fluorophore(s), provided that fura-2-AM was loaded at $\leq 1 \, \mu M$.

**Effects of changes in pH**

Although changes in pH$_i$ are known to affect HVA Ca$^{2+}$ currents in rat hippocampal neurons (Tombaugh and Somjen 1997), in the present study changes in pH$_i$ at a constant pH$_o$ modulated the slow AHP in the absence of marked changes in the priming Ca$^{2+}$ signal. This finding is in agreement with previous conventional sharp microelectrode and whole cell recordings in CA1 neurons in slices (Church 1999; Kelly and Church 2004; see also Church et al. 1998), where reductions in the slow AHP and $s_{ahp}$ evoked by decreases in pH$_i$ at a constant pH$_o$ occurred in the absence of significant changes in Ca$^{2+}$-dependent depolarizing potentials or $I_{ca}$ and were significantly attenuated when internal buffering power was raised by the inclusion of high concentrations of H$^+$ buffers in the recording electrode. Also in support of pH$_i$ being an important modulator of the slow AHP is the observation that high pH$_i$-induced increases in the potential were correlated with the accompanying increases in pH$_i$ but not the accompanying increases in [Ca$^{2+}$], transients. This result is also entirely consistent with previous conventional whole cell recordings from CA1 neurons in slices (Kelly and Church 2004), where in addition it was found that the effects of high pH$_i$ to augment the slow AHP and $s_{ahp}$ were significantly attenuated by increasing internal H$^+$ buffering capacity in the absence of any change in the preceding Ca$^{2+}$ potentials.

The apparent dissociation between the priming Ca$^{2+}$ signal and the magnitude of the slow AHP when pH$_i$ is changed at a constant pH$_o$ could reflect the effects of changes in pH$_i$ on the slow AHP and, possibly, other K$^+$ conductances (see Church et al. 1998) that, in turn, would act to offset any direct effect of changes in pH$_i$ on Ca$^{2+}$ influx, resulting in little net effect on the magnitude of [Ca$^{2+}$]$_i$ transients. Alternatively, the relatively small changes in pH$_i$ used in the present experiments may have been insufficient to appreciably affect the activities of the L- and N-type HVA Ca$^{2+}$ channels in large part mediate depolarization-evoked increases in [Ca$^{2+}$]$_i$, and the subsequent activation of the slow AHP in rat hippocampal neurons (see Borde et al. 2000; Church et al. 1994, 1998; Kelly and Church 2004; Shah and Haylett 2000; Tanabe et al. 1998). Importantly, L- and N-type Ca$^{2+}$ channels in rat hippocampal neurons exhibit similar sensitivities to changes in pH$_i$ and pH$_o$ (Church et al. 1998; Tombaugh and Somjen 1996, 1997), indicating that the lack of correlation between the amplitude of depolarization-evoked [Ca$^{2+}$]$_i$ transients and the slow AHP when pH$_o$ is increased or when pH$_i$ is changed at a constant pH$_o$ is unlikely to reflect the possibility that changes in pH might be affecting Ca$^{2+}$ entry through a Ca$^{2+}$ channel subtype that does not contribute to the activation of the slow AHP.

In agreement with previous reports (e.g., Church 1999; Church et al. 1998; Ou-Yang et al. 1994a), decreasing pH$_o$ from 7.2 to 6.5 decreased pH$_i$, reduced the magnitude of depolarization-evoked [Ca$^{2+}$]$_i$ transients, and inhibited the subsequent slow AHPs. In contrast to results obtained at pH$_o$ 7.5, however, the decrease in the slow AHP at pH$_o$ 6.5 was correlated with a low pH$_i$-dependent decrease in the priming Ca$^{2+}$ signal rather than a low pH$_i$-induced decrease in pH$_i$. This observation parallels previous findings in conventional whole cell patch-clamped CA1 neurons in slices (Kelly and Church 2004), where low pH$_i$-induced reductions in the slow AHP and $s_{ahp}$ were accompanied by decreases in depolarization-evoked Ca$^{2+}$-dependent potentials and $I_{ca}$ and were not significantly affected by increasing internal H$^+$ buffering capacity.

The differences between the mechanisms by which reductions and increases in pH$_i$ modulate the slow AHP may be explained by a number of factors. For example, the marked decrease in [Ca$^{2+}$]$_i$ transients observed at pH$_i$ 6.5, which is consistent with the pK$_s$ for the effects of pH$_o$ on L- and N-type HVA Ca$^{2+}$ currents and depolarization-evoked [Ca$^{2+}$]$_i$ transients in rat hippocampal neurons (pH$_i$ 7.1–7.2; Church et al. 1998; Tombaugh and Somjen 1996), may have reduced the slow AHP to such an extent that modest decreases in pH$_i$ consequent on decreases in pH$_o$ failed to exert an additional inhibitory effect. In contrast, the augmented slow AHP at pH$_o$ 7.5 occurred despite the likelihood that the underlying channels were already saturated with Ca$^{2+}$ (see Abel et al. 2004; Gerlach et al. 2004; Shah and Haylett 2000). Although this is consistent with the possibility that internal protons modulate the slow AHP by an allosteric site on the channel complex (see Laurido et al. 1991), the possibility remains that protons may compete with Ca$^{2+}$ ions at regulatory binding sites to modulate channel activity (see Church et al. 1998; Copello et al. 1991; Kume et al. 1990; Peitersen et al. 2006). Because microdomains of [Ca$^{2+}$]$^{2+}$/Ca$^{2+}$ and/or pH$_i$ in the immediate vicinity of the channels underlying the slow AHP may differ from values measured in bulk cytoplasm (e.g., Ro and Carson 2004; Vaughan-Jones et al. 2006; Willoughby and Schwiening 2002; Willoughby et al. 2005), simultaneous near-membrane [Ca$^{2+}$], and pH$_i$ measurements may help shed further light on the relationships between Ca$^{2+}$ and H$^+$ in the regulation of the slow AHP. Nevertheless, it must be noted that the mechanism(s) whereby Ca$^{2+}$ activates the channels underlying the slow AHP remain unknown (cf. BK- and SK-type Ca$^{2+}$-activated K$^+$ channels; see Sah and Faber 2002; Stocker 2004; Vogalis et al. 2003) and, consistent with a role for a cytoplasmic intermediate between Ca$^{2+}$ and the gating of the channels underlying the slow AHP, activation of the slow AHP has been reported to require elevations in bulk cytosolic [Ca$^{2+}$] rather than at the membrane (Abel et al. 2004; see also Lasser-Ross et al. 1997; Lee et al. 2005). Kinetic studies at the single-channel level will be required to determine the precise mechanism(s) whereby protons interact with Ca$^{2+}$ ions to modulate the slow AHP.

We did not address the possibilities that changes in pH$_o$ might act directly on the channels underlying the slow AHP or...
that changes in pHo and/or pHi might affect processes downstream from Ca\(^{2+}\) influx that could potentially modulate the magnitude of the Ca\(^{2+}\) signal responsible for their activation. Nevertheless, changes in pHi fail to affect the unitary properties of BK-, IK-, or SK-type Ca\(^{2+}\)-activated K\(^+\) channels in a variety of cell types (Church et al. 1998; Jäger and Grissmer 2004; Kume et al. 1990; Pedersen et al. 2000) and, in the present study (see also Kelly and Church 2004), pH\(_i\)-induced changes in the slow AHP were never observed in the absence of parallel changes in the magnitude of [Ca\(^{2+}\)]\(_{\text{cyt}}\) transients. In addition, although changes in pHi can affect Ca\(^{2+}\) handling by intracellular stores, internal Ca\(^{2+}\) buffering, and the activities of [Ca\(^{2+}\)]\(_{\text{cyt}}\), extrusion mechanisms (e.g., Hoyt and Reynolds 1998; Ou-Yang et al. 1994b; Thomas 2002; Zucker 1981), the relatively modest changes in pHo, and/or pHi, used here (also Church et al. 1998) were not associated with marked changes in resting [Ca\(^{2+}\)]\(_{\text{cyt}}\) or the generation of depolarization-independent [Ca\(^{2+}\)]\(_{\text{cyt}}\) transients, and changing pH\(_i\) at a constant pHo failed to significantly affect the magnitude of depolarization-evoked [Ca\(^{2+}\)] transients.

Functional implications

In summary, simultaneous measurements of [Ca\(^{2+}\)], pHo, and V\(_{\text{m}}\) indicate that changes in pHo modulate the slow AHP in rat hippocampal neurons in a manner that depends on the direction of the pH\(_o\) change; moderate reductions in pHo inhibit the slow AHP primarily by reducing Ca\(^{2+}\) influx, whereas moderate increases in pHo augment the slow AHP primarily through an increase in pHi. In addition, changes in pHi at a constant pHo modulate the slow AHP independent from changes in the priming Ca\(^{2+}\) signal, further supporting a role for pHi in the regulation of the slow AHP.

The sensitivity of the slow AHP to the changes in pHo and pHi, used here, which are within the pathophysiological range seen in vivo (see Chesler 2003), may have a number of implications for neuronal function. Reductions in pH, for example, may contribute to the inhibition of the slow AHP observed during oxygen deprivation (Kulik et al. 2002) and, if pronounced, may in this way promote neuronal injury. Conversely, an increase in the slow AHP may help to limit the increases in neuronal excitability and epileptiform activity observed during increases in pH (Balestrino and Somjen 1988; Church and McLennan 1989; also Kelly and Church, unpublished observations), especially if Ca\(^{2+}\) influx is increased to such an extent that the underlying channels become saturated (see Canepari et al. 2000; Sinha et al. 1995). The present results also raise the possibility that activity-induced reductions in pHo\(_{\text{crit}}\) could limit the potential therapeutic effects of agents designed to enhance the slow AHP, and it will be important to assess whether such agents, like the neuronal SK channel enhancer EBIO (Petersen et al. 2006; see also Pedarzani et al. 2005), retain their ability to augment the slow AHP at the low pH values associated with pathological events such as ischemia.

GRANTS

This work was funded by a Grant-in-Aid from the Heart and Stroke Foundation of British Columbia and Yukon and an Operating Grant from the Canadian Institutes of Health Research.

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