Metrifonate Increases Neuronal Excitability in CA1 Pyramidal Neurons from Both Young and Aging Rabbit Hippocampus

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The effects of metrifonate, a second generation cholinesterase inhibitor, were examined on CA1 pyramidal neurons from hippocampal slices of young and aging rabbits using current-clamp, intracellular recording techniques. Bath perfusion of metrifonate (10–200 μM) dose-dependently decreased both postburst afterhyperpolarization (AHP) and spike frequency adaptation (accommodation) in neurons from young and aging rabbits (AHP: p < 0.002, young; p < 0.050, aging; accommodation: p < 0.024, young; p < 0.001, aging). These reductions were mediated by muscarinic cholinergic transmission, because they were blocked by addition of atropine (1 μM) to the perfusate. The effects of chronic metrifonate treatment (12 mg/kg for 3 weeks) on CA1 neurons of aging rabbits were also examined ex vivo. Neurons from aging rabbits chronically treated with metrifonate had significantly reduced spike frequency accommodation, compared with vehicle-treated rabbits. Chronic metrifonate treatment did not result in a desensitization to metrifonate ex vivo, because bath perfusion of metrifonate (50 μM) significantly decreased the AHP and accommodation in neurons from both chronically metrifonate- and vehicle-treated aging rabbits. We propose that the facilitating effect of chronic metrifonate treatment on acquisition of hippocampus-dependent tasks such as trace eyeblink conditioning by aging subjects may be caused by this increased excitability of CA1 pyramidal neurons.

Key words: afterhyperpolarization; aging; atropine; carbachol; cholinesterase inhibitor; eserine; hippocampal slice; metrifonate; rabbits; spike frequency adaptation

The learning and cognitive deficits observed in normal aging and in Alzheimer’s disease (AD) patients are hypothesized to be partly caused by the loss of cholinergic neurons in the basal forebrain. It has been suggested that these deficits may be alleviated by improving cholinergic function (Bartus et al., 1982; Hal-lak and Giacobini, 1989). Currently, tacrine and donepezil (United States) and exelon and galantamine (Europe) are clinically approved cholinesterase inhibitors (ChEI) used for treating AD. Further research is ongoing to develop more effective ChEIs with fewer side effects (Cummings et al., 1998; Morris et al., 1998; Pettigrew et al., 1998).

One new generation ChEI currently in phase III clinical trials is metrifonate, an organophosphate compound that is considered to be a prodrug, because it is transformed nonenzymatically to 0.0-dimethyl 2,2-dichlorovinyl phosphate, which produces the long-lasting inhibition of both acetylcholinesterase (AChE) and butyrylcholinesterase (Nordgren et al., 1978; Schmidt et al., 1998). More importantly, extended treatment with metrifonate has been shown to result in an increased acetylcholine (ACh) level with fewer, less severe side effects than other ChEIs (Soin-innen et al., 1990; Becker et al., 1991; Sihver et al., 1997). Behavioral experiments have demonstrated that metrifonate treatment improved cognitive performance in AD patients (Cummings et al., 1998; Morris et al., 1998; Pettigrew et al., 1998), reduced both scopolamine- and basal forebrain lesion-induced deficits in water maze and passive avoidance tasks in rats (Itoh et al., 1997), and rescued object recognition in aging rats (Scali et al., 1997). Recently, our laboratory demonstrated that metrifonate treatment facilitated acquisition of trace (hippocampus-dependent) eyeblink conditioning in aging rabbits (Kronforst-Collins et al., 1997a,b).

Although there are numerous behavioral experimental reports concerning metrifonate, there is no electrophysiological literature concerning the effects of metrifonate, which may be relevant to the mechanism of actions mediating its potentially valuable therapeutic benefits. Several in vitro experiments have demonstrated that application of ACh, muscarinic agonists, or anticholinesterases increased neuronal excitability [reduced postburst afterhyperpolarization (AHP) and spike frequency adaptation (accommodation)] of hippocampal pyramidal neurons (Bernardo and Prince, 1981, 1982; Cole and Nicoll, 1983, 1984a,b; Madison and Nicoll, 1984; Halliwell, 1990; Taylor and Griffith, 1993; Pedarzani and Storm, 1996). Furthermore, both the AHP and accommodation were reduced in CA1 neurons from young and aging rabbits that acquired eyeblink conditioning, but not in trained rabbits that did not learn (Disterhoft et al., 1986, 1988, 1996; Coulter et al., 1988; de Jonge et al., 1990; Moyer et al., 1996; Thompson et al., 1996b). Also, both the AHP and accommodation are greater in CA1 neurons from aging rabbits (Moyer et al., 1992) and rats (Landfield and Pitzer, 1984; Potier et al., 1992) as compared with that from young animals.

The current study was designed to determine (1) the effects, (2) the effective concentrations of bath application of metrifonate on CA1 neurons from hippocampal slices of young and aging rabbits,
(3) whether chronic metrifonate treatment in aging rabbits alters basal CA1 excitability ex vivo, and (4) if the chronic metrifonate treatment has a saturating, desensitizing effect.

MATERIALS AND METHODS

Subjects. Young (<3 month) and aging (>36 month) female New Zealand albino rabbits (Oryctolagus cuniculus) were used as subjects. We chose to study these two age groups for the following reasons: (1) previous work in our laboratory has established that rabbits, 30+ months old, are impaired in acquiring the trace eyelblink conditioning task (Thompson et al., 1997a,b), and similar impairments have been observed in aging humans (Woodruff-Pak and Thompson, 1988); (2) we have demonstrated that hippocampal pyramidal neurons from aging rabbits are less excitable than those from young animals and possibly contribute to the age-related learning deficits (see results; Moyer et al., 1992); (3) the effects of bath application of metrifonate in vivo have not been explored in either age group; and (4) previous work has demonstrated that metrifonate in vivo unequally increased the ACh levels in the hippocampus of young and aging subjects (Scali et al., 1997), thus, bath application of metrifonate in vitro may also yield unequal effects on the hippocampal slices from young and aging subjects. Each subject was housed in an individual cage in a climate-controlled room on a 12 hr light/dark cycle with ad libitum access to food and water. The animal care was provided and managed by the animal care personnel of Northwestern University after the guidelines established by the university and the United States Department of Agriculture.

Slice preparation. Hippocampal slices were made using procedures previously described (Moyer et al., 1996; Thompson et al., 1996b). The rabbits were anesthetized with halothane in a fume hood and killed by decapitation. The brain was quickly exposed, hemisected in situ, removed, and immersed in an ice-cold (<1°C) oxygenated sucrose–artificial CSF (aCSF) that minimizes anoxic impact during slice prepara-

Electrophysiological recording and data analysis. Intracellular recordings were made from CA1 pyramidal neurons using an Axoclamp 2A amplifier (Axon Instruments, Foster City, CA) and previously published protocols (Moyer et al., 1996; Thompson et al., 1996b). Microelectrodes were made from a thin-walled capillary glass and filled with 3 M KC1 (30–50 Ml). Slices were individually transferred to a submersion chamber and superfused (at 1.75 ml/min) with aCSF heated to 31°C. A cell was classified as a CA1 neuron and included in the study if it had little spontaneous activity at rest, a stable resting membrane potential less than −60 mV, an action potential duration >1.2 msec from rise threshold to recrossing the resting potential, an input resistance ≥20 Ml, and an action potential amplitude >80 mV from rest. After the neuron had stabilized for 5 min after the initial impalement, the biophysical properties were measured with the neuron held near −68 mV (using less than ±0.2 nA) to ensure that the differences observed were not caused by voltage-dependent membrane properties.

The baseline membrane properties were measured in normal aCSF. The current–voltage (I–V) relations were studied by using 400 msec current steps from −0.2 nA to +0.2 nA. The input resistance was calculated by measuring the plateau voltage deflection during the last 75 msec of a 400 msec, −0.2 nA hyperpolarizing step. The AHP was studied using a 100 msec depolarizing current step that reliably elicited a burst of four action potentials. The duration of the AHP was measured as the time required for the membrane potential to return to the baseline potential for at least 10 msec from the 100 msec depolarizing current step offset. The peak AHP amplitude was calculated as the maximum negative voltage deflection from the baseline potential during the first 250 msec after the current offset. The integrated area of the AHP was calculated from the current offset for the entire duration of the AHP. A total of five AHP measurements were made from each neuron at 30 sec intervals.

Accommodation was studied using an 800 msec depolarizing current step of the same stimulus intensity used to evoke the AHP. The number of action potentials elicited was noted for three trials at 30 sec intervals.

After the baseline measurements were recorded, the perfusate was changed to an aCSF containing carbachol (500–1000 nM), eserine (500–5000 nM), metrifonate (1–200 μM), atropine (1 μM; by itself or added to the previously mentioned drugs), or vehicle (normal aCSF). The experimenter was blind to the identity of the perfusate until the end of data collection. The neuron was held near −75 mV for 10 min and allowed to stabilize during the changing of the perfusate. After the 10 min interval, the biophysical measurements were repeated. In some cases, the neuron was subject to another perfusate, or a wash-out of the perfusate was attempted (blind procedures were used). At the end of the experiment, the resting membrane potential was determined as the difference in the membrane potential before and after the microelectrode withdrawal from the neuron. The slice was changed if a cell was lost during an experiment, if more than five penetrations were made, or after the completion of the experiment. All data were digitized and analyzed on-line using a Lab NB or NB-MIO-16H and DMA-2000 boards (National Instruments, Austin, TX) interfaced to Power Macintosh computers using software routines written in LabView (National Instruments). Analog-to-digital sampling rates were 10 kHz for I–V, AHP, and accommodation measurements and 1–2 kHz for the resting membrane potentials. Complete analyses were performed off-line using procedures developed with LabView. Statistical analyses were performed using paired t tests and ANOVA (StatView; Abacus Concepts, Berkeley, CA). Significant main effects were evaluated using Scheffe’s post hoc tests. All data are reported as the mean ± SEM.

Chronic treatment with metrifonate. Using similar procedures to those previously published (Kronforst-Collins et al., 1997b), aging (>40 month) rabbits received 15 oral doses (5 of treatment followed by 2 of no treatment repeated for three weeks) of either 12 mg/kg metrifonate dissolved in a 100 mM sodium citrate vehicle (n = 4; mean age, 41.31 ± 0.06 months) or vehicle alone (n = 3; mean age, 41.00 ± 0.13 months). Previously, 12 mg/kg metrifonate was found to produce optimum facilitation of trace eyelink conditioning in aging rabbits (Kronforst-Collins et al., 1997a,b). Blood samples were taken from all subjects 1 day before the start of treatment and −2 hr before killing. Twenty-four hours after the last treatment, hippocampal slices were prepared, and CA1 neurons were recorded from as described above. AHP, accommodation, and input resistance were measured sequentially as described above in aCSF, aCSF with 50 μM metrifonate, and aCSF with 50 μM metrifonate plus 1 μM atropine perfusates with 10 min intervals between changes in the perfusate. The experimenter was blind to the identity of the chronic treatment during the daily administration of drug sampling, electrophysiological recordings, and data reductions until the end of the experiment.

ChE inhibition measurement. The level of ChE inhibition was measured using procedures described by Kronforst-Collins et al. (1997a,b). The subjects were given fentanyl citrate and droperidol anesthesia (0.5 mg/kg, i.m.) before blood sampling. Each blood sample was collected in two 1.5 ml aliquot tubes each containing 50 μl of heparin. The samples were centrifuged at 1000 × g, 4°C for 15 min. The plasma and red blood cells (RBCs) were separated and stored at −80°C until ChE inhibition assays were performed. It has been previously demonstrated that the level of RBC AChE activity in the rabbit is significantly correlated with that of the ChE of the brain (Kronforst-Collins et al., 1997a). Thus, the mean percentages of ChE inhibition values were calculated for the RBCs for each subject. The ChE inhibition values were analyzed with ANOVA and unpaired t tests.

Drugs. Metrifonate was a gift from Bayer Corporation (West Haven, CT). All other drugs used were purchased from Sigma (St. Louis, MO). Eserine and atropine stocks were made and used in near darkness. Stock solution of metrifonate (pH = 4.0) was prepared weekly and refrigerated (−3°C) along with the other stock solutions.

RESULTS

Metrifonate decreased the AHP amplitude and area

Metrifonate significantly decreased the AHP peak amplitude and integrated area in CA1 neurons from both young and aging subjects (Figs. 1D, 2A; Table 1). Significant reductions of the AHP amplitude and area were produced with 10 μM metrifonate in the neurons from young rabbits (p < 0.006; p < 0.046, respec-
tively). No depression was observed in the neurons from aging rabbits at this concentration ($p_{H11022} = 0.309$; $p_{H11022} = 0.178$, respectively). Instead, a significant reduction of the AHP amplitude was observed with 50 μM metrifonate in the neurons from aging rabbits ($p_{H11021} = 0.009$); the AHP area was not significantly reduced, although a trend toward the reduction was observed ($p_{H11021} = 0.074$).

Decrements in both AHP peak amplitude and integrated area for the neurons from aging rabbits were observed with 100 μM metrifonate ($p_{H11021} = 0.002$; $p_{H11021} = 0.001$, respectively). The neurons from young rabbits depolarized to levels at which regular bursts of spontaneous action potentials made the biophysical measurements impossible at 100 μM metrifonate, and eventual cell death occurred in all but one of five neurons attempted. No such effects were observed in neurons from aging rabbits (tested up to 200 μM).

Metrifonate reduced the AHP peak amplitude in the neurons from both young and aging rabbits in a dose-dependent manner ($F_{(4,20)} = 6.048$, $p < 0.002$; $F_{(3,31)} = 2.920$, $p < 0.050$; respectively). Scheffe’s post hoc test revealed that the percent AHP peak amplitude reduction was significantly greater at both 50 and 100 μM metrifonate compared with 1 μM in the neurons from young rabbits ($p < 0.019$; $p < 0.026$, respectively). Also, ANOVA revealed that there was a significant difference in the mean baseline AHP peak amplitudes between the neurons from young (3.76 ± 0.33; $n = 25$) and aging (4.90 ± 0.30; $n = 37$) rabbits ($F_{(1,60)} = 6.29$; $p < 0.015$) (Moyer et al., 1992).

Metrifonate also shortened the duration of the AHP. However, this effect was significant only at 100 and 200 μM metrifonate for neurons from the aging rabbits ($p < 0.001$; $p < 0.025$, respectively).

**Metrifonate decreased spike frequency accommodation**

Metrifonate significantly decreased the accommodation of CA1 neurons from both naive young and aging subjects. In neurons from the young rabbits, significant accommodation decrement was observed at 50 μM metrifonate ($p < 0.001$). Lower concentrations of metrifonate were not effective; a higher concentration (100 μM tested) caused instability and cell death in neurons from young animals. In neurons from the aging rabbits,
significant reductions were observed at concentrations of 50 μM and higher metrifonate (Table 1). The accommodation decrement was dose-dependent in neurons from the aging rabbits (F(3,32) = 7.11; p < 0.001) with greater reduction observed at 200 μM compared with 10 and 50 μM metrifonate (p < 0.035; p < 0.013, respectively). A significant dose interaction was also observed for the neurons from young rabbits (F(4,20) = 3.57; p < 0.024). Finally, there was a significant difference of mean baseline accommodation measures between the neurons from young and aging rabbits (F(1,60) = 10.60; p < 0.002) (Moyer et al., 1992).

**Atropine partially reversed the effects of metrifonate**

Atropine, by itself, had no significant effect on the biophysical measurements (see Table 3). Atropine (1 μM) significantly reversed the action of metrifonate on accommodation (Fig. 1; see Fig. 4). In the neurons from young rabbits, when atropine was added to 50 μM metrifonate, the accommodation was returned to that observed during baseline (p < 0.028). However, atropine did not return the AHP measures back to baseline (p > 0.192).

**Chronic metrifonate treatment increased neuronal excitability**

CA1 neurons from chronically metrifonate-treated rabbits were more excitable than those from the vehicle-treated rabbits (Fig. 3). Typically, the neurons from aging rabbits exhibit a strong accommodation, as illustrated in Figure 1A (see also Fig. 3A, bottom panel; Table 1). The baseline accommodation measurement was significantly reduced in neurons from the metrifonate-

### Table 1. Biophysical properties of CA1 neurons from young and aging rabbits after bath application of metrifonate in vitro

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<th>Metrifonate (10 μM)</th>
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<td>Baseline</td>
<td>Postdrug</td>
<td>p value</td>
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<td>Young</td>
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<tr>
<td>Postburst AHP</td>
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<tr>
<td>Amplitude (mV)</td>
<td>−3.62 ± 0.68</td>
<td>−3.07 ± 0.72</td>
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<td>Area (mVsec)</td>
<td>−3.09 ± 1.31</td>
<td>−2.01 ± 0.98</td>
<td><strong>0.046</strong> (6)</td>
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<td>Duration (sec)</td>
<td>1.80 ± 0.54</td>
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<td>Accommodation</td>
<td>10.75 ± 1.64</td>
<td>11.62 ± 1.33</td>
<td>0.439 (6)</td>
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<td>Input resistance (MΩ)</td>
<td>43.04 ± 4.80</td>
<td>39.65 ± 5.09</td>
<td>0.299 (6)</td>
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Aging

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<tr>
<th></th>
<th>Metrifonate (50 μM)</th>
<th>Metrifonate (100 μM)</th>
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<td></td>
<td>Baseline</td>
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<td>Postburst AHP</td>
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<tr>
<td>Amplitude (mV)</td>
<td>−4.94 ± 0.47</td>
<td>−4.12 ± 0.46</td>
<td><strong>0.009</strong> (17)</td>
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<tr>
<td>Area (mVsec)</td>
<td>−5.43 ± 1.06</td>
<td>−4.16 ± 0.84</td>
<td>0.074 (17)</td>
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<tr>
<td>Duration (sec)</td>
<td>3.46 ± 0.34</td>
<td>3.34 ± 0.44</td>
<td>0.775 (17)</td>
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<tr>
<td>Accommodation</td>
<td>7.02 ± 0.56</td>
<td>8.81 ± 0.91</td>
<td><strong>0.014</strong> (17)</td>
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<td>Input resistance (MΩ)</td>
<td>43.53 ± 2.54</td>
<td>47.53 ± 2.31</td>
<td><strong>0.034</strong> (16)</td>
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The number in parentheses indicates the number of cells recorded. The data for young rabbits at 100 μM metrifonate is based on the one of five neurons that survived through the drug application. The numbers in bold type indicate statistical significance using paired t test. The measurements are the mean ± SEM.

**Figure 3.** Chronic, oral treatment with metrifonate (12 mg/kg daily) in aging subjects significantly reduced the accommodation in CA1 neurons. A depicts a typical example of the differing response to an 800 msec depolarizing current pulse used to obtain four action potentials in the first 100 msec observed in CA1 neurons from chronically metrifonate- (top) or vehicle-treated (bottom) subjects. B, Mean baseline accommodation was significantly reduced in neurons from the chronic metrifonate-treated subjects as compared with those from vehicle-treated subjects (mean ± SEM; **p < 0.01, unpaired t test).
treated aging rabbits ($F_{(1,45)} = 9.789; p < 0.003$). The decreased accommodation brought the baseline accommodation measurements of these neurons to values similar to those from young, untreated subjects; 8.76 ± 0.71 action potentials vs 8.73 ± 0.33 action potentials for the aging, treated and young, untreated rabbits, respectively ($F_{(1,102)} = 0.0019; p = 0.965$). The baseline AHP measurements were not significantly different between the metrifonate- and vehicle-treated aging rabbits ($F_{(1,45)} = 0.358; p = 0.553$).

The CA1 neurons from chronically metrifonate-treated aging rabbits were not desensitized to the effects of bath application of metrifonate ex vivo. Addition of metrifonate (50 μM) to the perfusate significantly decreased the AHP peak amplitude and accommodation in neurons from both chronically metrifonate- and vehicle-treated aging rabbits ($F_{(1,45)} = 0.358; p = 0.553$). The decrements were reversed with 1 μM atropine in the perfusate ($** p = 0.01; ** p = 0.003$).

**Figure 4.** Bath application of 50 μM metrifonate to CA1 neurons from chronic metrifonate-treated subjects further significantly decreased the accommodation ($B$). This effect was significantly reversed by adding 1 μM atropine to the perfusate ($C$) (same neuron in $A-C$). The mean increase (with 50 μM metrifonate) and decrease (with 50 μM metrifonate and 1 μM atropine) in the number of action potentials elicited in CA1 neurons from the chronic metrifonate-treated subjects are illustrated in $D$ (mean ± SEM; **$p < 0.01$; paired $t$ tests).

The CA1 neurons from chronically metrifonate-treated aging rabbits were not desensitized to the effects of bath application of metrifonate ex vivo. Addition of metrifonate (50 μM) to the perfusate significantly decreased the AHP peak amplitude and accommodation in neurons from both chronically metrifonate-treated aging rabbits ($p < 0.004; p < 0.009$, respectively) and vehicle-treated rabbits ($p < 0.012; p < 0.011$, respectively) (Figs. 4, 5; Table 2). The integrated area of the AHP was significantly reduced for the neurons from metrifonate-treated rabbits ($p < 0.033$). The decrements were reversed with 1 μM atropine in the perfusate ($p < 0.028$ for AHP; $p < 0.033$ for accommodation and $p < 0.033$ for area; Figs. 4, 5; Table 2).

**Chronic metrifonate treatment reduced cholinesterase activity**

The chronic metrifonate treatment significantly decreased the RBC AChE activity of aging rabbits by 25% (24.62 ± 3.62%; $n = 4; p < 0.021$); no inhibition of AChE activity was observed in the vehicle-treated, control subjects ($p > 0.241; n = 3$). The RBC AChE activity was significantly different between the two treatment groups ($F_{(1,5)} = 28.79; p = 0.003$).

**Carbachol increased neuronal excitability**

Bath application of carbachol significantly increased the excitability of CA1 neurons from both young and aging subjects, as reported previously (Bernardo and Prince, 1982; Cole and Nicoll, 1983, 1984a,b; Potier et al., 1992; Taylor and Griffith, 1993; Pedarzani and Storm, 1996). At 500 nM, the AHP amplitude was reduced by 1.0 mV in CA1 neurons from both young (1.18 ± 0.30 mV; $p = 0.003$) and aging (1.17 ± 0.27 mV; $p < 0.001$) subjects (Fig. 6, Table 3). The integrated area of the AHP was significantly reduced for CA1 neurons from both age groups ($p < 0.009$ for young; $p < 0.005$ for aging). The duration of the AHP was also significantly shortened for CA1 neurons from aging subjects ($p < 0.034$), and a trend for such a reduction was observed in young ($p > 0.106$).

Accommodation was significantly reduced in CA1 neurons from both age groups. In CA1 neurons from aging subjects, the number of action potentials elicited by a long depolarizing pulse was nearly doubled after carbachol (500 nM) application ($p = 0.0002$). The effect on accommodation in CA1 neurons of young subjects was not nearly as dramatic, although it was significantly reduced, ($p = 0.0010$; Fig. 6, Table 3). This may be caused by the fact that the baseline accommodation was stronger in CA1 neu-
metrifonate significantly decreases neuronal excitability in vitro; (2) chronic metrifonate treatment increases the basal level of neuronal excitability ex vivo; (3) neurons from chronically metrifonate-treated subjects remain sensitive to bath application of metrifonate, i.e., no "saturating" effect is observed with chronic treatment; and (4) the excitability changes observed with metrifonate mimic that observed with the cholinergic agonist carbachol and the cholinesterase inhibitor eserine, and could be reversed by atropine, thus metrifonate acts primarily via muscarinic cholinergic neurotransmission.

A higher concentration of metrifonate was necessary to significantly increase the neuronal excitability of CA1 neurons of aging rabbits compared with that of the young (Fig. 2, Table 1). This suggests an apparent shift in the efficacy of metrifonate for the neurons of aging rabbits. Likewise, a ceiling effect of metrifonate (tested up to 200 μM) was not observed in the neurons from aging rabbits. In the neurons from young rabbits, 100 μM metrifonate depolarized the neurons with high-frequency bursts of action potentials, leading to membrane potential instability and eventual cell death. However, no such effects were observed in the neurons from aging rabbits. If anything, a greater reduction of AHP and accommodation was observed with higher concentration of metrifonate in the neurons from aging rabbits. One possible contributing factor to these effects may be the reduced level of endogenous ACh in vivo in aging subjects (Scali et al., 1997; Vannucchi et al., 1997), which may translate into less endogenous ACh present in the tissue of aging subjects in vitro. Scali and colleagues (1997) report that 80 mg/kg of metrifonate given to young and aging rats produced unequal increase in ACh levels in the hippocampus in vivo: a threefold increase in the young compared with only a 30% increase in the aged. Thus, a higher dose of metrifonate may be needed to allow cholinergic reduction of the larger AHPs and stronger accommodations that are generally observed in CA1 neurons from aging subjects (see Results; Landfield and Pitler, 1984; Moyer et al., 1992).

Chronic metrifonate treatment caused a clear reduction of RBC AChE inhibition similar to the effect previously reported by Kronforst-Collins et al. (1997b). CA1 neurons from the metrifonate-treated rabbits were found to be more excitable, resembling neurons from the young rabbits. Furthermore, the AHP and accommodation of neurons from chronically metrifonate-treated rabbits remained sensitive to the bath application of metrifonate (Fig. 4). This observation is consistent with the results obtained by Hinz et al. (1998) that chronic metrifonate treatment did not (1) alter the ACh synthesis rate, (2) change the muscarinic or nicotinic receptor binding, or (3) affect the monoamines in the brain (Soininen et al., 1990). This is contrary to the effects of chronic treatment with tacrine, which increased dopaminergic and serotoninergic metabolism (Soininen et al., 1990) and decreased binding to high-affinity choline uptake and nicotinic and muscarinic receptors (Silver et al., 1997).

Numerous experiments have examined the modulatory role of cholinergic inputs in various learning and memory tasks, especially in aging subjects. For example, experiments with the central cholinergic blocker, scopolamine, demonstrated that humans (Solomon et al., 1993) and rabbits (Harvey et al., 1983) were impaired in learning delay (nonhippocampus-dependent) eyelink conditioning when pretreated with scopolamine. This impairment is hypothesized to be mediated by the hippocampus, because scopolamine treatment did not impair hippocampal-lesioned subjects from acquiring delay eyelink conditioning (Solomon et al., 1983), but did prevent rabbits from acquiring trace (hippocampus-dependent) eyelink conditioning (Kaneko and
As mentioned in Materials and Methods, the neurons were given 10 min of rest/stabilization periods during each change of the bath perfusate. They posit that the learning rate is increased and recall of novel associations have been emphasized by Hasselmo et al., 1989; Moyer et al., 1992; Kowalska and Disterhoft, 1994). This is presumably mediated through the neocortex and/or associated circuitry. Similar results were obtained with the L-type calcium channel blocker, nimodipine, providing further support for this neuronal excitability hypothesis. Nimodipine facilitated acquisition of trace eyeblink conditioning (Kim et al., 1995). Changes in neuronal excitability were observed immediately after acquisition, and the neuronal changes returned to that of the controls (presumed basal levels) within 7 d, whereas the behavioral performance remained asymptotic for months (Moyer et al., 1996; Thompson et al., 1996b). By that time memory storage was served with chronic metrifonate treatment in aging rabbits that acquire the trace eyeblink conditioning task, but not in the control animals or in animals that are trained but fail to acquire the task (Disterhoft et al., 1996; Moyer et al., 1996; Thompson et al., 1996b). Both are reduced in hippocampal neurons from rabbits that acquire the trace eyeblink conditioning task, but not in the control animals or in animals that are trained but fail to acquire the task (Disterhoft et al., 1996; Moyer et al., 1996; Thompson et al., 1996b). The AHP and accommodation changes also parallel the time course of hippocampal function. The hippocampus is required for the acquisition, but not long-term recall, of trace eyeblink conditioning (Kim et al., 1995). Changes in neuronal excitability were observed immediately after acquisition, and the neuronal changes returned to that of the controls (presumed basal levels) within 7 d, whereas the behavioral performance remained asymptotic for months (Moyer et al., 1996; Thompson et al., 1996b). By that time memory storage was presumably mediated through the neocortex and/or associated circuitry. Similar results were obtained with the L-type calcium channel blocker, nimodipine, providing further support for this neuronal excitability hypothesis. Nimodipine facilitated acquisition of trace eyeblink conditioning and increased neuronal excitability (reduced the AHP and accommodation) (Deyo et al., 1989; Moyer et al., 1992; Kowalska and Disterhoft, 1994). This is similar to the situation that we find here with chronic metrifonate treatment: enhanced acquisition rate of trace eyeblink conditioning, tightly coupled with increased CA1 excitability ex vivo.

The importance of cholinergic modulation of synaptic transmission in the hippocampus and neocortex during learning and recall of novel associations have been emphasized by Hasselmo et al. (1992), Hasselmo and Bower (1992), and Hasselmo and Schnell (1994). They posit that the learning rate is increased and

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<td>Vehicle</td>
</tr>
<tr>
<td>Metrifonate (50 μM)</td>
</tr>
<tr>
<td>Metrifonate (50 μM) + Atropine (1 μM)</td>
</tr>
</tbody>
</table>

The neurons were subjected to a bath application of metrifonate (50 μM) followed by metrifonate plus atropine (1 μM) after the initial baseline measurements were recorded. The measurements are the mean ± SEM.

Figure 6. Bath application of 500 nm carbachol significantly increased CA1 excitability in neurons from both young and aging subjects. The mean AHP peak amplitude was significantly reduced by 500 nm carbachol (striped bars) as compared with the baseline (open bars) measurements (A). The mean accommodation was significantly reduced by 500 nm carbachol (striped bars) as compared with the baseline (open bars) measurements (B) (mean ± SEM; **p < 0.01; ***p < 0.001; paired t tests).
the maintenance of memory is prolonged by cholinergic suppression of the intrinsic fiber synapses in the hippocampus and cortex during learning. They further postulate that there is a direct regulation of ACh concentration in the hippocampus: during learning of new associations, the cholinergic input is high; during recall of learned associations, the cholinergic input is low (Has-selmo and Schnell, 1994). Their hypotheses of the cholinergic modulation fits very well with what may be occurring in the hippocampus of rabbits during trace eyeblink conditioning. As the rabbits are initially learning to associate the tone with the airpuff (novel association), the cholinergic input into the hippocampus would be increased, thereby increasing the neuronal excitability of the pyramidal neurons (AHP and accommodation reductions). This boost of increased neuronal excitability with ACh may be crucial for aging rabbits that are usually impaired in learning the trace eyeblink conditioned response (Thompson et al., 1996a) to learn associative tasks: slow EPSPs and CA1 population EPSPs are significantly reduced in aging subjects compared with the young (Landfield et al., 1986; Barnes et al., 1992; Potier et al., 1992; Taylor and Griffith, 1993). Soon after acquiring the tone–airpuff association, the cholinergic input to the hippocampus would begin to decrease, but the neuronal excitability of the pyramidal neurons would remain increased for a period of time (Moyer et al., 1996; Thompson et al., 1996b). With the steady decline of the cholinergic input to the hippocampus, the neuronal excitability of the pyramidal neurons would return to the basal state. But the association would be maintained over time, because by that point the memory for the association would have been stored in the neocortex, as rabbits tested months after the training sessions perform asymptotically on trace eyeblink conditioning (Moyer et al., 1996; Thompson et al., 1996b). In the behavioral experiments conducted by Kronforst-Collins et al. (1997b), metrifonate treatment clearly facilitated acquisition of trace eyeblink conditioning and decreased AChE activity (thus, increased ACh throughout the brain) in aging rabbits. After acquiring the task and after metrifonate treatment was ceased, the metrifonate-treated rabbits still displayed asymptotic behavior (Kronforst-Collins et al., 1997b), suggesting that the cholinergic input was important for acquisition but not the recall of the association.

Recently, Zhang et al. (1997) have demonstrated that AChE inhibition led to an increased (as long as 30 sec) temporal “window” for ACh to modulate the current underlying the slow AHP (I_{AHP}) (i.e., reduce the AHP) in CA1 neurons in vitro after a train of cholinergic afferent stimulation in CA1 stratum radiatum; without the AChE inhibitors, the window was <5 sec. They further speculated that the window may be shorter in vivo. They have also demonstrated that the stimulation of the cholinergic afferents must precede the CA1 depolarization (activity) by 400–1500 msec to achieve the I_{AHP} reduction (Zhang et al., 1996). It is possible that metrifonate in vivo increases the ACh level in the synapses of pyramidal neurons, prolonging the temporal window for ACh modulation of pyramidal neuronal activity and resulting in an increased signal-to-noise ratio that facilitates the acquisition of various behavioral tasks in aging and cholinergic-deficient (AD and medial–septal or scopolamine lesioned) subjects. In trace eyeblink conditioning, where a stimulus-free trace period separates the conditioned and unconditioned stimuli, the AChE inhibition should increase the ACh present in the synapses of hippocampal pyramidal neurons during the trace period. This would help the subject to associate the conditioned and unconditioned stimuli and enhance the acquisition rate in aging subjects.

In summary, the increased neuronal excitability (as evidenced by the reduced postburst afterhyperpolarization and spike frequency adaptation) of hippocampal pyramidal neurons may be one of the underlying mechanisms by which the hippocampus stores information during associative learning. As previously demonstrated, the AHP and accommodation are reduced in hippocampal neurons when a subject learns an associative task but not in subjects that did not learn the same associative task (Moyer et al., 1996; Thompson et al., 1996b). Furthermore, the AHP and accommodation are greater in neurons from aging subjects than from the young (Landfield and Pitler, 1984; Moyer et al., 1992); in the same aging population, a severe impairment in learning associative tasks (such as trace eyeblink conditioning) is observed (Thompson et al., 1996a). The learning deficit observed in the aging population is alleviated by pharmacological agents, such as metrifonate (Kronforst-Collins et al., 1997a,b) and the L-type calcium channel blocker nifedipine (Deyo et al., 1989; Kowalska et al., 1994). These same two compounds have been demonstrated to increase hippocampal pyramidal neuronal excitability (reduced AHP and accommodation) in vitro (see results and Moyer et al., 1992). Thus, increased neuronal excitability of hippocampal pyramidal neurons may be one mechanism by which memory is transiently stored in the hippocampus during the early stages of learning before being transferred to the neocortex for

<table>
<thead>
<tr>
<th>Postburst afterhyperpolarization</th>
<th>n</th>
<th>Amplitude (mV)</th>
<th>% AHP Δ</th>
<th>Area (mVsec)</th>
<th>Duration (sec)</th>
<th>Accommodation (# of action potentials)</th>
<th>Input resistance (MO)</th>
</tr>
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<tbody>
<tr>
<td>Young</td>
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<td></td>
<td></td>
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<tr>
<td>aCSF</td>
<td>13</td>
<td>−0.40 ± 0.35</td>
<td>−10.44 ± 4.93</td>
<td>−0.78 ± 0.79</td>
<td>0.37 ± 0.71</td>
<td>0.63 ± 0.40</td>
<td>1.28 ± 2.20</td>
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<tr>
<td>Atropine (1 μM)</td>
<td>5</td>
<td>0.11 ± 0.48</td>
<td>−1.89 ± 11.93</td>
<td>−0.14 ± 0.83</td>
<td>0.27 ± 0.28</td>
<td>1.25 ± 0.69</td>
<td>2.56 ± 7.05</td>
</tr>
<tr>
<td>Carbachol (500 nm)</td>
<td>11</td>
<td>−1.18 ± 0.30**</td>
<td>−29.25 ± 7.30</td>
<td>−1.74 ± 0.54**</td>
<td>−1.23 ± 0.69</td>
<td>4.74 ± 0.99***</td>
<td>3.45 ± 2.52</td>
</tr>
<tr>
<td>Eserine (500 nm)</td>
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<td>0.75 ± 0.45</td>
<td>−12.16 ± 11.53</td>
<td>−0.36 ± 0.64</td>
<td>0.91 ± 0.39</td>
<td>2.11 ± 0.60**</td>
<td>2.41 ± 3.48</td>
</tr>
<tr>
<td>Eserine (1 μM)</td>
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<td>−0.38 ± 0.12**</td>
<td>−16.38 ± 4.31</td>
<td>−0.51 ± 0.36</td>
<td>−0.50 ± 0.36</td>
<td>2.12 ± 0.48***</td>
<td>3.65 ± 1.30*</td>
</tr>
<tr>
<td>Eserine (5 μM)</td>
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<td>−1.07 ± 0.33*</td>
<td>−27.62 ± 7.07</td>
<td>−1.83 ± 0.62*</td>
<td>−0.85 ± 0.40</td>
<td>4.81 ± 0.69***</td>
<td>−2.03 ± 2.38</td>
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<tr>
<td>aCSF</td>
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<td>14.87 ± 10.05</td>
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<td>1.11 ± 0.78</td>
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<td>Carbachol (500 nm)</td>
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<td>−1.17 ± 0.27***</td>
<td>−22.74 ± 5.82</td>
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<td>−0.89 ± 0.38*</td>
<td>4.80 ± 0.90***</td>
<td>3.30 ± 2.27</td>
</tr>
</tbody>
</table>

n indicates the number of cells recorded. The numbers in bold type indicate statistical significance using paired t test. The measurements are the mean ± SEM (*p < 0.05; **p < 0.01; ***p < 0.001; paired t tests).
more permanent storage. Pharmacological agents that cause an excitability increase comparable to that which occurs during the acquisition process in young subjects appear to enhance learning in normal aging subjects and perhaps in those undergoing a neurodegenerative process such as Alzheimer's disease.

REFERENCES


