Corticosteroid Regulation of Ion Channel Conductances and mRNA Levels in Individual Hippocampal CA1 Neurons

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Overexposure to corticosteroid hormones is harmful to hippocampal neuronal integrity, likely by perturbation of calcium homeostasis. To identify molecular mechanisms at the single-cell level, we characterized mRNA expression corresponding to voltage- and ligand-gated Ca channels in individual dissociated CA1 neurons in response to long-term corticosterone (CORT) exposure. Predominant mineralocorticoid receptor occupation (ADC-HI group) resulted in increased amplitudes, particularly of L-type Ca channel mRNAs, high levels of GluR-2 versus GluR-1, and a high ratio of NMDAR-2A to NMDAR-2B mRNA. Corresponding alterations in protein expression were consistent with the restriction of Ca influx. In contrast, additional glucocorticoid receptor occupation (ADC-HI group) altered the expression of these mRNAs in a manner consistent with enhanced Ca influx; interestingly, qualitatively similar alterations were seen in control ADX neurons. Electrophysiological data from the same neurons indicate that Ca current amplitudes also are modulated by CORT, although on a shorter time scale. Finally, principal components analysis (PCA) suggests that neuronal AMPA and NMDA receptor composition may be regulated by MR and GR activation in a complex manner. Therefore, our data implicate molecular events by which CORT may regulate Ca influx into CA1 hippocampal neurons.

Key words: corticosterone; dissociated CA1 Neurons; aRNA amplification; expression profile; mRNA; ion channels; hippocampus

Adrenal corticosteroid hormones such as corticosterone (CORT) mediate a wide range of metabolic, immune-suppressive, and anti-inflammatory responses to stress (Bohus et al., 1982; Funder, 1987). CORT binds with high affinity (Kd = 0.5 nM) to mineralocorticoid receptors (MRs) and with 10-fold lower affinity (Kd = 5 nM) to glucocorticoid receptors (GRs) in rat hippocampal CA1 neurons (Hollenberg et al., 1985; Reul and de Kloet, 1985; Arriza et al., 1987; Chao et al., 1989; Herman et al., 1989). These ligand-bound receptors interact with sequence-specific genomic response elements and regulate transcriptional activity (Rousseau, 1984; McEwen et al., 1986; Funder, 1987). Therefore, the actions of CORT on MRs and GRs alter the expression of cellular mRNAs and proteins and influence fundamental properties of hippocampal neurons.

Selective occupation of MRs by CORT is associated with small amplitudes of both T- and L-type Ca currents in CA1 neurons from slice preparations (Karst et al., 1994). Additional activation of GRs resulted in increased amplitudes, particularly of L-type currents (Kerr et al., 1992; Karst et al., 1994). Other Ca-dependent phenomena, e.g., the accommodation in cell firing and slow after hyperpolarization observed with brief depolarization of CA1 neurons, also displayed a comparable dependency on the extent of MR and GR occupation (Joëls and de Kloet, 1989, 1994; Kerr et al., 1989). Corticosteroid receptor-mediated effects on Ca conductances and other neuronal properties were slow in onset and required protein synthesis (Karst and Joëls, 1991; Kerr et al., 1992), suggesting a genomic component of action.

Many reports in the literature suggest that MR activation can be neuroprotective in the hippocampus (Landfield et al., 1981; Gould et al., 1990; Joëls and de Kloet, 1996). Chronically, absence of circulating CORT in ADX rats, as well as exposure to high doses of CORT, increases neuronal vulnerability, sometimes resulting in cell death (Sapolsky et al., 1985; Sloviter et al., 1989; Gould et al., 1990; Woolley et al., 1990), particularly when concomitant excitatory challenges are present (Sapolsky and Pulsinelli, 1985; Sapolsky et al., 1988). Chronically elevated Ca influx associated with a high dose of CORT is thought to endanger hippocampal neurons (Elliot and Sapolsky, 1993; Elliot et al., 1993) via NMDA receptor-dependent processes (Choi, 1988; Armanini et al., 1990; Tymianski et al., 1994).

In the present study we examined whether prolonged changes in average plasma CORT levels affect mRNA expression for proteins contributing to Ca influx and Ca-mediated synaptic transmission in CA1 pyramidal neurons. These included the α1 subunits of neuronal Ca channels, e.g., α1A, α1B, and α1C/D, encoding P-, N- and L-type Ca channels, respectively (Varadi et al., 1995), subunits of the AMPA (GluR-1, R-2, R-3, and R-4) and NMDAR-1, R-2A, R-2B, and R-2C receptors. To this end, we used the technique of single-cell antisense RNA amplification and expression profiling (Eberwine et al., 1992), subsequent to electrophysiological recording of voltage-gated Ca currents from
isolated CA1 neurons (Kay and Wong, 1986). Thus, regulation of Ca conductances, as well as expression of mRNAs encoding Ca channel proteins, was characterized in the same neurons. Four experimental groups were investigated: (1) rats that were adrenalectomized 3–4 weeks before the experiments so that MRs and GRs were chronically unoccupied (ADX); (2) ADX rats that received a low dose of CORT (20 μg/ml) in their drinking water so that mostly MRs, but not GRs, were activated (ADC-LO) (Jacobson et al., 1988); (3) ADX rats that received a very high CORT dose (300 μg/ml), resulting in activation of GRs in addition to MRs (ADC-HI); and (4) sham-operated controls (SHAM). These treatments were performed for 4 weeks.

MATERIALS AND METHODS

Treatment paradigm. Groups of adult male Sprague Dawley rats (average weight, 100–200 gm) were adrenalectomized bilaterally or sham-operated and treated for 4 weeks with 0.9% saline and 1.5% ethanol in their drinking water. The following additions were made to their water 1–2 d after the operation: ADX, none; SHAM, none; ADC-LO, CORT (20 μg/ml); ADC-HI, CORT (300 μg/ml). On the day of the experiment, rats were exposed for 30–60 min to a novel environment and then either anesthetized or killed. Their brains were dissected quickly and stored at 0°C in artificial CSF of the following composition (in mM): NaCl 120, KCl 5, CaCl 2 1, MgCl 2 1, and D-glucose 25, pH 7.0, was added (final concentration 100–200 μg/ml RNase inhibitor). Subsequently, reagents were added for first-strand cDNA synthesis. The composition and approximate final concentrations (because of variations in volume of the aspirated cell content) of these reagents were (in mM) 130 K-gluconate, 10 KCl, 6 MgCl 2 , and 10 HEPES with 500 ml RNase inhibitor, and tRNA, NaCl (final concentrations 25 ng/ml and 0.1 M each dATP, dCTP, dGTP, and TTP, plus 1.5 ng/ml oligo-dT-17 primer. This mixture was heated to 90°C for 10 min and then cooled to 39°C, after which the AMV Seikagaku reverse transcriptase was added (final concentration 1 U/ml) in the presence of which the mixture was incubated for another 90 min (at 39°C). Next, the mixture was cooled to 0°C and tRNA, NaCl (final concentrations 25 ng/ml and 0.1 M each dATP, dCTP, dGTP, and TTP, respectively), and ethanol were added to precipitate first-strand cDNA.

Antisense RNA amplification: first and second round. The single-stranded cDNA population was purified by phenol-chloroform extraction and ethanol precipitation. Then it was made double-stranded by the Gubler–Hoffmann method; the hairpin loop was excised and blunt-ended by standard procedures. The cDNA template was purified by dialysis before incubation with the following reagents at 37°C for 3.5 hr to generate 32P-labeled antisense RNA (aRNA) (in mM): 40 Tris buffer, pH 7.5, 7 MgCl 2 , 10 NaCl, 2 spermidine, and 8 DTT plus 250 μM ATP, GTP, and UTP, 25 μM CTP, 30 μCi of [32P] CTP, 20 U of RNase inhibitor, and 1000 U of T7 RNA polymerase. A fraction of the 32P-labeled aRNA was electrophoresed on a denaturing formaldehyde gel. Then the gel was washed in 10% w/v trichloroacetic acid (TCA), blotted dry for 12 hr, and apposed to x-ray film to assess the size distribution of the first-round aRNA population.

Purified first-round aRNA was incubated at 37°C with 100 ng of random primers, 40 U of RNase inhibitor, 40 U of Seikagaku AMV reverse transcriptase, and 250 μM dNTPs under appropriate buffer conditions (in mM) 50 Tris, pH 8.0, 6 MgCl 2 , 120 KCl, and 7 DTT to generate the corresponding complementary DNA population. The resulting cDNA population was subjected to the sense orientation relative to the original poly(A) RNA population and had poly(dA) tails. The oligo-dT-17 amplification primer was used to synthesize double-stranded cDNA by using T4 DNA polymerase. This cDNA population was made blunt-ended as before, dialyzed, and amplified to generate a 32P-labeled aRNA population antisense in orientation to the original poly(A) population. Under optimal conditions, resulting aRNA was amplified linearly over the latter by greater than a million-fold. This was used to probe slot blots loaded with candidate cDNA clones (below).

Preparation of slot blots and probe addition. Amersham's Hybond-N nylon membrane (Arlington Heights, IL) was wetted with distilled deionized H 2 O and 10× SSC before use in the slot blot apparatus. One microgram of each linearized plasmid cDNA in 10× SSC corresponding to each candidate mRNA sequences was heat-denatured (85°C for 7 min) and loaded per slot under gravity. The cDNAs were fixed to the nylon membrane by UV-cross-linking.

Slot blots were prehybridized in heat-sealed plastic bags containing 50% v/v formamide, 5× SSC, 5× Denhardt's reagent, 0.5% w/v SDS, 100 μg/ml salmon sperm DNA, and 1 μM Na pyrophosphate for 12 hr.

32P-labeled aRNA probe (not less than 10 7 counts) was heat-denatured and added to the bags. After a hybridization period of ~48 hr, blots were washed two times for 15 min in 2× SSC, 0.1% w/v SDS at 42°C, followed by two times for 15 min washes in 0.1× SSC, 1% w/v SDS at 55°C. The single-wash was repeated until the signal corresponding to 1 μg of plasmid vector cDNA on the slot blots that represented nonspecific binding of the aRNA probe was judged to be sufficiently low. The blots were air-dried and apposed to a phosphor screen, which then was analyzed on a phosphorimager with ImageQuant software.

Data analysis. The density of bands corresponding to candidate cDNA clones was measured by analysis on the phosphorimager. The specific signal (above background) of probe bound to each clone was expressed as a percentage of the sum of specific signals for the α-subunits of the neuronal GABA A receptor—the “inhibitory component” of CA1 neurons. With the exception of α1 mRNA, there were no alterations in other α-subunit mRNAs among treatment groups. Because α1 mRNA represented between 2 and 4% of total α-subunit mRNA signal in all neurons examined, this difference is unlikely to have an impact on our analysis.

Use of internal references minimized variations caused by differences in specific activity of the probe and absolute quantity of the probe present. Relative abundances of multiple candidate mRNAs in individual CA1 neurons from each treatment group were characterized. Further, alterations in relative steady-state mRNA levels in CA1 neurons from ADC-LO and ADC-HI treatment groups were quantified relative to the ADX-V and SHAM control groups. Such an analysis of concomitant expression of multiple candidate mRNAs is called expression profiling.

Data from the four groups were tested first with a one-way ANOVA, followed by a post hoc multiple comparison of the means, using a Student's post hoc test. Principal components and univariate ratio analyses. Uni- and multivariate statistical procedures were performed on mRNA expression data for 10 genes generated from individual SHAM CA1 neurons. Alterations in the relative levels of gene expression may be correlated with changes in
The four experimental groups comprised a group of 4 week adrenalectomized rats (ADX), a group of ADX rats treated with 300 µg/ml CORT in the drinking water (ADC-HI), and the sham-operated controls (SHAM).

As expected, SHAM animals showed the greatest increase in weight, whereas ADC-HI animals showed minimal weight gain over the same treatment period. Average daily intake of CORT per animal was calculated from daily water intake; when plasma CORT levels were measured at the start of the experiment, both untreated ADX and ADC-LO groups exhibited significantly lower amounts than the ADC-HI group or SHAM-operated controls (p < 0.01). Plasma CORT levels in SHAMs were also significantly higher than in the ADC-HI group (p < 0.05).

Next, univariate analyses were performed on the ratio conveyed by the selected PC. The ratio was constructed mathematically in SAS (Statistical Analysis Systems, 1990), using only those genes with coefficients that dominated the PC structure (≥0.2). The univariate procedures calculated the means of this ratio for each treatment group and generated comparisons of their statistical significance (p < 0.05) across treatment groups, using the least-squares means option in the general linear models (GLM) procedure extension of ANOVA. Statistical significance for all of the univariate analyses was set at the α 0.05 (p < 0.05) level (Sokal and Rohlf, 1981). The Hartley’s F max statistic was used to test for equality of variances among treatment groups for the ratio of gene expression (Mason et al., 1989). All statistical computations, with the exception of the Hartley’s F max statistic, were performed by SAS.

RESULTS

We examined the effects of chronic exposure to CORT on electrophysiological and molecular markers of neuronal Ca influx in individual CA1 neurons. As an experimental approach to alter chronically the daily CORT intake, we adrenalectomized rats and offered them low or high CORT doses in their drinking water. Previous studies indicate that in this paradigm rats still exhibit a considerable degree of variation in circulating CORT levels during the day because of differences in activity levels and drinking patterns, similar to what occurs in adrenally intact rats, but not with pellet administration of CORT (Akana et al., 1988; Jacobson et al., 1988). From their fluid intake it is clear that CORT was ingested successfully by both the ADC-LO (0.76 mg/rat per day) and the ADC-HI groups (10.28 mg/rat per day) (Table 1). Further, the average weight gain in the ADC-HI group was significantly lower than the weight gains in the other treatment groups (Table 1), suggesting that the former group ingested high levels of CORT.

The four experimental groups differed with respect to their daily CORT intake (Table 1). In addition, because daily variations in circulating CORT levels previously have been observed (Akana et al., 1988; Jacobson et al., 1988), we thought it necessary to examine how mRNA expression and Ca current properties correlate both with circulating CORT levels several hours before the experiment as well as with the average daily CORT intake shown in Table 1. Interestingly, plasma CORT levels at the start of the experiment observed for the ADX and ADC-LO groups were extremely low for both, although daily CORT exposure is different for these two groups. This is likely attributable to the time of sampling relative to drinking patterns in the context of the

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Table 1. Correlations among treatment groups, average daily CORT intake, measured plasma CORT levels, and Ca conductance properties

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Initial weight</th>
<th>Final weight</th>
<th>[CORT] intake</th>
<th>[CORT]-RIA</th>
<th>Capacitance</th>
<th>Input impedance</th>
</tr>
</thead>
<tbody>
<tr>
<td>SHAM</td>
<td>159.3 ± 4.4</td>
<td>305.3 ± 6.8</td>
<td>0.0 ± 0.0</td>
<td>11.6 ± 3.0</td>
<td>8.8 ± 0.6</td>
<td>502 ± 46</td>
</tr>
<tr>
<td>ADX</td>
<td>156.0 ± 8.5</td>
<td>262.1 ± 9.2</td>
<td>0.3 ± 0.8</td>
<td>7.3 ± 0.5</td>
<td>8.0 ± 147</td>
<td>9.118</td>
</tr>
<tr>
<td>ADC-LO</td>
<td>166.0 ± 12.1</td>
<td>290.0 ± 14.5</td>
<td>0.5 ± 0.2</td>
<td>8.6 ± 0.5</td>
<td>633 ± 69</td>
<td>7.127</td>
</tr>
<tr>
<td>ADC-HI</td>
<td>146.5 ± 3.9</td>
<td>154.5 ± 12.3</td>
<td>4.1 ± 1.0</td>
<td>8.5 ± 0.7</td>
<td>827 ± 124</td>
<td>6.916</td>
</tr>
</tbody>
</table>

Four experimental groups were administered CORT or vehicle daily in their drinking water ([CORT] intake); their weights were recorded before (Initial weight) and after the treatment period (Final weight). Average daily [CORT] ingested was calculated from fluid intake. On the day of the experiment, they were tested for plasma corticosterone levels ([CORT]-RIA). Cellular capacitance and input impedance were determined before whole-cell patch-clamp recording. The number of animals subjected to the indicated treatment (N), cells used for the single cell RNA profiling (n), and electrophysiology (n) are indicated in the last column; molecular and electrophysiological data were not obtained in all animals in which plasma CORT was determined. The four experimental groups comprised a group of 4 week adrenalectomized rats (ADX), a group of ADX rats treated with 300 µg/ml CORT in the drinking water (ADC-HI), and the sham-operated controls (SHAM).
extremely short plasma half-life of CORT (Jacobson et al., 1988; Sloviter et al., 1989; Shors et al., 1990). Further, plasma CORT levels in sham-operated animals are greater than those in ADX animals treated with high CORT, likely attributable to the acute stress related to their exposure to a novel environment 30–60 min before their deaths. Acutely dissociated CA1 neurons were examined first for effects of steroid treatment on whole-cell Ca conductance properties, as described elsewhere (Vreugdenhil and Wadman, 1992). Actual Ca current amplitudes correlated with plasma CORT levels shortly before the experiment, rather than with changes in daily average CORT intake (see Whole-Cell Calcium Currents below).

Next, cellular contents were aspirated into the recording pipette. Single-cell antisense RNA amplification and expression profiling were performed to characterize simultaneous expression of candidate mRNAs as a function of long-term CORT treatment in individual CA1 neurons (Eberwine et al., 1992). All data analysis was performed with ImageQuant software. Specific hybridization to each cDNA clone was calculated by subtracting nonspecific hybridization (to control vector sequence) from total hybridization. Within each blot, specific signals for cDNA clones corresponding to voltage-gated Ca channels and NMDA receptor subunits were expressed relative to the “inhibitory component,” i.e., the sum of specific signals for α-subunits (α1–α6) comprising the GABA_A receptor in the same blot. Recent in situ hybridization data suggest that neither adrenalectomy nor CORT replacement (100 mg/ml in drinking water, i.e., 5× the CORT concentration administered to the ADC-HI group) led to alterations in levels of α-subunit mRNAs comprising the GABA_A receptor in the CA1 (Orchinik et al., 1994). Levels of AMPA receptor mRNAs were expressed relative to the sum of the specific signals for all AMPA receptor mRNAs. Principal component (PCA) and univariate ratio analyses additionally were performed on expression data from individual samples in each treatment group. These procedures were performed jointly to identify mRNA combinations that reflect cellular activities involved in normal CA1 neuronal behavior and their possible regulation by exposure to CORT.

Voltage-gated calcium channel mRNAs

Hybridization to cDNAs corresponding to α1A, α1B, and α1C/D subunit mRNAs (encoding P/Q-, N-, and L-type voltage-gated Ca channels, respectively) was detected in every individual CA1 subunit mRNAs (encoding P/Q-, N-, and L-type voltage-gated Ca

Calcium Currents below).

In the present study, individual CA1 neurons coexpressed various genes for AMPA receptor subunits (Glur1 through Glur-4) were characterized as a function of differential corticosteroid receptor occupation. Glur1 mRNA was, relatively speaking, the most abundant transcript, whereas Glur-4 mRNA was barely detectable in neurons from sham-operated animals. Thus, Glur1 through Glur-3 subunit mRNAs were coexpressed in individual CA1 neurons.

In the present study the relative levels of Glur1 and Glur-3 subunit mRNAs were comparable for all treatment groups (Fig. 3A) (data not shown). Glur-2 mRNA expression was not altered after 4 weeks of adrenalectomy relative to sham controls. However, ADC-LO neurons expressed ~74% more Glur-2 mRNA relative to untreated ADX and SHAM neurons (p < 0.001; Fig. 3B). This increase was reversed in ADC-HI neurons, which expressed ~49% less Glur-2 mRNA relative to the low CORT-treated group.

Selective occupation of MRs increased the relative abundance of Glur-2 mRNA, such that the ratio of Glur1 to Glur-2 mRNA levels was ~2:1. Lack of occupation of MRs and GRs in untreated ADX neurons, as well as their simultaneous occupation in ADC-HI neurons, decreased the relative abundance of Glur-2 mRNA, resulting in a Glur1 to Glur-2 mRNA ratio of ~3:1.

NMDA receptor subunit mRNAs

In the present study, individual CA1 neurons coexpressed NMDA receptor mRNAs for the glutamate-responsive NR1, NR2A, 2B, and 2C subunit proteins. In SHAM neurons, the relative order of abundance was NR1 mRNA (35%), NR2A (27.5%), NR2B (17.5%), and NR2C (20%).

No alterations were detected in levels of NR1 (Fig. 4) or NR2C mRNAs among any of the experimental groups. Similarly, no differences were observed in NR2A and NR2B expression in

Figure 1. Expression profile of various candidate genes in hippocampal pyramidal cells. One microgram each of cDNA containing plasmid for various genes was linearized with EcoRI or HindIII, heat-denatured, and gravity-applied to a nylon membrane filter, using a slot blower. The filter was UV-cross-linked with a Stratalinker (Stratagene, La Jolla, CA), followed by prehybridization as described (Craig et al., 1997). Radiolabeled aRNA from an individual neuron was hybridized for 2 d as described (Craig et al., 1997). The filters were washed with increasing stringencies to a final wash stringency of 0.1× SSC, 0.1% SDS at 55°C for 30 min. The filters either were placed on a phosphomager screen for analysis or were apposed to film for 3 d, followed by developing in an automated film developer.
ADX neurons when compared with SHAMs. ADC-LO neurons expressed significantly higher levels of NR2A mRNA than neurons in either the SHAM or ADX group \((p < 0.05)\); no further alteration was observed in ADC-HI neurons (Fig. 4B). On the other hand, NR2B mRNA levels were significantly lower in ADC-LO neurons relative to the SHAM and ADX groups \((p < 0.05)\) and to the ADC-HI group \((p < 0.001;\) Fig. 4C). The ratios of NR2A to NR2B mRNA levels in ADX and ADC-HI neurons were 1.6:1 and 1.8:1, respectively. However, in low CORT-treated neurons this ratio was 6:1 (Fig. 4D).

**Whole-cell calcium currents**

Ca currents were evoked in acutely dissociated neurons by the voltage protocol depicted in Figure 5A. As described elsewhere,
these currents mainly consisted of L- and N-type Ca currents (Thompson and Wong, 1991). From each cell the membrane input impedance, capacitance, and the current–voltage relation-ship for the peak amplitude of the Ca current were determined (Table 1).

Average peak Ca current amplitude for a voltage step to 10 mV was not different in the ADC-LO group (1.1 ± 0.1 nA) when compared with the untreated ADX group (1.0 ± 0.1 nA; SHAM-operated controls, 1.3 ± 0.1 nA). As shown in Figure 5B, we also tested peak Ca current amplitudes as a function of plasma CORT levels at the start of the experiment. Ca current amplitudes were relatively high when considered in the group of animals with extremely low (0–1 μg CORT/100 ml plasma) or moderately high (>5 μg/100 ml plasma) circulating CORT levels, whereas significantly lower Ca current amplitudes were observed in neurons from animals with intermediate (2–5 μg/100 ml plasma) CORT levels (Fig. 5B).

The data suggest that, in the present experimental paradigm, peak Ca current amplitudes may correlate with circulating CORT levels several hours before the start of the experiment rather than with average daily CORT intake.

**Principal components and univariate ratio analyses**

SHAM mRNA expression data for the 10 genes described above were consolidated into eigenvector (coefficient)-based variables called principal components or PCs (see Materials and Methods). PC1 has the highest variance (32.8%; data not shown) relative to the remaining PCs, because it expresses the largest percentage of variability in mRNA expression in SHAM neurons; each successive PC seeks to decrease this variability. We specifically selected PC10 for further analysis because it suggested a biologically plausible scenario in addition to accounting for the smallest percentage of variability, i.e., <0.01%. The coefficients with the highest values (≥0.2) were identified; the gene combination (GluR2.NMDAR-2A)/(GluR3.NMDAR-1) dominated PC10, making it an excellent candidate for a “diagnostic” molecular fingerprint. This gene subset is used in all further discussions of PC10, with genes represented by positive coefficients (GluR-2 and
NMDAR-2A) in the numerator and those with negative coefficients (GluR-3 and NMDAR-1) in the denominator.

In Figure 6, the PCs plot and bar graph represent the treatment group relationships determined by this initial analysis. CA1 neurons from each treatment group are plotted against the numerator and denominator genes that characterize PC10 (Fig. 6b), revealing cellular distributions within each treatment group. SHAM CA1 neurons are clustered closely around their mean, indicating that this ratio was expressed consistently from one neuron to the next. ADC-HI CA1 neurons also demonstrated a linear relationship, whereas ADX and ADC-LO neurons were clustered in nonlinear groups, implying erratic expression of these genes in the latter. The variance test determined that the distributions of the neuron groups were statistically distinct from one another (p, 0.05), with the exception of the comparison between ADC-HI and SHAM neurons.

Chronic ADX treatment did not alter the mean PC10 ratio (Fig. 6b) relative to SHAMs. However, ADC-LO neurons expressed a mean PC10 value distinct from the means of the other three groups (p < 0.05), whereas ADC-HI neurons were not distinguishable from SHAM or ADX neurons (Fig. 6b). There are also significant slope differences among SHAM, ADC-LO, and ADC-HI groups (Fig. 6b), i.e., CORT treatment has a disruptive effect on the relationship between the numerator and denominator genes expressed in SHAM neurons.

The neuronal interrelationships illustrated in Figure 6a, together with the least-squares mean ratio expression values (Fig. 6b) and Hartley’s F_{max} test for variance equality (data not shown), distinguished the neuron groups. From Figure 6b it is clear that an inverted U-shaped dose relationship exists between the extent of MR and GR occupation and the PC10 ratio of gene expression in CA1 neurons.

DISCUSSION

In this study we observed that chronic activation of MRs and GRs by CORT regulates expression levels of mRNAs encoding voltage- and ligand-gated ion channels in individual CA1 neurons. This suggests that CORT may exert receptor-dependent actions on fundamental properties of these neurons, including ion influx and synaptic transmission, as well as susceptibility to neurodegenerative processes.

We first examined the α1 subunits of voltage-gated Ca channels, the sequences for which have been cloned recently (Dunlap et al., 1989). Of these, the α1A subunits constitute channels with properties of the P/Q-type Ca channels, α1B subunits of N-type channels, and α1C/D subunits of L-type Ca channels (Varadi et al., 1995). In our study chronically altered CORT intake affects relative mRNA expression levels of subunits of the P/Q- and L-type channels, but not of the N-type Ca channel. Specifically, in the absence of steroid, increased expression was observed for the L-type channel subunit mRNA and, to a lesser extent, for the P/Q-type channel subunit mRNA. However, P/Q- and L-type Ca...
channel subunit mRNA levels were low in ADC-LO neurons (animals that chronically received low CORT in their drinking water). Finally, chronic exposure to high CORT led to high levels of expression for the P/Q-type and L-type channel subunit mRNAs in ADC-HI neurons relative to ADC-LO neurons. P-type Ca channels are associated predominantly with stimulus-coupled exocytosis in the mammalian brain (Dunlap et al., 1989) and are present at both excitatory and inhibitory synapses (Luebke et al., 1993; Takahashi and Momiyama, 1993; Castillo et al., 1994) in the hippocampus (Llina´s et al., 1992; Mintz et al., 1992). Therefore, if the mRNA changes observed in our study (see Fig. 2) lead to alterations in protein expression, chronically negligible or high CORT may increase the relative contribution of P/Q-type Ca current and consequently may modulate exocytosis mediated by these channels. Conversely, our data imply that chronic low CORT may lead to an opposite shift in P/O-type Ca current-mediated function. Expression levels of mRNAs encoding α-CgTx-sensitive N-type Ca channels are not altered by either dose of CORT after 4 weeks. This fraction of the Ca current is associated with cooperative regulation of stimulation-coupled secretion events in hippocampal neurons, along with P/O-type Ca channels (Dunlap et al., 1989; Wheeler et al., 1994). Therefore, CORT may not regulate N-type current-mediated synaptic transmission by actions on either MRs or GRs in CA1 neurons. Further, chronic absence of CORT or high CORT may increase the relative numbers of L-type Ca channels, whereas low CORT results in mRNA levels that correlate with decreased expression. L-type channels regulate levels of cytosolic calcium (Dunlap et al., 1989; Hell et al., 1993) and can activate other channels directly, regulate gene expression (Morgan and Curran, 1986; Murphy et al., 1991), and activate a number of Ca-dependent protein kinases and phosphatases (Kennedy, 1989). Differential corticosteroid receptor activation therefore may modulate Ca-dependent signal transduction mechanisms mediated by L-type channels in CA1 neurons.

Interestingly, although the potential for Ca influx in animals subjected to low daily CORT intake may be downregulated, measured Ca currents in ADC-LO were seen to be comparable to those from SHAM controls. Accordingly, whole-cell Ca current amplitudes correlated with plasma CORT levels at the start of the experiment (see Fig. 5)—as opposed to average daily CORT intake—displaying a U-shaped dose dependency, as has been observed previously for the steroid regulation of Ca currents and other electrical responses of hippocampal cells in slices (Joëls and de Kloet, 1994; Karst et al., 1994). This may be because acutely dissociated neurons lack part of the dendritic tree where steroid-sensitive Ca channels have been shown to be located (Karst et al., 1993, 1994); MR-mediated events also have been shown to be less apparent in this neuronal preparation (Werkman et al., 1997). Further, Ca currents recorded in the whole-cell configuration show fewer differences in their dynamic range (Zhu and Ikeda, 1994) so that downregulation of Ca channel number may have been less apparent in terms of whole-cell Ca conductances. Other ionic conductances, either not regulated by CORT or not examined in this study, also may contribute to measured whole-cell Ca currents in CA1 neurons. Finally, observed differences between electrophysiological and molecular data may be attributable to the actions of CORT at different regulatory sites, e.g., regulation of protein synthesis to modulate Ca currents (Karst and Joëls, 1991; Kerr et al., 1992) in addition to regulation of mRNA expression levels over the long term. The precise

Figure 6. Plots of the numerator and denominator genes comprising the PC10 ratio derived from the expression data (additive coefficients are based on log-transformed cpm's) for AMPA and NMDA receptor subunit mRNAs. a. Each point represents a single neuron from SHAM, ADX, ADC-LO, and ADC-HI animals. Numbers on the ordinate and abscissa for each plot reflect the scale resulting from construction of separate linear combinations of PC10 coefficients representing the numerator (GluR-2 + NMDAR-2A) and denominator (GluR-3 + NMDAR-1), respectively. The tight clustering of SHAM CA1 data points along its line indicates a unified cellular response to the ratio. Dispersal from the central line represents an alteration, i.e., a “disruption” in the unified cellular response to the ratio. *Significant slope differences relative to SHAM neurons. b. Shown are mean expression level values and SE of the PC10 ratio across treatment groups; means with the same symbol are not significantly different (p > 0.05).
Ionotropic glutamate receptors responsive to AMPA mediate most of the rapid synaptic excitatory transmission in the CNS (Sommer and Seeburg, 1992). Assembly of the GluR-2 subunit has been shown to be responsible for the very low Ca permeability and linear current–voltage (I–V) relationship exhibited by native AMPA receptors (Hollmann et al., 1991; Verdoorn et al., 1991; Jonas and Sakmann, 1992). In our study the contribution of GluR-2 relative to GluR-1 mRNA was increased significantly in ADC-LO animals, as compared with the ADX and ADC-HI groups (see Fig. 3). This suggests that predominant activation of MRs may restrict AMPA receptor-gated current, whereas lack of activation of either MRs or GRs or their simultaneous activation may enhance this current in CA1 neurons. Therefore, the extent of occupation of MRs and GRs may modulate resting Ca conductance and fast synaptic transmission mediated by AMPA receptors. GluR-1, R-2, and R-3 subunit mRNAs are expressed in alternatively spliced forms in the hippocampus (Sommer et al., 1990), whereas GluR-2 mRNA undergoes RNA editing (Hollmann et al., 1991). We currently are investigating the potential roles of RNA editing and alternative splicing of GluR-2 mRNA in the context of our observations.

The NMDA glutamate-responsive receptor is distributed widely in the CNS. This ligand-gated ion channel mediates cationic synaptic transmission and plays a pivotal role in developmental synaptic plasticity and excitotoxic cell death (Mayer and Westbrook, 1987; Tymianski et al., 1994). In previous experiments NR1, NR2A, and NR2B mRNAs were shown to be expressed at moderate to high levels in pyramidal neurons in the hippocampus (Monyer et al., 1992, 1994). These investigators detected NR2C transcripts only in a subset of hippocampal neurons, presumably interneurons, at much lower levels. In our study, however, individual CA1 neurons were seen to coexpress NR1, 2A, 2B, and 2C mRNAs (see Fig. 1). Our ability to detect NR2C mRNA in every CA1 neuron that was examined may be attributable to the increased sensitivity of detection afforded by single-cell aRNA analysis relative to in situ hybridization studies previously performed on whole-brain sections (Eberwine et al., 1992). This study represents the first observation that NR1, NR2A, 2B, and 2C mRNAs are coexpressed in individual CA1 neurons from the postnatal rat hippocampus.

CA1 neurons with predominant activation of MRs (ADC-LO) expressed a significantly higher ratio (6:1) of NR2A to NR2B mRNA relative to neurons in which corticosteroid receptors were either simultaneously activated (ADC-HI) or simultaneously inactivated (ADX) (1.6:1 and 1.8:1, respectively) (see Fig. 4). In previous experiments, recombined heteromeric NR1–NR2A channels from rat brain exhibited faster (4×) offset decay times but similar reversal potentials relative to recombinant NR1–NR2B channels (Monyer et al., 1992, 1994; Burnashev et al., 1995); theoretically, this should result in relatively restricted Ca influx via the former class of channels. Indeed, in Xenopus oocytes, recombinant mouse NR1–NR2B receptors gated Ca currents that were approximately twice as large as those carried by NR1–NR2A receptors (Kutsuwada et al., 1992; Meguro et al., 1992; Scheetz and Constantine-Paton, 1994); further, NR1–NR2B receptors were approximately twice as sensitive to L-glutamate as NR1–NR2A receptors were. Finally, evoked NMDA receptor-mediated postsynaptic currents in slices from neonatal rats (which express the NR1 and NR2B subunits) were of longer duration relative to those from adult rats that expressed the NR2A subunit in addition to NR1 and NR2B (Kato et al., 1991; Hestrin, 1992).

Therefore, our data suggest that NMDA receptor subunit composition may be modulated by CORT. Chronic MR activation may result in significantly restricted Ca influx in CA1 neurons relative to those from ADX and ADC-HI animals. This regulation is likely to have two components: an altered sensitivity of the heteromeric NMDA receptor to L-glutamate and altered Ca influx as a function of channel kinetics, both of which result in smaller currents being gated by NMDA receptors in ADC-LO neurons. Some investigators, however, have found no differences between NR1–NR2A and NR1–NR2B heteromeric channels in human embryonic kidney (HEK 293) cells, both at the level of single-channel conductances and whole-cell Ca currents (Stern et al., 1992; Anegawa et al., 1995). This may reflect differences in the properties of their expression system relative to neurons. Recent studies also have shown that NMDA receptors may exist as complex heteromeric receptors containing both NR2A and NR2B subunits, a condition that complicates interpretation of our data (Sheng et al., 1994).

The above data show that chronically low daily CORT intake results in low expression levels of mRNAs encoding P and L-type Ca channel subunits, high mRNA expression of the GluR-2 versus GluR-1 subunit, and a significantly higher ratio of NR2A to NR2B mRNA when compared with animals with no circulating CORT or who are exposed to a high daily CORT intake. Our data therefore suggest that expression of these neuronal mRNAs is regulated reciprocally by MR activation relative to no activation or simultaneous activation of MRs and GRs in CA1 neurons. If these changes are, indeed, translated into protein expression, MR activation may result in a tighter regulation of Ca influx, whereas chronic activation of both MRs and GRs may lead to enhanced Ca influx (Table 2). It is possible that alterations in mRNA expression observed in ADC-LO neurons arise in part because of chronic lack of GR activation rather than solely as a functional consequence of chronic MR activation. In the present study we cannot distinguish between these two effects. Lack of receptor occupation in ADX samples leads to levels of mRNA expression similar to the ADC-HI group.

The combined results of PC and univariate statistical analyses are consistent with the above observations (see Fig. 6). Normal physiology of SHAM CA1 neurons is maintained by complex interactions between expression levels of numerous neuronal proteins and mRNAs that encode them. We feel that univariate methods of analysis are insufficient to identify and characterize these interactions, both in normal and altered physiological states. However, multivariate approaches, e.g., PCA, have been used for such purposes before (Bennett et al., 1997; Craig et al., 1997). It is used in the present study as an exploratory tool to identify possible interrelationships among expression of the 10 neuronal genes already described. For this purpose, mRNA expression data first were consolidated mathematically into coefficient-based variables known as PCs (see Materials and Methods). PC1 has a high variance relative to the remaining PCs; each successive PC consecutively explains the remainder of the variability. Higher order PCs are less variable, e.g., in the present analysis PC10 represents the least variability (0.0%) and therefore (arguably) a cohesive cellular response in SHAM neurons. For this reason, we chose to analyze further the variables that comprise PC10.

The ratio of expression levels of GluR-2 and NMDAR-2A relative to GluR-3 and NMDAR-1 was found to be the dominant variable in PC10. This suggests that individual SHAM neurons...
around the mean (see Fig. 6). This scatter is also evident to a lesser extent in ADC-LO neurons, but not in ADC-HI neurons. Further, sole activation of MRs or activation of MRs plus GRs modulates coordinate regulation of these four genes in CA1 neurons differentially relative to the SHAM state and from each other (Fig. 6). Both lack of occupation of MRs and GRs and their concomitant activation have similar effects, whereas sole MR activation results in a markedly different ratio, leading to an inverted U-shaped dose–response relationship between the actions of CORT and this ratio in CA1 neurons. This suggests that AMPA and NMDA receptor subunit mRNAs may be regulated by the extent of MR and GR occupation. The exact mechanism, as well as the consequences of MR and GR activation on functions and properties of AMPA and NMDA receptors, needs to be demonstrated directly by future experiments.

The above results mimic at the molecular level what has been observed previously for regulation of neurotransmitter responses and ionic conductances by glucocorticoids (Joëls and de Kloet, 1994); predominant MR activation and MR plus GR activation usually lead to opposite effects. Further, adrenalectomy without steroid replacement induces effects that are remarkably similar to those seen with MR plus GR activation. Thus, steroid regulation of neuronal membrane and neurotransmitter responses usually appears as a U-shaped dose–response curve. Electrophysiological experiments in tissue from mutant mice lacking a functional GR indicate that interactions between MR and GR proteins may be important for the U shape of the curve (Hesen et al., 1996). Homozygous mutants that only express MRs did not show the characteristic depression of Ca current amplitude seen in wild-type controls in the presence of low to moderate levels of circulating CORT.

In summary, MR activation relative to simultaneous lack or presence of activation of MRs and GRs reciprocally regulates coordinate expression of mRNAs that may play a critical role in Ca permeability and synaptic transmission in CA1 neurons. In this regard, predominant activation of MRs may play a neuroprotective role (Landfield et al., 1981; Gould et al., 1990; Joëls and de Kloet, 1994) relative to conditions in which neither MRs nor GRs are activated or in which both are activated simultaneously in individual CA1 neurons (Elliott and Sapolsky, 1993; Elliott et al., 1993; Stein-Behrens et al., 1994). These molecular events occurring in individual CA1 neurons are in agreement with previous observations that prolonged aberrations in circulating CORT levels are harmful to neuronal integrity in the hippocampus.

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