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Melanocortins regulate the electric waveforms of gymnotiform electric fish

Michael R Markham^{1,2}, Susan J. Allee¹, Anna Goldina¹, and Philip K. Stoddard¹

¹ Department of Biological Sciences, Florida International University, Miami, FL, USA

² Section of Neurobiology, The University of Texas at Austin, Austin, TX, USA

Abstract

The hypothalamic-pituitary-adrenal/interrenal axis couples serotonergic activity in the brain to the peripheral regulators of energy balance and response to stress. The regulation of peripheral systems occurs largely through the release of peptide hormones, especially the melanocortins (adrenocorticotrophic hormone [ACTH] and alpha melanocyte stimulating hormone [α -MSH]), and beta-endorphin. Once in circulation, these peptides regulate a wide range of processes; α -MSH in particular regulates behaviors and physiologies with sexual and social functions. We investigated the role of the HPI and melanocortin peptides in regulation of electric social signals in the gymnotiform electric fish, *Brachyhyppopomus pinnicaudatus*. We found that corticotropin releasing factor, thyrotropin-releasing hormone, and α -MSH, three peptide hormones of the HPI/HPA, increased electric signal waveform amplitude and duration when injected into free-swimming fish. A fourth peptide, a synthetic cyclic- α -MSH analog attenuated the normal circadian and socially-induced EOD enhancements *in vivo*. When applied to the electrogenic cells (electrocytes) *in vitro*, only α -MSH increased the amplitude and duration of the electrocyte discharge similar to the waveform enhancements seen *in vivo*. The cyclic- α -MSH analog had no effect on its own, but blocked or attenuated α -MSH-induced enhancements in the single-cell discharge parameters, demonstrating that this compound functions as a silent antagonist at the electrocyte. Overall, these results strongly suggest that the HPI regulates the EOD communication signal, and demonstrate that circulating melanocortin peptides enhance the electrocyte discharge waveform.

The melanocortin peptides serve a broad range of functions in the central nervous system and the periphery (Hadley and Haskell-Luevano, 1999). This family of peptides includes adrenocorticotrophic hormone (ACTH) and alpha-, beta-, and gamma-melanocyte stimulating hormones (MSHs), which bind with varying affinities to five G-protein coupled melanocortin receptors (MC1R–MC5R). Among the melanocortin peptides, ACTH and α -MSH play prominent roles in the periphery when released from the pituitary into general circulation. While α -MSH does not induce cortisol release in most vertebrates, it nonetheless regulates key peripheral processes including energy homeostasis, melanocyte dispersion, melanin synthesis, exocrine activity, inflammation, sexual receptivity, sexual performance, and sexual signaling (rev. Strand, 1999). Alpha -MSH appears to be a key regulator of vertebrate social signals, such as aggression pheromones in mice (Morgan, Thomas and Cone, 2004) and socially-induced skin darkening response in subordinate Arctic charr (Hoglund et al., 2000).

Correspondence: Michael R. Markham, The University of Texas at Austin, Section of Neurobiology, 1 University Station Stop C0920, Austin, TX 78712, Phone: 512-968-0443, Email: E-mail: markham@mail.utexas.edu.

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Expanding the suite of known social signal functions for melanocortins, we have shown that melanocortin peptides are capable of regulating the electric navigation/communication signals in some electric fish by modulating the activity of the excitable cells that produce the electric signals (Markham and Stoddard, 2005).

The electric organ discharge (EOD) of weakly electric gymnotiform fish is a dual-purpose signal used to navigate and communicate in total darkness. The EOD is a model system particularly well suited for investigating neuroendocrine control of communication signals in response to social stimuli (Stoddard et al., 2006). The EOD is produced by electrocytes, specialized excitable cells in the peripheral electric organ. The EOD waveform of *Brachyhyopomus pinnicaudatus* is a biphasic sinusoidal pulse that varies in amplitude and in the duration of the second phase. Dynamic variation in EOD waveform is readily observed between and among individuals on a circadian rhythm, and in response to social and environmental stimuli (Franchina et al., 2001; Stoddard et al., 2007). Social encounters and environmental stimuli modulate the EOD waveform by altering the membrane biophysics and discharge waveforms of the electrocytes, resulting in either an enhanced or diminished waveform (Franchina et al., 2001; Hagedorn and Zelick, 1989).

We have shown that peripheral injections of serotonin (5-HT) or the melanocortin peptide adrenocorticotrophic hormone (ACTH) modulate the EOD waveform in a manner similar to social encounters in *B. pinnicaudatus*, but only ACTH acts directly on electrocytes to modify their discharge waveforms (Allee et al., 2008; Markham and Stoddard, 2005; Stoddard et al., 2003). Serotonin's lack of direct effect on the electrocytes led us to hypothesize that 5-HT modulates the EOD indirectly, acting centrally to stimulate the release of melanocortin peptides into peripheral circulation. We proposed that the hypothalamic-pituitary-interrenal (HPI) axis is the circuit connecting central serotonin activity and modulation of signals emitted by the electrocytes in the periphery.

In teleost fishes, 5-HT interacts specifically with 5HT_{1A} and 5HT_{2A/2C} receptors in the hypothalamus and/or pituitary to stimulate the HPI axis, inducing the biosynthesis and release of corticotropin-releasing factor (CRF) and thyrotropin-releasing hormone (TRH) (Calogero et al., 1990; Calogero et al., 1993; Contesse et al., 2000; Høglund et al., 2002; Larsen et al., 1998; Van de Kar et al., 2001; Winberg et al., 1997). These peptidergic hormones then stimulate pituitary corticotropes and melanotropes to release into circulation ACTH and α -MSH, respectively (Rotllant et al., 2000). The presence of ACTH in turn stimulates the production and release of glucocorticosteroids (cortisol), adjusting peripheral responses to stressors. Alpha-MSH may act as an alternate corticotrope in some teleosts, but this function for α -MSH is not universal and its taxonomic range is unknown (Lamers et al., 1992; Lamers et al., 1994; Metz et al., 2005). Glucocorticosteroids, the terminal products of the HPI cascade, regulate situation-appropriate behaviors and promote the allocation and distribution of energy required to sustain those behaviors. This connection between brain 5-HT and cortisol production defines the HPI as a key neuroendocrine pathway connecting central brain serotonergic activity and the expression of behaviors within social contexts, particularly in those where dominance and subordination are expressed. Responsiveness of the EOD waveform to social environment is consistent with the hypothesis that 5-HT is involved in producing contextually appropriate changes seen in the EOD waveform via activation of the HPI.

Our objective in this study was to investigate the role of the HPI in regulation of electric social signals. Based on the direct actions of ACTH on the electrocytes, we predicted that α -MSH, the other dominant melanocortin, and CRF and TRH, the upstream hypothalamic secretagogues of endogenous pituitary melanocortins, should also augment the waveform by acting in the pituitary to release melanocortins into the periphery. We tested the ability of these peptide

hormones to modulate the EOD waveform *in vitro*, applied to electrocytes in a dish, and *in vivo* by peripheral injection into free-swimming fish. All three peptides modulated the EOD waveform *in vivo*, but only α -MSH modulated the electrocyte discharge. We also found that a fourth peptide, a synthetic cyclic-MSH analog, inhibits circadian and socially-induced EOD enhancements *in vivo*, and blocks the effects of α -MSH on the electrocyte discharge *in vitro*. We conclude from these data that the HPI mediates EOD enhancements by regulating release of peripheral melanocortins, and that circulating melanocortin peptides contribute to EOD modulation by enhancing the electrocyte discharge.

Materials and Methods

Animals

We used captive-bred adult male *Brachyhypopomus pinnicaudatus* from colonies maintained at Florida International University (Miami, FL) and The University of Texas at Austin. Experiments were approved in advance by the FIU IACUC and the UT IACUC, and complied with the "Principles of Animal Care" publication No. 86-23, revised 1985, of the National Institutes of Health.

Solutions and Reagents

The normal saline for injections and *in vitro* recordings contained (in mM): 114 NaCl, 2 KCl, 4 CaCl₂•2H₂O, 2 MgCl₂•6H₂O, 5 HEPES, 6 glucose; pH to 7.2 with NaOH. We purchased thyrotropin releasing hormone (TRH), corticotropin releasing factor (CRF), and alpha-melanocyte stimulating hormone (α -MSH) from Sigma-Aldrich (St. Louis, MO). We obtained the cyclic- α -MSH analog, [Ac-Cys4,DPhe7,Cys10] α -MSH(4-13)-amide (hereafter cyclic-MSHa), from American Peptide (Sunnyvale, CA). All peptides were dissolved in water at a stock concentration of 100 μ M, stored in single use aliquots at -20 C, then thawed and diluted in saline to working concentrations immediately before use.

EOD Recordings, Injections, and Social Challenges

Our system for recording calibrated EODs from freely swimming fish and procedures for injecting fish are described in detail elsewhere (Stoddard et al., 2003). Fish were placed in an automated measurement tank, 120 × 44 × 44 cm, located in a light- and temperature-controlled room on a 12L:12D light cycle. EODs were amplified and digitized from carbon electrodes at opposite ends of the tank only when the fish passed through or was resting in an unglazed ceramic tube centered between the recording electrodes. EODs were recorded at intervals of ~1 min round the clock. We recorded baseline EODs for ~24 h before performing injections or social challenges.

We prepared the injection solutions to produce the desired dose when injected intramuscularly at 1 μ l g⁻¹ body weight (bw). Saline injections (1 μ l g⁻¹ bw) served as a control condition for handling and injection effects. We selected dosages for each of the pharmacological compounds based on our earlier experiments

All injections of CRF, TRH, and α -MSH were given midday (12:00–15:00). We selected dosages for these compounds based on our previous finding (Stoddard et al., 2003) that EOD response to serotonin saturates at 2.5 nM g⁻¹ body weight (bw). Accordingly, the dosages of CRF and TRH were 2.5 nM g⁻¹ and α -MSH was delivered at a dose of 3 nM g⁻¹ bw. We injected cyclic-MSHa at a much higher dose (10 nM g⁻¹ bw) to allow this compound to compete with the endogenous melanocortin ligands. Fish were quickly netted from the recording tank, injected in the hypaxial muscle, and then returned to the recording tank where EOD recordings continued at ~1 min intervals. Handling time from capture to replacement in the tank was usually less than 30 s.

To determine involvement of endogenous melanocortins with waveform enhancement that accompanies social encounters (Franchina et al., 2001) we initiated social challenges midday (13:00) when EOD parameters are low. We injected the focal fish with cyclic-MSHa or saline control, returned it to its ceramic hiding/recording tube. Within 15 min we added a second male fish from the breeding colony to the same hiding tube, blocking the ends with polyester filter fiber to keep the fish from relocating. After 45 minutes the challenger was removed while the focal fish remained in the recording tank to record ongoing changes in its waveform. Individuals were distinguished by 5–10% differences in body length. Their waveform recordings were distinguished by continuity from the time before the 2nd fish was added. In one of six trials, the waveform amplitude recordings were confounded by the presence of the 2nd fish and the trial was deleted in the final analysis. To determine effects of endogenous melanocortins on the evening rise, we injected focal fish 50–60 min before the normal lights-out time with cyclic-MSHa or saline control, then recorded data for an additional 5 hours. For both social and evening rise effects, each of the 6 male subjects served as its own control, with half of the individuals receiving cyclic-MSHa and half receiving saline. Injection dates were spaced two days apart to allow recovery from treatment. Amplitude recordings for one of the evening rise trials were removed from the final analysis due to excessive missing data points.

Electrophysiology

We recorded the discharges of single electrocytes (μ EODs) and electrocyte action potentials (APs) as described in detail elsewhere (Markham and Stoddard, 2005). Electrocytes of *B. pinnicaudatus* are large disc-shaped cells (approx. 250 μ m width \times 800 μ m diameter) innervated at the apex of the posterior membrane (Hopkins et al., 1990) (Fig. 1). Both the posterior- and anterior-face membranes are excitable and the sequential APs of these two excitable membranes in series produce the μ EOD (Bennett, 1961). The nerve initiates the μ EOD, which is produced by sequential APs: the posterior face spikes first (AP1), followed within 80 μ s by a spike from the non-innervated anterior face (AP2). These two APs sum to produce the biphasic μ EOD recorded differentially across the whole cell. The single-membrane APs are typical monophasic spikes, while the μ EOD is a biphasic sinusoidal discharge that resembles the EOD.

We cut short sections (1.0 – 1.5 cm) of the tail filament, removed the skin on one side to expose the cells of the electric organ, and pinned the tissue in a Sylgard-coated recording chamber (800 μ l total volume) containing normal saline at room temperature ($23 \pm 1^\circ\text{C}$).

We recorded AP1, AP2, and the μ EOD with multi-electrode current clamp procedures detailed elsewhere (Bennett, 1961; Markham and Stoddard, 2005). A depolarizing current step passed through one intracellular pipette initiates the μ EOD. A second intracellular pipette records intracellular potential, and two extracellular pipettes placed within 50 μ m of the anterior and posterior membranes yield recordings of the extracellular anterior and posterior potentials. Offline subtraction of the posterior extracellular record from the intracellular record yields AP1, subtraction of the anterior extracellular record from the intracellular record yields AP2, and subtraction of the posterior extracellular record from the anterior extracellular record yields the μ EOD.

Sharp micropipettes were pulled from thin wall borosilicate glass to resistances of 0.8–1.2 M Ω when filled with 3M KCl. Extracellular pipettes were broken to resistances of 400–600 K Ω filled with normal saline. For intracellular stimulation and recording we used an Axoclamp 900A amplifier (Molecular Devices, Union City, CA) in current clamp mode. Extracellular recordings were made using both channels of a Dagan TEV200A amplifier (Dagan Corp., Minneapolis, MN) in current clamp mode. We controlled the experiment and acquired data at 125 KHz with a Digidata 1440 digitizer and pClamp 10 software (Molecular Devices, Union

City, CA). Data were analyzed with custom-developed code and resident functions in MATLAB (The Mathworks Inc, Natick, MA).

Recording and Perfusion

We recorded only from cells with stable resting potentials ($V_{\text{rest}} < -90$ mV) and stable input resistances. After placement of all electrodes, depolarizing current steps of 6 ms were delivered and the current magnitude was adjusted manually until the current step reliably elicited the μ EOD. We then elicited μ EODs at 60 s intervals for the remainder of each experiment. The recording chamber was perfused at a rate of ~ 5 ml/h with normal saline or with saline containing one of the test compounds. We changed solutions during the interstimulus interval by rapidly perfusing 5 ml of the new solution then slowing perfusion of the new solution to the normal flow rate of 5 ml/h. We first recorded at least 20 min of baseline data in normal saline before changing the solution to one of the test solutions. Saline controls were exposed to continued perfusion of normal saline.

Data Treatment and Analysis

The calibrated EOD recorded *in vivo* is a biphasic sinusoidal pulse (Fig. 1) that varies in amplitude and duration (Franchina and Stoddard, 1998), with duration changing primarily in the second phase (Hopkins et al., 1990). We analyzed *in vivo* EODs as described previously (Stoddard et al., 2003; Stoddard et al., 2007). We measured amplitude peak-to-peak and measured duration of P2 as the repolarization time constant of the second phase (P2) by fitting an exponential curve to the P2 repolarization segment from 50% amplitude to 5% amplitude (hereafter τ_{P2}).

We measured amplitude and τ_{P2} of μ EODs recorded *in vitro* as done for the EOD *in vivo*. Amplitudes of AP1 and AP2 were measured from peak to resting potential. We also measured AP spike width at half-amplitude (half-width), and the delay between AP1 and AP2 (AP1-AP2 delay) as the interval between the peak of AP1 and the peak of AP2.

Changes in the EOD waveform resulting from *in vivo* injections of CRF, TRH, and α -MSH are superimposed upon endogenous circadian cycles in waveform parameters (Stoddard et al., 2007). We therefore mathematically isolated drug-induced changes in these measures as reported previously (Stoddard et al., 2003). This analysis allowed us to extract the changes in EOD amplitude and τ_{P2} that were caused by the challenge trials and not a function of the normal circadian modulations in EOD. Consistent with previous reports, we quantified EOD amplitude and τ_{P2} responses as the peak increase in each waveform parameter within 2h following injection (Stoddard et al., 2003). Statistical analyses and data plotting were performed with MATLAB (Mathworks, Natick, MA) or Prism (Graphpad, La Jolla, CA). Averaged data are reported as mean \pm SEM. All statistical analyses were compared to a significance level set at $p < 0.05$. Experiments with only two treatment conditions were analyzed with Student's t-tests. Experiments with three or more experimental conditions were analyzed first with one-way ANOVA. Significant omnibus ANOVAs were further analyzed with post-hoc pairwise comparisons of each treatment condition against a single control condition (saline carrier control) using Dunnett's pairwise multiple comparison t-test (Dunnett, 1955; 1964) to maintain experiment-wise alpha at 0.05.

For social treatments, changes in EOD waveform following cyclic-MSHa and saline control were normalized to the values at the time of injection and social treatment. For evening treatments, these changes were normalized to values at the time of lights-out. These data were compared at 5-minute intervals from 60–90 min post treatment using repeated measures analysis of variance, then again at the center of this period (75 min post treatment) using a

paired T-test. P-values are one-tailed following a prediction of decline in values for treatment with the antagonist.

Results

Effects of hypothalamic and pituitary peptide hormones on EOD waveform

Intramuscular injection of the hypothalamic releasing hormones CRF (2.5 nM g⁻¹; n=6) and TRH (2.5 nM g⁻¹; n = 8), as well as the melanocortin α -MSH (3 nM g⁻¹; n = 8) elicited marked increases in EOD amplitude compared to saline control injections (n = 7; Fig. 2; $F_{[3, 26]} = 19.28$, $p < 0.0001$). Injection of CRF, TRH, and α -MSH also increased EOD τ_{P2} compared to saline controls (Fig. 2; $F_{[3, 26]} = 19.33$, $p < 0.0001$).

Cyclic-MSHa attenuates social and circadian modulation of the EOD in vivo

We were surprised to find during pilot experiments that cyclic-MSHa, previously shown to function as a potent and long-lasting MCR agonist at the mammalian MC1R (Cody et al., 1985), appeared to suppress rather than enhance endogenous EOD modulation in *B. pinnicaudatus*. We hypothesized that this compound was functioning as an antagonist of endogenous melanocortin activity, and therefore tested whether injections of cyclic-MSHa indeed would suppress circadian and socially-induced EOD enhancements *in vivo*.

Injecting cyclic-MSHa 60 min prior to lights-out caused an initial decrease in EOD amplitude (n = 5) and τ_{P2} (n = 6) compared to saline controls (Fig. 3). Following lights out, fish pretreated with cyclic-MSHa showed a delayed and diminished increase in τ_{P2} compared to saline controls (MANOVA 60–90 min: $F_{[5, 1, 83]} = 6.626$, $P = 0.025$; paired T-test at 75 min: $T = 3.22$, $P = 0.012$). In contrast, EOD amplitude showed only a delayed increase in cyclic-MSHa treated animals with a non-significant mean difference (albeit in the predicted direction) in the 60-90 min comparison window ($F_{[4, 1, 69]} = 2.08$, $P = 0.11$; $T = 1.37$, $P = 0.16$). Cyclic-MSHa significantly reduced socially-induced EOD enhancements. Injecting the focal fish midday with cyclic-MSHa 15 min prior to adding a second fish to the recording tube depressed EOD τ_{P2} relative to saline controls ($F_{[5, 1, 83]} = 11.45$, $P = 0.01$; $T = 4.09$, $P = 0.005$). Cyclic-MSHa depressed EOD amplitude relative to saline, again with effects not as clearly pronounced as for τ_{P2} ($F_{[4, 1, 69]} = 3.50$, $P = 0.07$; $T = 2.40$, $P = 0.03$).

Effects of hypothalamic and pituitary peptide hormones on electrocyte discharges in vitro

Application of CRF (100 nM; n = 6) and TRH (100 nM; n = 6) had no effect on μ EOD waveform whereas α -MSH (100 nM; n = 6) produced rapid and pronounced increases in μ EOD amplitude compared to saline controls (Fig. 4c; $F_{[6, 36]} = 15.23$, $p < 0.0001$). We found a similar pattern for τ_{P2} . Cells treated with α -MSH showed significantly higher τ_{P2} than saline controls whereas cells treated with CRF or TRH showed no difference in τ_{P2} compared to controls (Fig. 4d; $F_{[6, 36]} = 19.59$, $p < 0.0001$).

Cyclic-MSHa antagonized the effects of α -MSH action on the electrocyte discharge. When applied to electrocytes, cyclic-MSHa alone (200 nM; n=6) did not alter the μ EOD amplitude or τ_{P2} (Fig. 4). In contrast, pretreating electrocytes with cyclic-MSHa for 30 min (200 nM; n=6) completely blocked the ability of α -MSH to augment μ EOD τ_{P2} (combined bath of 100 nM α -MSH and 200 nM cyclic-MSHa) (n = 6; Fig. 4b,d). In contrast, cyclic-MSHa reduced but did not completely abolish the effects of α -MSH on μ EOD amplitude. A delayed and attenuated increase in μ EOD amplitude occurred after approximately 20 minutes (n = 6; Fig. 4a,c).

The antagonist effects of cyclic-MSHa (100nM) were quickly and completely reversible on washout. After pretreating cells for 30 min with cyclic-MSHa, switching to a bath solution

containing only α -MSH produced increases in μ EOD amplitude and τ_{P2} comparable in timecourse and magnitude to cells that were not pretreated with cyclic-MSHa (Fig. 4).

The changes in AP timing and waveform underlying the α -MSH-induced increases in μ EOD amplitude and τ_{P2} are the same as we reported for ACTH-induced modulations of the μ EOD (Markham and Stoddard, 2005). As with ACTH, application of α -MSH selectively broadened the width of AP2 but not AP1 (Fig. 5), the change responsible for the increase in P2 amplitude and τ_{P2} seen in the whole-cell μ EOD. The half-width of AP2 increased significantly in the presence of α -MSH compared to saline controls ($t = 5.33$, $df = 12$, $p < 0.001$), whereas AP1 half-width was not affected by ACTH (Fig. 5; unpaired t-test: $p > 0.9$). The enhanced μ EOD P1 amplitude occurs through an increase in the delay between the electrocyte's two action potentials (AP1-AP2 Delay; Fig. 5), unblocking AP1, which is otherwise partially attenuated by temporal overlap with AP2. We found that α -MSH increased AP1-AP2 delay compared to saline controls ($t = 3.47$, $df = 12$, $p < 0.01$).

Discussion

Taken together these results support our hypothesis that the hypothalamic-pituitary portion of the hypothalamic-pituitary-interrenal axis dynamically regulates the EOD waveform of the gymnotiform electric fish *B. pinnicaudatus*. First, intramuscular administration of the hypothalamic hormones corticotropin-releasing factor (CRF) and thyrotropin-releasing hormone (TRH) led to rapid and dramatic enhancements of EOD amplitude and τ_{P2} . The action of these hypothalamic peptides to augment the EOD *in vivo* without affecting electrocytes *in vitro* proves that endogenous downstream target hormones (presumably melanocortins) are capable of augmenting the waveforms. Second, the pituitary melanocortin α -MSH also induced large waveform enhancements, concurring with our previous finding that the melanocortin ACTH is a potent enhancer of EOD waveform both *in vivo* and *in vitro* (Markham and Stoddard, 2005). These data collectively demonstrate that the hormones found at hypothalamic and pituitary levels of the HPI cascade are pharmacologically relevant modulators of the EOD in our fish. Third, injection of cyclic-MSHa *in vivo* attenuated EOD enhancements that normally follow lights-out or interaction with conspecifics. Because cyclic-MSHa acts as a silent antagonist of α -MSH activity *in vitro*, this result supports our hypothesis (Markham and Stoddard, 2005) that endogenous melanocortin action is at least partially responsible for the circadian and socially induced enhancement of the EOD documented previously (Franchina and Stoddard, 1998; Franchina et al., 2001; Silva et al., 1999; Stoddard et al., 2007). We should note that we have never been able to induce in *B. pinnicaudatus* any waveform modulations with cortisol over a wide range of doses, which indicates that the HPI fills a unique role in signaling behavior of some electric fish apart from its role as a regulator of cortisol release.

It is not entirely surprising that cyclic-MSHa functioned as a potent antagonist of α -MSH *in vitro* but did not completely block circadian and socially induced EOD enhancements *in vivo*. Our *in vitro* data strongly suggest that cyclic-MSHa is competitively binding with α -MSH at the electrocyte melanocortin receptor (MCR). Despite pretreating cells with cyclic-MSHa then applying it together with α -MSH at a 2:1 concentration ratio, we still observed a moderate increase in the μ EOD amplitude. Furthermore, cyclic-MSHa appears to bind to electrocyte melanocortin receptors with low affinity. The effects of this compound were eliminated completely within minutes after washout. It is possible then, that during our *in vivo* experiments excretion or metabolism of cyclic-MSHa from a single bolus dose quickly reduced its plasma concentration while concentrations of endogenous α -MSH simultaneously increased in response to lights out or social challenge. Such an account might also explain why we observed an initial drop in EOD amplitude and τ_{P2} during the 60 minutes between the cyclic-MSHa injection and lights out, but not during the shorter 15-min interval between injection of

cyclic-MSHa and the social challenges (Fig. 3). Alternatively, circulating hormones other than melanocortins might contribute to circadian and socially induced EOD enhancements, a possibility given that other 7-transmembrane receptors can induce the cAMP-PKA increase previously shown to drive intracellular changes in electrocyte action potentials of *B. pinnicaudatus* (Markham and Stoddard, 2005).

We have frequently observed that EOD amplitude and τ_{p2} do not change synchronously *in vivo*, suggesting that these waveform parameters are somehow independently regulated. In particular, amplitude typically begins its nocturnal rise shortly after noon while τ_{p2} is still in decline (Franchina and Stoddard, 1998). The present experiments provide our first evidence that these parameters can be dissociated *in vitro*. Exposing electrocytes to a saturating concentration of α -MSH increased both μ EOD amplitude and τ_{p2} with identical time courses indicating that this peptide is responsible for modulating both parameters. However, pretreatment with cyclic-MSHa completely blocked the α -MSH induced τ_{p2} increase, but only delayed and attenuated the increase in μ EOD amplitude. We have shown previously that ACTH increases both amplitude and τ_{p2} via the cyclic-AMP/protein kinase A (PKA) pathway, consistent with the known signal transduction pathways regulated by MCRs. It seems unlikely, therefore, that the differential response of μ EOD amplitude and τ_{p2} would occur through different MCRs or transduction pathways. Instead, we speculate that the cellular mechanisms responsible for setting μ EOD amplitude are sensitive to lower concentrations of PKA than the mechanisms that control τ_{p2} .

Taken together, our results solidify roles for the HPI axis and circulating melanocortins in regulating the EOD waveform. That CRF and TRH modulate the EOD waveform is consistent with our proposal that the HPI mediates serotonergic control of EOD waveform (Allee et al., 2008), since serotonin is a confirmed regulator of these peptide hormones in teleosts (Winberg et al., 1997). Moreover, the EOD is an energetically expensive signal and enhancing waveform parameters in response to social stimuli is particularly costly (Salazar and Stoddard, 2008). Therefore, any stimulus that results in reduction or enhancement of the EOD likely involves the HPI to regulate simultaneously the shape of the electric waveform and the energy made available for adaptive social response.

While the present experiments and our previous reports confirm that the pituitary melanocortins ACTH and α -MSH both can modulate the EOD and μ EOD in a similar manner, we have yet to show which endogenous melanocortins are responsible for EOD modulation *in vivo*, since both are co-released by CRF and TRH in other teleosts (Rotllant et al., 2000). In addition, we have not confirmed which of the melanocortin receptor isoforms is responsible for μ EOD modulation. Preliminary evidence from ongoing cloning and gene expression studies points to MC5R as the functional receptor (Stoddard et al., 2006; J. Tackney & P. Stoddard, unpublished observations). Further, the compound SHU9119 enhances the EOD *in vitro* (Stoddard et al., 2006) and has been shown to act as an agonist at the MC5R and an antagonist at the MC4R of rainbow trout (Haitina et al., 2004). In addition, electrocytes are derived from myocytes during development so MC5R receptors, present in skeletal muscle (Fathi et al., 1995), are likely candidates for expression in electrocytes.

Activation of MC5R in skeletal myocytes upregulates fatty acid oxidation (An et al., 2007). Melanocortin peptides also exert neurotropic effects at the neuromuscular junction, increasing amplitude of compound muscle action potentials by enhancing excitability and neurotransmitter release at the motor endplate (Davies and Smith, 1994; Gonzalez and Strand, 1981). These roles of melanocortins in the skeletal muscle system suggest that they might, under stressful conditions, simultaneously increase energy availability in the muscle while enhancing neurotransmission at the neuromuscular junction, a combined effect that might help

to overcome muscle fatigue. This suggests the interesting question of whether electric fish have harnessed these mechanisms to enhance the μ EOD discharge in myogenic electrocytes.

It is noteworthy, but not unusual, that cyclic-MSHa is reported to be a potent agonist of MC1R receptors (Cody et al., 1985), but functions as a silent antagonist of melanocortin activity at electrocytes. Synthetic MCR ligands often have agonist effects at one receptor subtype while functioning as antagonists for other subtypes as is the case for the widely used α -MSH analog SHU9119 mentioned above.

While the hypothalamic-pituitary-gonadal axis (HPG) is known to be an important mediator of sexual and aggressive signaling, the hypothalamic-pituitary-interrenal/adrenal axis (HPI/HPA) has surprised us with its prominence. Melanocortins, intermediate hormones in the HPI/HPA axis, are known to regulate (1) synthesis and release of aggression-mediating preputial pheromones in nocturnal rodents (Morgan, Thomas and Cone, 2004; Morgan, Thomas, Ma et al., 2004; Nowell and Wouters, 1975), (2) skin pigmentation in teleosts, frogs, and lizards (e.g., Castrucci et al., 1984; Hoglund et al., 2000), and now (3) electric waveform characters in a gymnotiform electric fish. Given the morphologic and taxonomic variation in these signal structures, it appears that evolution has persistently reused melanocortins for control of social signals. It will be interesting to explore common life history characteristics that favor use of melanocortins for control of social signaling. Clues may be found in their pleiotropic effects, in particular their deep involvement in the physiology of energy management.

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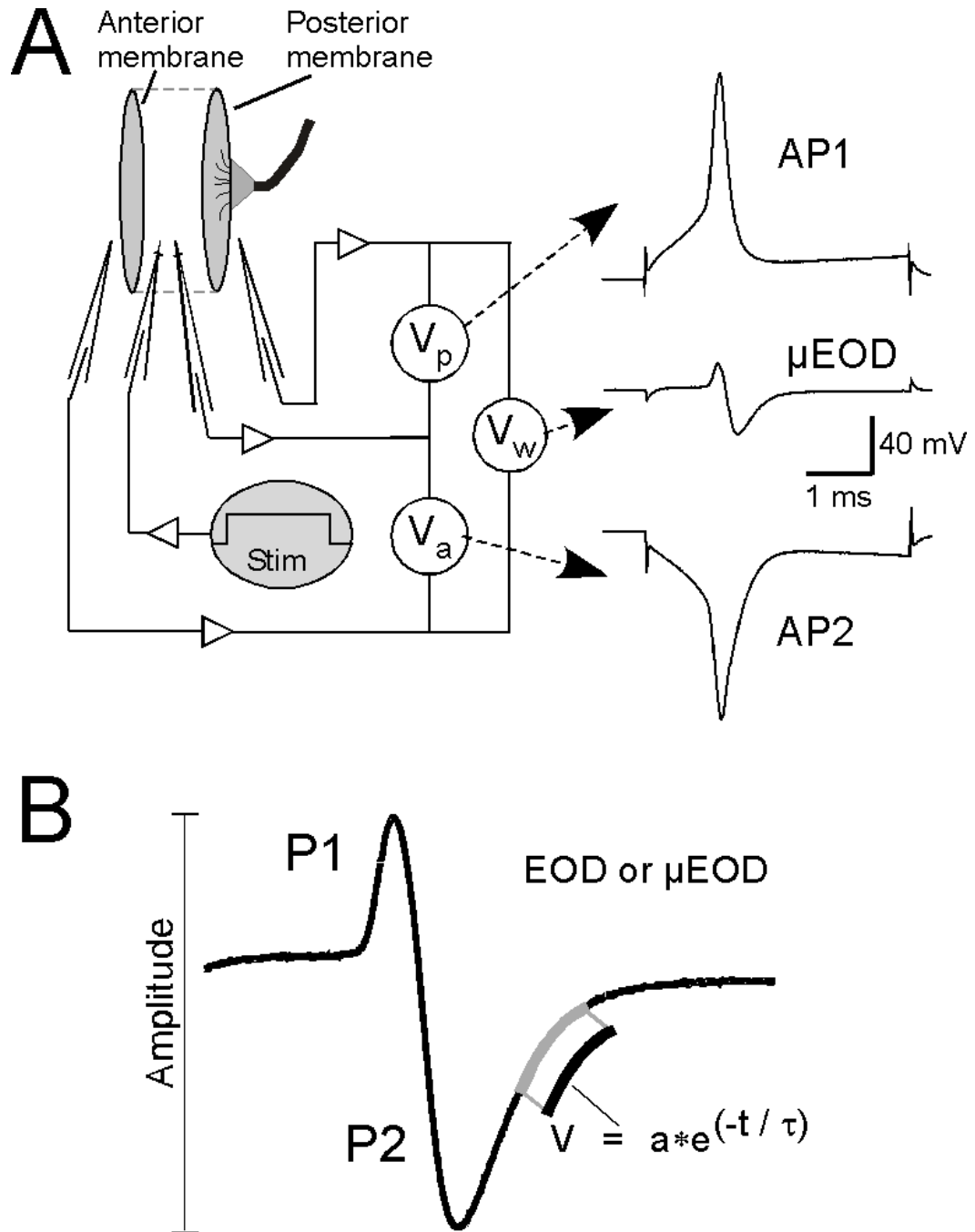


Figure 1.

Recording of electrocyte action potentials and μ EODs and analysis of EOD waveform parameters. **A)** We delivered intracellular current steps to electrocytes *in vitro* to elicit μ EODs while simultaneously recording voltage from one intracellular and two extracellular micropipettes. Sequential action potentials on the innervated posterior face (AP1) and the noninnervated anterior face (AP2) of the electrocyte sum to produce the biphasic μ EOD. Potentials are recorded from one intracellular micropipette and two extracellular pipettes. Offline subtraction of the posterior extracellular record from the intracellular record yields the posterior-membrane voltage (V_p) and AP1, the posterior-face action potential. Subtraction of the anterior extracellular record from the intracellular record yields the anterior-membrane

voltage (V_a) and AP2, the anterior-face action potential. Subtraction of the posterior extracellular record from the anterior extracellular record yields the whole-cell voltage (V_w) and the μ EOD. B) EOD and μ EOD amplitude was measured peak-to-peak. The time constant of the second phase (P2) repolarization, τ_{P2} , was estimated by exponential fit to the repolarization segment of P2.

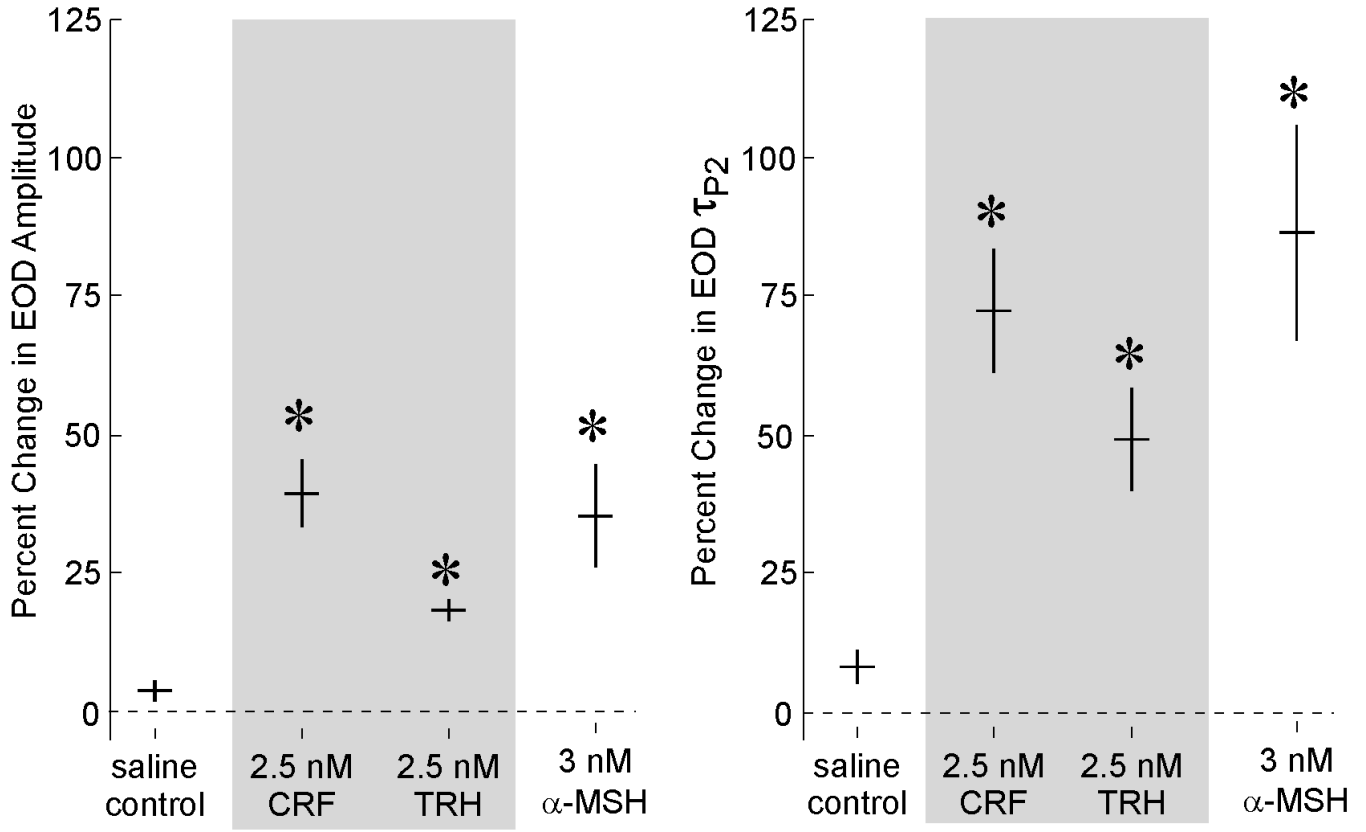


Figure 2. Injections of CRF, TRH, and α -MSH enhance EOD waveforms *in vivo*. Injections of saline caused little or no change in amplitude or τ_{P2} , whereas injections of CRF, TRH, and α -MSH caused clear increases in EOD amplitude (left panel) and τ_{P2} (right panel). Horizontal bars represent means and vertical bars represent SEM. Asterisks indicate conditions statistically different from saline ($p < 0.01$),

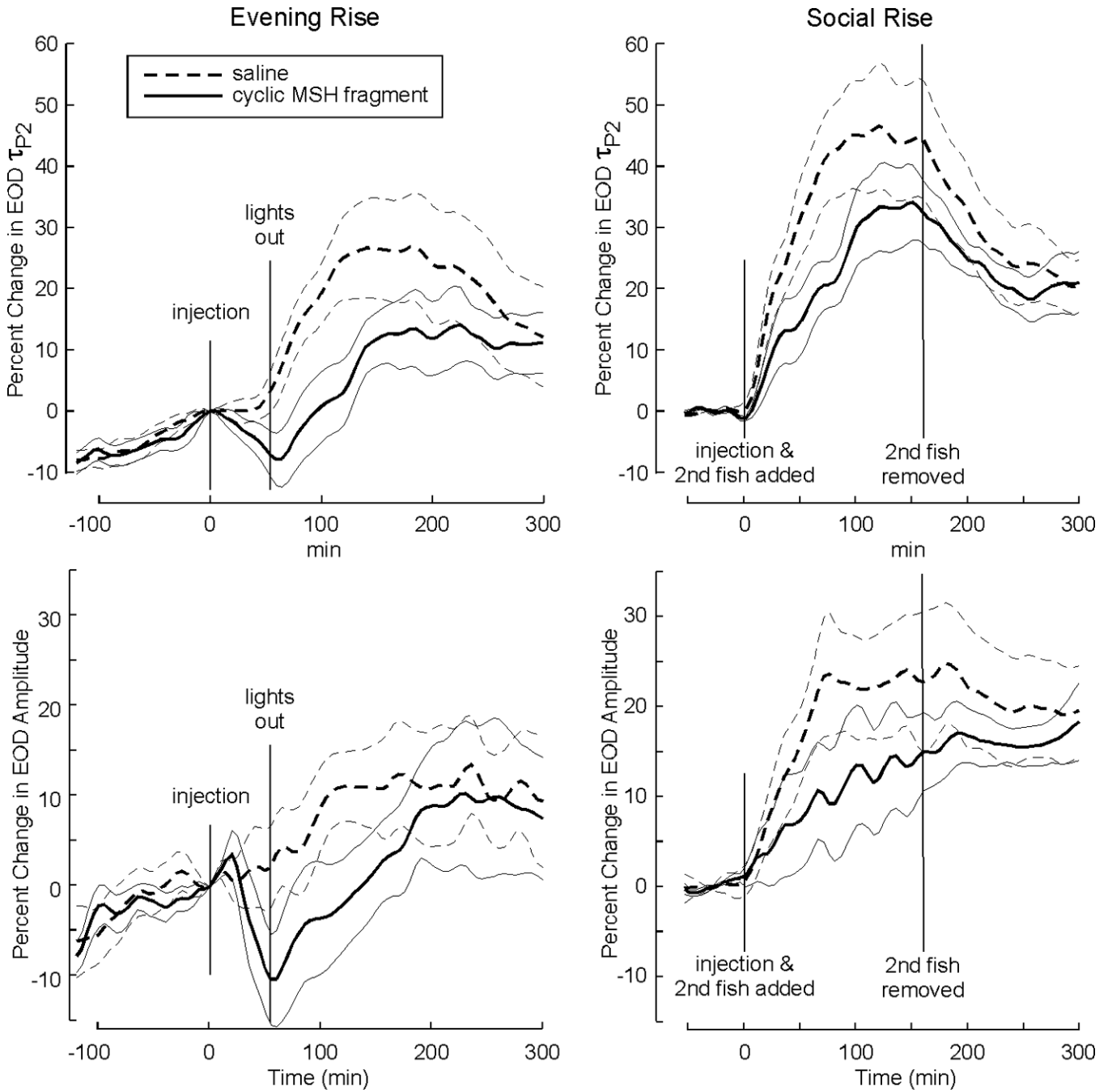


Figure 3. Injections of cyclic-MSHa *in vivo* attenuate circadian and socially-induced EOD waveform enhancements. Bold lines represent means and dashed lines 95% confidence intervals. Left panels show that injection of cyclic-MSHa 60 min before lights out delays and attenuates the circadian rise in EOD τ_{P2} (top) and amplitude (bottom). Right panels show that cyclic-MSHa injections just before addition of a conspecific delay and attenuate the socially- induced increases in EOD τ_{P2} (top) and amplitude (bottom). All records are rescaled to the magnitude at the time of injection. Over the entire 350 min record, the parameter values show an overall rise rather than a return to baseline: the rise reflects the endogenous circadian rhythm (Stoddard et al., 2007) in the parameter after the passing of the treatment effect.

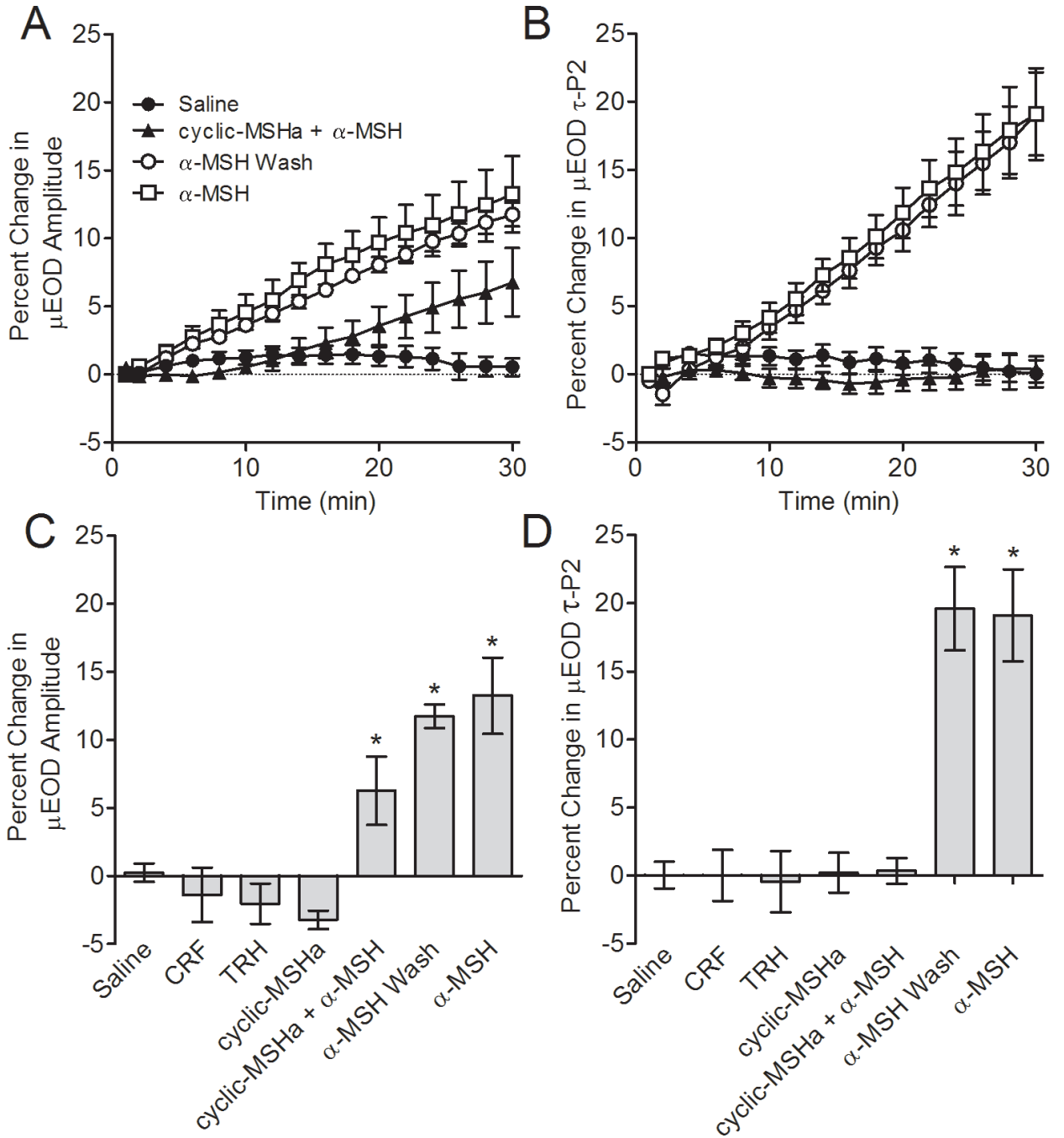


Figure 4. Effects of peptide and peptidergic factors on the μ EOD *in vitro*. Symbols or bars represent means and error bars show SEM. **A)** Time course of μ EOD amplitude changes in single electrocytes. Alpha-MSH caused rapid μ EOD amplitude increases. Cells pretreated with cyclic-MSHa showed a delayed and attenuated μ EOD amplitude response in pretreated cells. Following washout of cyclic-MSHa, application of α -MSH rapidly increased μ EOD amplitude. **B)** Time course of μ EOD τ_{P2} changes. Alpha-MSH caused rapid and large increases in μ EOD τ_{P2} . Cells pretreated with cyclic-MSHa showed no change in τ_{P2} in response to α -MSH application. Following washout of cyclic-MSHa, application of α -MSH increases μ EOD amplitude and τ_{P2} in a manner indistinguishable from cells that were not pretreated with cyclic-

MSHa. **C)** Percent change in μ EOD amplitude after 30 minutes exposure to experimental conditions. Responses to TRH, CRF and cyclic-MSHa were indistinguishable from saline controls after 30 minutes. In contrast, α -MSH caused a large increase in μ EOD amplitude. Cells pretreated with cyclic-MSHa showed no change in τ_{p2} in response to α -MSH application, whereas α -MSH produced an attenuated μ EOD amplitude response in pretreated cells. Following washout of cyclic-MSHa, application of α -MSH increased μ EOD amplitude. **D)** Percent change in μ EOD τ_{p2} after 30 minutes exposure to experimental conditions. Responses to TRH, CRF and cyclic-MSHa were indistinguishable from saline controls after 30 minutes. α -MSH caused a large increase in μ EOD τ_{p2} , whereas cells pretreated with cyclic-MSHa showed no change in τ_{p2} in response to α -MSH application. Following washout of cyclic-MSHa, application of α -MSH increased μ EOD τ_{p2} .

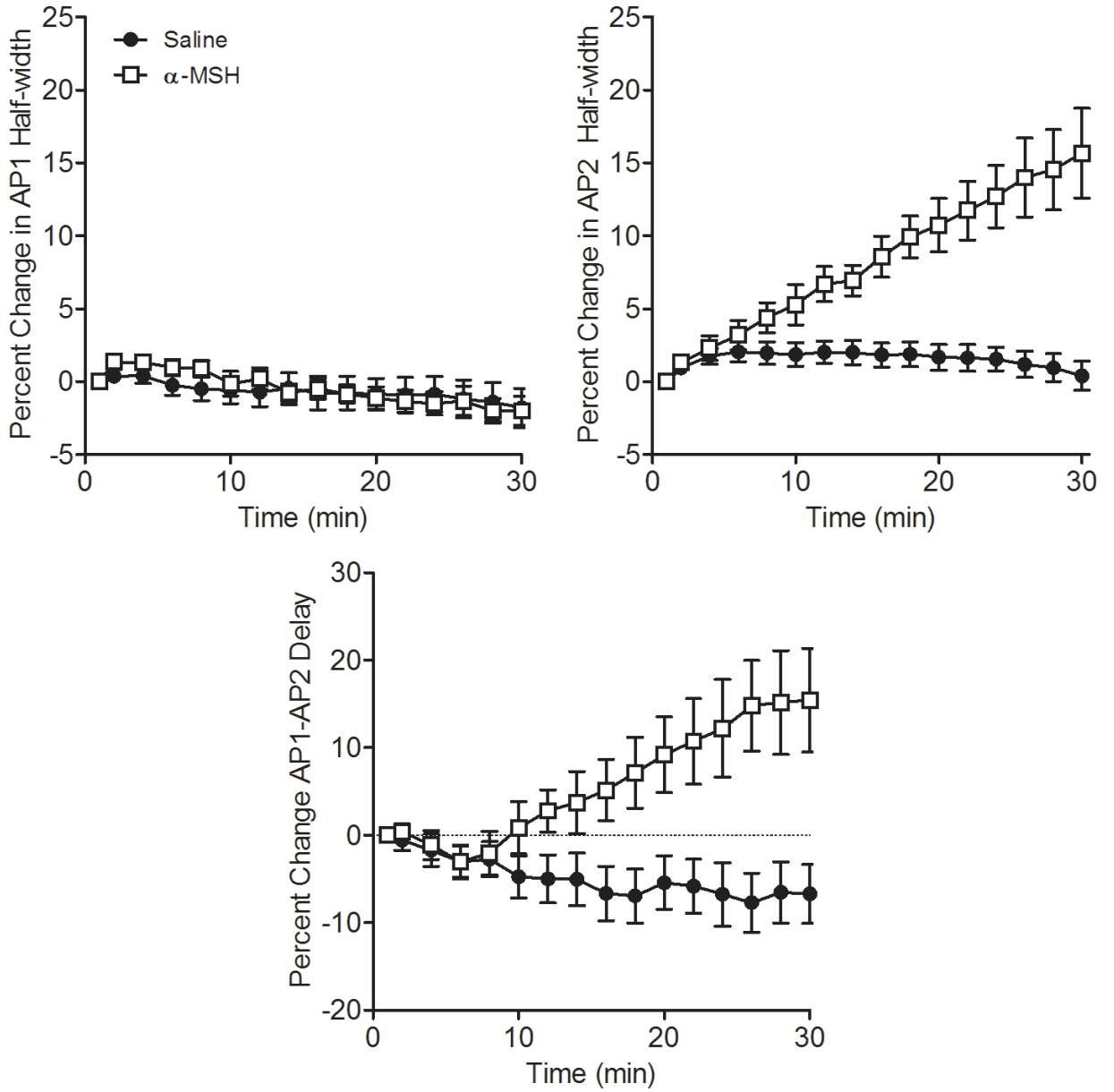


Figure 5. Alpha-MSH selectively broadens the electrocyte AP2 waveform and increases the temporal offset of AP1 and AP2. The half width of AP1 spikes remained constant in electrocytes treated with α -MSH or saline with no differences between groups even after 30 minutes. Symbols represent means and error bars show SEM.