Stress elevates corticotropin-releasing factor (CRF) and CRF-binding protein mRNA levels in rainbow trout (Oncorhynchus mykiss)

C Doyon, V L Trudeau and T W Moon

Centre for Advanced Research in Environmental Genomics (CAREG), Department of Biology, University of Ottawa, Ottawa, Ontario K1N 6N5, Canada

Abstract

The objectives of this study were to characterize rainbow trout (Oncorhynchus mykiss) corticotropin-releasing factor (CRF)-binding protein (CRF-BP) cDNA and to examine the variations in CRF-BP and CRF mRNA levels in response to different intensities of stress. Trout were physically disturbed by a single or three consecutive periods of chasing until exhaustion followed by 2 h of recovery. The pituitary CRF-BP and preoptic area CRF1 mRNA contents were significantly increased only after repeated chasing events. Physical disturbance increased plasma cortisol levels with the largest change occurring in the group of trout that were exposed to repeated chasing events. Trout were also individually isolated in 120 l tanks or confined to 1·5 l boxes for 4, 24 or 72 h. CRF-BP mRNA levels in confined fish were greater than those of isolated fish at 72 h although there were no differences compared with the control group. CRF1 mRNA levels in the preoptic area were greater and remained elevated for a longer period in confined compared with isolated trout. Isolation led to a transient increase in plasma cortisol levels, but the higher cortisol values developed in the confined fish suggest that this treatment was more stressful than isolation. These results demonstrate that the intensity and duration of stress are important factors regulating CRF and CRF-BP mRNA levels in rainbow trout. We hypothesize that pituitary CRF-BP is involved in regulating the activity of the stress axis, possibly by reducing access to CRF1 receptors in the corticotropes.

Journal of Endocrinology (2005) 186, 123–130

Introduction

The primary stress response involves the activation of hypothalamic neurons producing corticotropin-releasing factor (CRF), an initial step in the cascade that leads to the synthesis and release of glucocorticoids (Wendelaar Bonga 1997). The activity of the stress axis can be modulated at several levels including pituitary changes in the expression of the CRF1 receptor or CRF-binding protein (CRF-BP). The function of CRF-BP is still unclear although studies do support an inhibitory role for CRF-BP on CRF action (Seasholtz et al. 2002). Rodent CRF-BP blocks CRF-induced adrenocorticotropic hormone (ACTH) release from pituitary cells in vitro (Potter et al. 1991, Cortright et al. 1995). Also, CRF released by the human placenta rises exponentially during pregnancy (McLean et al. 1995), but most plasma CRF is bound to its binding protein and therefore unable to bind to its receptors (Zhao et al. 1997). The binding of CRF to plasma CRF-BP would prevent inappropriate stimulation of the stress axis by placental CRF (Petraglia et al. 1993). CRF-BP is present in corticotropes of the anterior pituitary of rodents, implying that it could modulate the action of receptor-bound CRF (Seasholtz et al. 2002). In a line of transgenic mice overexpressing CRF-BP in the anterior pituitary only, it was hypothesized that the elevated hypothalamic CRF expression observed may have a role to maintain normal levels of free CRF and normal activity of the stress axis (Burrows et al. 1998). The levels of CRF-BP mRNA in the rat pituitary were increased by stress and decreased by adrenalectomy, suggesting a possible involvement of pituitary CRF-BP in regulating the activity of the stress axis (McClenen et al. 1998). Recent evidence supports a function for CRF-BP other than simply inhibiting CRF action in the central nervous system of rodents. The CRF-BP ligand inhibitor, CRF6–33, activated some neurons that express CRF-BP but not CRF receptors, implicating CRF-BP in signalling by CRF-related peptides (Chan et al. 2000). Also, CRF required CRF-BP to potentiate N-methyl-D-aspartate (NMDA) receptor-mediated synaptic transmission in dopamine neurons (Ungless et al. 2003). Fish may represent a good model to study the regulation and role of pituitary CRF-BP because of the direct innervation of corticotropes within the pituitary pars distalis by CRF-containing nerve fibers (Yulis & Lederis 1987, Olivereau & Olivereau 1988). Although CRF-BP
was not detected in the plasma of tilapia (Pepels et al. 2004), CRF-BP-like immunoreactivity was localized in the pituitary gland of common carp (Huising et al. 2004). The objectives of this study were therefore to characterize rainbow trout CRF-BP cDNA and to examine the effects of different intensities of stress on CRF-BP and CRF1 mRNA levels. We recently characterized the cDNA encoding two paralogous forms of rainbow trout (Oncorhynchus mykiss) CRF, CRF1 and CRF2, and we demonstrated that social stress increased CRF1 mRNA levels in the preoptic area of the trout brain (Doyon et al. 2003). CRF1 mRNA content was examined in the preoptic area, which contains the highest levels of brain CRF mRNA (Doyon et al. 2003). CRF-BP mRNA levels were measured in the pituitary gland, one potential site for CRF-BP regulation of the stress axis activity. Various intensities of stress were achieved by physically disturbing trout with single or repeated chasing events and by isolating trout individually in large tanks or small confinement boxes for 4, 24 or 72 h. CRF-BP and CRF1 mRNA levels were measured by a ribonuclease protection assay (RPA). Based on results from previous studies (Doyon et al. 2003), we hypothesized that stress would increase pituitary CRF-BP and preoptic area CRF1 mRNA contents.

Materials and Methods

Animals

Immature female rainbow trout, Oncorhynchus mykiss, were obtained from Linwood Acres Trout Farm (Campbellcroft, ON, Canada). They were acclimated to tanks of well-aerated dechloraminated City of Ottawa tap water for at least 6 weeks in 1275 l fiberglass tanks (Campbellcroft, ON, Canada). They were acclimated to were obtained from Linwood Acres Trout Farm in December 2002 to March 2003 and used a total of 40 trout weighing between 35 and 97 g (63·2 ± 2·3 g). Over this period, 14 trout were netted quickly at a rate of two to four fish per sampling day from a 1275 l holding tank that had not been disturbed by netting for at least 2 days. These fish were immediately anaesthetized with high sequence identity between the cDNA encoding two paralogous forms of rainbow trout (Oncorhynchus mykiss) CRF, CRF1 and CRF2, and we demonstrated that social stress increased CRF1 mRNA levels in the preoptic area of the trout brain (Doyon et al. 2003). CRF1 mRNA content was examined in the preoptic area, which contains the highest levels of brain CRF mRNA (Doyon et al. 2003). CRF-BP mRNA levels were measured in the pituitary gland, one potential site for CRF-BP regulation of the stress axis activity. Various intensities of stress were achieved by physically disturbing trout with single or repeated chasing events and by isolating trout individually in large tanks or small confinement boxes for 4, 24 or 72 h. CRF-BP and CRF1 mRNA levels were measured by a ribonuclease protection assay (RPA). Based on results from previous studies (Doyon et al. 2003), we hypothesized that stress would increase pituitary CRF-BP and preoptic area CRF1 mRNA contents.

Hormone measurements

Fish were netted quickly and terminally anaesthetized in 150 mg l⁻¹ benzocaine (ethyl p-aminobenzoate; Sigma-Aldrich, St Louis, MO, USA). All fish were anaesthetized between 2 and 4 p.m. to avoid possible daily rhythms of plasma cortisol. Blood was collected in heparinized syringes by caudal puncture and centrifuged. The plasma was removed and stored at −80 °C for later analysis of cortisol levels using a commercially available RIA kit (Cortisol125I RIA Kit; ICN Diagnostics, Costa Mesa, CA, USA).

Amplification of CRF-BP-related cDNA fragments and sequencing

Poly(A)⁺ RNA was extracted from the preoptic area of the trout brain using the Straight As mRNA Isolation System (Novagen, Madison, WI, USA). Oligonucleotide primers (Invitrogen, Burlington, ON, Canada) were designed on the basis of high sequence identity between the Xenopus laevis (Brown et al. 1996) and the Japanese pufferfish CRF-BP sequences (Table 1). First-strand cDNA was synthesized from 400 ng poly(A)⁺ RNA with Superscript II (Invitrogen) and oligo(dT). Amplification was carried out using Taq polymerase (Invitrogen) and a Mastercycler.
Gradient (Eppendorf; Mississauga, ON, Canada) with the following program: 5 min denaturation at 94 °C, followed by 35 cycles of denaturation at 94 °C for 30 s, annealing at 55 °C for 30 s and extension at 72 °C for 1 min. After the last cycle, further extension was performed at 72 °C for 10 min. The PCR reaction contained 4 µl cDNA in a 50 µl PCR reaction.

The 3' and 5' ends of trout CRF-BP were obtained by rapid amplification of cDNA ends (RACE; Invitrogen). For 3' RACE, first-strand cDNA was synthesized from 400 ng poly(A)+ RNA with Superscript II and the 3' RACE adapter primer (Invitrogen). CRF-BP-related cDNA fragments were amplified using the sense primer of the initial cloning procedure (Table 1), the abridged universal amplification primer (AUAP; Invitrogen) and the program described above. For 5' RACE, total RNA was extracted from the preoptic area using TRIzol reagent (Invitrogen). First-strand cDNA was synthesized from 700 ng total RNA with Superscript II and the antisense (Invitrogen). CRF-BP-related cDNA was purified after each step of the 5' primer of the initial cloning procedure (Table 1). The abridged universal amplification primer (AUAP; Invitrogen) and the antisense primer of the initial cloning procedure (Table 1), the program described above. For 5' RACE, total RNA was extracted from the preoptic area using TRIzol reagent (Invitrogen). First-strand cDNA was synthesized from 700 ng total RNA with Superscript II and the antisense primer of the initial cloning procedure (Table 1). The cDNA was purified after each step of the 5' RACE with the QIAquick PCR Purification Kit (Qiagen). Tailing of the 5' end of the cDNA was performed using dCTP and terminal transferase (Roche, Laval, QC, Canada). The first PCR reaction was performed using the antisense primer of the initial cloning procedure (Table 1). Two semi-nested PCR reactions were performed using gene-specific primers (GSP1 and GSP2; Table 1) that were based on initial sequencing results.

Amplified CRF-BP products were extracted from a 1% agarose gel with the QIAquick Gel Extraction Kit (Qiagen), ligated into the pCR II-TOPO cloning vector (TOPO TA cloning kit; Invitrogen), and transformed into *Escherichia coli* competent cells (One Shot TOP10 Chemically competent cells; Invitrogen). Single colonies were cultured and plasmids were recovered with Wizard Plus (Promega, Madison, WI, USA). All procedures were carried out according to the manufacturer’s instructions. Both strands of at least three different cloned inserts were sequenced (Canadian Molecular Research Services, Ottawa, ON, Canada) to obtain the reported consensus sequence. Nucleotide sequences were submitted to BLAST (http://www.ncbi.nlm.nih.gov/blast/blast.cgi; Altschul et al. 1997) for comparison with sequences present in the GenBank nucleotide sequence database.

### RPA

Total RNA was extracted from the preoptic area of the brain (which also included part of the anterior hypothalamus) and pituitary glands using TRizol reagent. Levels of CRF1 and β-actin mRNA in the preoptic area were analyzed using the RPA (RPA III; Ambion, Austin, TX, USA) described previously (Doyon et al. 2003). Levels of CRF-BP and β-actin mRNA in individual pituitaries were determined using a separate RPA (RPA III). The DNA templates for the preparation of antisense RNA probes were generated by PCR using primers with incorporated SP6 and T7 phage polymerase promoters on the sense and antisense primers, respectively (Table 1). DNA templates were amplified using the following program: 3 min denaturing at 94 °C, then 35 cycles of denaturation at 94 °C for 30 s, annealing at 58 °C for 30 s and extension at 72 °C for 1 min. After the last cycle, further extension was performed at 72 °C for 7 min. Amplified products were extracted from a 1% agarose gel with the QIAquick Gel Extraction Kit. Antisense RNA probes were prepared using the MAXIscript In vitro Transcription Kit (Ambion) and approximately 100 ng template DNA.

For CRF-BP, probes were prepared by incubating the DNA at 37 °C for 30 min with 40 µCi [α-32P]CTP at 800 Ci mmol⁻¹ (Amersham Bioscience, Baie d’Urfe, QC, Canada) in the presence of 0.05 mM unlabeled CTP. To limit the hybridization signal of β-actin, a highly expressed gene, reduced-specific-activity probes were prepared by incubating the DNA at 37 °C for 1 h with 40 µCi [α-32P]CTP at 800 Ci mmol⁻¹ (Amersham Bioscience) in the presence of 0.5 mM unlabeled CTP. The transcription products were incubated for 30 min at 37 °C with 5 units of RNase-free DNase 1 (Promega). Full-length probes were purified from a 5% acrylamide/bis-acrylamide (19:1)/8 M urea gel. Approximately 8 × 10⁶ c.p.m. probe was hybridized overnight at 42 °C to 10 µg total RNA. Non-hybridized transcripts were digested at 37 °C for 1 h.

| Table 1 Primer sequences used for cloning and amplification of template DNA for the RPA |
|---------------------------------|---------------------------------|
| **5’ Primer**                   | **Position**                    | **3’ Primer**                   | **Position** |
| Initial cloning GGTGTTTGATGGCTGGGTAA 3' to 5' | 395–414 | AGCTGCAGTTCCTGTGCTG 5' to 3' | 603–681 |
| 5’ RACE GSP1 ATTAGGTGACACTATAGAAGVC 5’ to 3’ | 615–630 | ATCATCTCCCAGACAC AGCTACAGGGAACATT 3’ to 5’ | 830–845 |
| RPA CRF-BP GATCCATATAGCGTTCCATCAA 5’ to 3’ | 274–292 | ATATACGACTCATACTAGGGAACATT 3’ to 5’ | 593–611 |
| RPA β-actin GGACACGGCATCGTCACC 5’ to 3’ | 593–611 | TAATACGACTCATACTAGGGAACATT 3’ to 5’ | 593–611 |

Nucleotides in italics represent the promoter sequences for T7 and SP6 RNA polymerases. GSP, gene-specific primer; RACE, rapid amplification of cDNA ends.
30 min with approximately 0.2 units RNase A and 7.5 units RNase T1 (Ambion). Protected fragments were resolved on a 5% acrylamide/bis-acrylamide (19:1)/8 M urea gel. An Image Screen-K was exposed for 18 h to the dried gel and the optical density of the bands was analyzed with Quantity One version 4.1.1 (Bio-Rad Laboratories, Mississauga, ON, Canada). Results were expressed in counts per mm², the arbitrary volume unit of Quantity One. The RPA was optimized by verifying the linearity of the relation between quantified mRNA and the amount of total RNA (Fig. 1). The standard curves for the optimization of the RPA demonstrate linearity and thus the levels of /afii9826-actin (Fig. 1B) and CRF-BP (Fig. 1C) mRNA can be quantified with this technique.

Statistics

Results are presented as mean values ± 1 S.E.M. For the isolation experiment, statistical differences between treatments and control were determined by one-way ANOVA. CRF1 mRNA values for the confined fish were transformed using a natural logarithm to meet the assumption of normality. For the physical-disturbance experiment, statistical differences between treatments were determined by one-way ANOVA. All parametric ANOVAs were followed by Tukey’s multiple-comparison tests. For both experiments, data on cortisol content did not meet the assumption of normality so differences between treatments were determined by Kruskal–Wallis one-way ANOVA on ranks followed by Dunn’s multiple-comparison test. All statistical analyses used SigmaStat (v2.0) software (SPSS, Chicago, IL, USA).

Results

Sequencing and characterization of trout CRF-BP

A complete coding sequence was obtained for the cDNA encoding rainbow trout CRF-BP (GenBank accession no. AY363677). The nucleotide sequence of CRF-BP (1609 nt) is composed of a 62 bp 5’-untranslated region (UTR), a 963 bp open reading frame (ORF) and a 583 bp 3’-UTR that contains two polyadenylation signals (AATAAA). The deduced amino acid sequence (521 amino acids) contains a putative signal peptide of 21 amino acids, as predicted using the SignalP program v2.0 (http://www.cbs.dtu.dk/services/SignalP-2.0/; Nielsen et al. 1997, Nielsen & Krogh 1998). The putative mature peptide contains 10 cysteine residues that are highly conserved throughout vertebrates (Huising et al. 2004) and are likely involved in the formation of five disulfide bonds (trout C60–C81, C104–C141, C183–C205, C239–C264, C277–C318).

Physical disturbance

The levels of plasma cortisol in controls (9.30 ± 0.66 ng/ml) were characteristic of unstressed trout (Fig. 2A; Barton & Iwama 1991). One period of physical disturbance led to a significant increase in plasma cortisol levels; levels were further increased after three consecutive chasing events (Fig. 2A). One chasing event was not sufficient to increase CRF1 mRNA content despite elevated cortisol levels (Fig. 2B). However, three chasing events significantly increased CRF1 mRNA levels compared with both the control and a single chasing event (Fig. 2B). A 2-fold increase in pituitary CRF-BP mRNA content was obtained after three chasing events (Fig. 2C).

Isolation

The levels of plasma cortisol in controls (1.38 ± 1.02 ng/ml) were characteristic of unstressed trout (Fig. 3A; Barton
Plasma cortisol levels in both treatments were significantly elevated compared with the control at 4 and 24 h of isolation but were back to control values at 72 h (Fig. 3A). The elevation in plasma cortisol at 4 h was significantly different from other treatments (P<0.05; n=12–16 per group).

Figure 2 Effects of single and repeated periods of physical disturbance on the levels of (A) plasma cortisol, (B) preoptic area CRF1 mRNA and (C) pituitary CRF-BP mRNA in juvenile female rainbow trout. Measurements were carried out after a single (Chasing) or three consecutive (Repeated) periods of physical disturbance followed by a 2-h recovery period in isolation. β-actin was used as a control to correct for the amount of total RNA used in the RPA. Treatments that do not share a common letter are significantly different from each other (P<0.05; n=12–16 per group).

Figure 3 Time course of changes in the levels of (A) plasma cortisol, (B) preoptic area CRF1 mRNA and (C) pituitary CRF-BP mRNA in juvenile female rainbow trout that were individually isolated in 120 l tanks (Isolation) or confined to 1.5 l boxes (Confinement). β-actin was used as a control to correct for the amount of total RNA used in the RPA. Treatments that do not share a common letter within isolation or confinement are significantly different from each other (P<0.05). Asterisks show statistical difference compared with control, whereas horizontal bars show difference between isolation and confinement (P<0.05; n=12–14 per group).
greater in trout confined in 1.5 l boxes compared with trout isolated in a 120 l tank. Although cortisol levels were highest at 4 h of isolation, CRF1 mRNA levels were different from control values in the confined fish only (Fig. 3B). At 24 h of isolation, both groups had elevated CRF1 mRNA levels compared with both the control and 4 h values. CRF1 mRNA levels remained elevated at 72 h in confined fish despite low cortisol values, whereas levels of CRF1 mRNA were back to control values in isolated fish. The levels of CRF-BP mRNA in the pituitary gland also showed trends that were similar to those of CRF1 mRNA (Fig. 3C). The difference in the levels of CRF-BP mRNA between isolation and confinement groups was only significant at 72 h although there was no difference compared with the control group.

Discussion

We report the complete coding sequence for the precursor cDNA of rainbow trout CRF-BP. The predicted amino acid sequence of trout CRF-BP shares 75% identity with the recently reported sequence of the common carp, *Cyprinus carpio* (Huising *et al.* 2004), and approximately 60% identity with mammalian sequences (Potter *et al.* 1991). Two distinct forms of the cDNA encoding CRF-BP were reported for the common carp, a tetraploid species (Huising *et al.* 2004). As the rainbow trout is also tetraploid, more than one form of CRF-BP would be predicted, and preliminary sequencing information of another trout CRF-BP (NJ Bernier, personal communication) appears to substantiate this prediction. The second CRF-BP band on the RPA is consistent with the presence of a second form of CRF-BP in trout (Fig. 1A).

The repeated physical-disturbance stress that led to the highest cortisol levels in this study increased CRF-BP mRNA levels in the trout pituitary gland. Trout exposed to three consecutive chasing events had a 2-fold increase in pituitary CRF-BP mRNA levels. This novel finding contrasts with a study in rats showing that exposure to repeated footshock stress increased CRF mRNA levels in the paraventricular nucleus, whereas a single footshock was not sufficient to produce this effect (Deutch *et al.* 1987, Imaki *et al.* 1991). However, 5 min of immobilization stress was sufficient to increase CRF heteronuclear RNA levels in the rat paraventricular nucleus, implicating activation of CRF transcription (Imaki *et al.* 1995, 1996). Despite a rapid increase in CRF heteronuclear RNA, elevated CRF mRNA levels were detectable only 60–120 min after the beginning of the stress (Imaki *et al.* 1992, 1995).

Placing trout individually in either 120 l tanks or 1.5 l boxes led to a transient increase in plasma cortisol levels despite continuous exposure to the stressor (4–72 h). Although several studies report transient increases in plasma cortisol with continuous confinement of rainbow trout (Pottinger & Moran 1993, Gamperl *et al.* 1994), this is the first study to examine these changes in plasma cortisol in parallel with those of CRF and CRF-BP mRNA. Also, isolation in 120 l tanks led to a transient increase in CRF1 mRNA levels in the preoptic area of the trout brain despite sustained stimulus. Although part of this increase in CRF1 mRNA may result from the initial netting and handling stress, this result implies a habituation to isolation. Such adjustments may contribute to maintaining the responsiveness of the hypothalamic-pituitary-interrenal (HPI) axis to a novel stressor in situations of chronic stress, as has been observed in chronically stressed mammals (Aguilera 1994).

Plasma cortisol levels were higher in confined fish (1.5 l) compared with fish isolated in a large water volume (120 l). Thus, confinement was more stressful than isolation, and this greater stress intensity was reflected in CRF1 mRNA levels. The magnitude and persistence of the elevation in CRF1 mRNA was greater in confined fish, suggesting a correlation between CRF mRNA levels and...
the intensity of stress in trout. Similarly, brief low-water exposure did not affect arginine vasotocin (AVT) mRNA, whereas acute confinement, which elicited a greater cortisol response, significantly increased AVT transcript levels in parvocellular neurons of the preoptic nucleus of the rainbow trout brain (Gilchriest et al. 2000). AVT co-localizes with CRF in some neurons of the preoptic nucleus in teleost fish (Yulis & Lederis 1987, Olivierau & Olivereau 1988) and AVT potentiates the effect of CRF on ACTH release in rainbow trout (Baker et al. 1996).

Isolation and confinement had relatively mild effects on CRF-BP mRNA levels. In confined fish, CRF-BP mRNA levels were greater than those of isolated fish at 72 h although there was no difference with the control group. Nevertheless, CRF-BP mRNA levels were positively correlated with those of CRF1 in both experimental protocols (physical disturbance, $R^2=0.26$, $P<0.001$; isolation, $R^2=0.21$, $P<0.001$). This result supports the idea that both transcripts are regulated by similar factors. However, unlike hypothalamic CRF, pituitary CRF-BP is under positive glucocorticoid control in rodents (Behan et al. 1995, McClennen et al. 1998).

In conclusion, we have characterized the precursor cDNA for rainbow trout CRF-BP. Our results demonstrate that both intensity and duration of stress are important factors determining the magnitude and persistence of the elevation in preoptic area CRF and pituitary CRF-BP mRNA levels in rainbow trout. We hypothesize that pituitary CRF-BP is involved in regulating the activity of the stress axis, possibly by sequestering CRF from its receptor and thus, promoting a more rapid return to homeostasis after stress.

**Acknowledgements**

This research was supported by research grants from NSERC-Canada to T W M and V L T. C D was supported by scholarships from NSERC-Canada, FCAR-Québec and the University of Ottawa. We wish to thank James Nickerson for providing the β-actin primers for the RPA, and Dr Nicholas J Bernier for his enlightening discussions of our work. The authors declare that there is no conflict of interest that would prejudice the impartiality of this scientific work.

**References**


Doyon C, Gilmour KM, Trudeau VL & Moon TW 2003 Corticotropin-releasing factor and neuropeptide Y mRNA levels are elevated in the preoptic area of socially subordinate rainbow trout. General and Comparative Endocrinology 133 260–271.


www.endocrinology-journals.org
Pottinger T & Moran T 1993 Differences in plasma cortisol and cortisone dynamics during stress in two strains of rainbow trout (Oncorhynchus mykiss). Journal of Fish Biology 43 121–130.

Received 18 April 2005
Accepted 22 April 2005
Made available online as an Accepted Preprint 6 May 2005