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Role of peripheral corticotropin-releasing factor and urocortin II in intestinal inflammation and motility in terminal ileum

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Corticotropin-releasing factor (CRF) and the closely related family of neuropeptides urocortins (Ucns) are ancient paracrine-signaling peptides secreted in both the central and peripheral neural circuits. CRF and Ucns released from the CNS (central) regulate a plethora of physiological processes that include food intake, inflammation, and bowel motility and permeability. In the gastrointestinal tract, CRF actions are largely proinflammatory, whereas the effects of the Ucn subtypes can be either pro- or antiinflammatory. Central (intracerebroventricular) or peripheral (i.p.) administration of CRF or Ucns inhibits gastric emptying and promotes colonic motility. To ascertain the role of peripherally expressed CRF and UcnII in gastrointestinal inflammation and motility, we generated ileum-specific phenotypic knockouts of these peptides by using RNA interference. Long dsRNA effectively silenced basal expression of CRF and UcnII in ileum. Control dsRNA or saline treatment did not affect CRF or UcnII expression. In an experimental model of toxin-induced intestinal inflammation, inhibition of CRF ablated the inflammatory response (measured by epithelial damage, mucosal edema, and neutrophil infiltration). UcnII dsRNA treatment did not affect the inflammatory response to toxin. Furthermore, ileal motility was increased after site-specific inhibition of both CRF and UcnII. Thus, we demonstrate that ileal-specific CRF promotes inflammation and both CRF and UcnII modulate bowel motility.

corticotropin-releasing factor  |  long dsRNA  |  RNA interference

CRF and Ucns are involved in stress-induced inflammatory responses. The GI tract is laden with commensal flora. The presence of a few pathogenic bacteria can adversely affect GI physiology. These effects include tissue damage, altered immune response, enhanced motility, and epithelial permeability. *Clostridium difficile* is a pathogen that causes a wide range of GI illnesses, ranging from mild diarrhea to life-threatening colitis. *C. difficile* produces two toxins, toxin A (TxA) (enterotoxin) and toxin B (cytotoxin), which are mainly responsible for tissue destruction (14). The acute inflammatory reaction elicited by *C. difficile* TxA has been associated with increased mucosal CRF, CRF-R1, and CRF-R2 expression. The inflammatory response to TxA can be blunted by CRF receptor antagonists (15, 16). A recent study demonstrated that, after TxA exposure, the *CRF-*−/− mice had decreased ileal inflammation as compared with wild-type mice (17). In these experiments, distinction between the roles of central vs. peripheral CRF in the regulation of acute inflammation could not be discerned.

In this study, we determined the role of peripherally expressed CRF and UcnII in inflammation and bowel motility by inhibiting their basal expression in a site-specific fashion by using long

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Abbreviations: CRF, corticotropin-releasing factor; Ucn, urocortin; RNAi, RNA interference; GI, gastrointestinal; TxA, toxin A; sRNA, small interfering RNA; IHC, immunohistochemistry.

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dsRNA-mediated RNA interference (RNAi). First, we investigated their role in an experimental model of *C. difficile* TxA-induced inflammation by measuring the effects of TxA on tissue damage and intestinal fluid secretion. Second, we studied the effect of inhibition of CRF and UcnII on GI motility under basal or unstressed conditions by measuring the rate of passage of a locally administered fecal marker. Our results provide direct evidence that locally synthesized CRF is essential for initiation, augmentation, and perpetuation of acute GI inflammation after TxA exposure.

**Materials and Methods**

**Animals and Experimental Design.** Male Sprague–Dawley rats weighing 225–250 g (B & K, Fremont, CA, or Simonsen Laboratories, Gilroy, CA; design 1 and 2, respectively) were individually housed in hanging wire cages in temperature- and light-controlled rooms. Rats had ad libitum access to chow and water unless otherwise stated. All procedures were in accordance with the Committee on Animal Research at the University of California, San Francisco.

**Cytokine Response to Naked Long dsRNA or Small Interfering RNA (siRNA) Treatment.** There have been conflicting reports as to whether long dsRNA or siRNA (18–25 bp) evoke indiscriminate innate immune responses. Both types of dsRNA have been shown to induce expression of certain specific cytokines in cultured cells (18, 19). However, this response has not been measured in vivo. To address this question, we injected the ileal wall with 15 μg of either long dsRNA (350–1,000 bp) or siRNA (20–25 bp) for CRF, UcnII, β-globin, or GFP. GFP and β-globin dsRNA served as nonspecific controls. Saline-injected rats served as additional controls. Three to 5 days after treatment, basal blood samples were collected by using the tail-nick method, and levels of 14 different cytokines were measured. In all subsequent experiments described below, long dsRNA was used.

**Experimental Design 1: Inflammation Studies.** Rats were injected with dsRNA in four treatment groups (*n* = 5–8): CRF, UcnII, GFP, and β-globin. Normal saline-injected rats served as additional controls. On day 0, rats were anesthetized with a mixture of ketamine and xylazine (75:1 mg per 100 g of body weight). The terminal ileum was identified and exteriorized. dsRNA (15 μg) was injected into the bowel wall of the distal ileum 3–4 cm proximal to the cecum. Ketoprofen (10 mg/kg s.c.) was given postoperatively. On day 4 or 6, a basal blood sample was taken by tail nick, and rats were fasted overnight with free access to water. The following morning, intestinal inflammation was induced by using the *C. difficile* TxA. Briefly, animals were anesthetized with isofluorane, the previous laparotomy incision was opened, and one 3- to 5-cm loop was created in the terminal ileum (encompassing the dsRNA injection site) (20). Tris-HCl buffer (50 mM, pH 7.4) or 5 μg of TxA in buffer was injected into the loop lumen in a 0.4-ml volume. Three hours later, rats were decapitated and trunk blood was collected. Ileal loops were removed. The weight and length of the loops and the volume of fluid secreted into the lumen were recorded. Ileal tissue was collected for histology, Western blotting, immunohistochemistry (IHC), and RNA isolation.

**Experimental Design 2: Motility Studies.** Rats were pretreated with long dsRNA as described in design 1. At the time of dsRNA injections, a silicon catheter (1.2 mm ID and 1.7 mm OD) was implanted 1 cm proximal to the injection site in the terminal ileum. The catheter was sutured to the ileal wall, s.c. tunneled, and exteriorized in the interscapular region. On days 3 and 4 after dsRNA treatment, 0.4 ml of Carmine red (1.2 g per 100 ml of saline) was injected into the catheter. Carmine red, a non-absorbable dye, is routinely used for measuring gut transit time in both humans and rodents (21–24). The number of times red mucus (or red-tinted fecal pellets) was excreted during a 5-h period after Carmine red injection was used to assess intestinal transit time and served as an indicator of intestinal motility.

**Synthesis of dsRNA.** Sense and antisense RNA were synthesized from cDNA inserts cloned in plasmid vectors by using a Mega-Script RNA kit (Ambion, Austin, TX) according to the manufacturer’s specification and as described elsewhere (25). GFP and rat β-globin sequences were used as nonspecific dsRNA controls. siRNA was prepared by digesting 50 μg of long dsRNA with RNase III (Ambion), resulting in a pool of siRNA ranging from 18 to 25 bp in length.

**cDNA Synthesis and RT-PCR.** RNA was isolated from tissues homogenized in Stat60 buffer. Random hexamers were used to reverse transcribe cDNA from 1 μg of total RNA followed by a 30-cycle PCR with use of gene-specific primers. The resulting products were cloned into pTOPO vector (Invitrogen). UcnII-specific forward and reverse primer sequences corresponded to nucleotide numbers 13–31 and 365–384, respectively (GenBank accession no. AY044835). The *T* termnus of GFP was subcloned into pBlueScript SK vector after restriction enzyme digestion of the parent pEGFP plasmid sequences. CRF and β-globin plasmid constructs have been described elsewhere (26). RT-PCR using gene-specific primers was also used to assess the degree of mRNA degradation after dsRNA injections.

**HIC and Western Analysis.** Ileum or colon tissue was fixed in 4% paraformaldehyde and 30% sucrose, embedded in OCT compound (Tissue-Tek, Sakura Finetek, Torrance, CA), sectioned (4 μm), and thaw-mounted onto Superfrost Plus (Fisher, Pittsburgh) slides. CRF was localized by indirect immunofluorescence (FITC) as described (26). Protein extraction, polyacrylamide gel electrophoresis, and Western blot analysis were performed as described elsewhere (27). Antibodies for CRF receptors (Santa Cruz Biotecnology) were used at a dilution of 1:1,000. Secondary antibody (donkey anti-goat) was diluted in 5% nonfat milk (1:4,000). β-Actin (1:4,000, Sigma) antibody was used for normalization.

**Histological Evaluation.** Ileal tissue was stained with hematoxylin/ eosin. The severity of inflammation was scored in a blinded fashion (by E.F.G.). The degree of (*i*) epithelial damage, (*ii*) hemorrhagic congestion and mucosal edema, and (*iii*) neutrophil infiltration was assessed as described (28). Each parameter was given a score of 0–7. Histological damage was calculated by adding scores from all these parameters.

**Measurement of Plasma Hormones and Cytokines.** Blood samples were centrifuged at 1,000 × g for 15 min at 4°C, and the plasma was aliquoted and stored at −80°C. Corticosterone levels (μg/dl) were measured by using an RIA kit (ICN) as described (29). An ELISA was used to measure circulating cytokine levels by using a multiplex rat cytokine panel at Linco Laboratories (St. Louis).

**Statistical Analysis.** Data were first analyzed by one-way ANOVA. A significant (*P < 0.05*) global effect of ANOVA was followed by post hoc tests of individual group differences (Fisher’s probable least-squares difference test).

**Results**

**Inhibition of Basal CRF and UcnII in Rat Ileum by dsRNA and Cytokine Response to dsRNA.** The expression pattern of CRF along the GI tract under basal or unstressed condition is incomplete. By using IHC, we determined that CRF is expressed in the nerve fibers of the submucosal plexus of the ileum (Fig. 1A), jejunum, and
duodenum, but not the colon (data not shown). In the ileum, low levels of CRF were detected in the lamina propria and Paneth cells. No CRF was detectable in the epithelial cells (Fig. 1A Left). Long dsRNA for CRF (dsCRF) and UcnII (dsUcnII) injected into the muscle wall effectively inhibited expression of basal CRF and UcnII mRNAs. By using RT-PCR, we first established that dsRNA effectively inhibited their cognate mRNAs. RT-PCR products for CRF or UcnII in groups treated with dsRNA for CRF or UcnII, respectively, were either very low or undetectable 5 days after dsRNA treatment (Fig. 1B). Both CRF and UcnII products were present in the GFP long dsRNA (dsGFP) group. Furthermore, UcnII mRNA was present in the dsCRF group and vice versa. Cyclophilin, a housekeeping gene, was detectable in all three groups, and its expression was unaffected by RNAi treatment. (A and C) CRF expression is induced 3 h after TxA treatment in the submucosa and lamina propria (LP). This induction is specifically blocked by 6 days of pretreatment with long dsRNA against CRF. CRF, but not UcnII, is proinflammatory. (D and E) TxA treatment markedly destroys tissue morphology in control or UcnII dsRNA groups (n = 4–6) as assessed by epithelial damage, edema, and neutrophil infiltration. Inhibition of CRF by dsRNA resulted in significant protection of tissue damage that appeared similar to buffer-treated naive animals. ANOVA indicated a global significant effect [F(3, 12) = 5.5; P < 0.02]. (F) Reduced tissue damage also resulted in reduced fluid secretion in dsCRF animals as a measured loop ratio (weight of the loop/length). ANOVA indicated a global effect [F(4, 17) = 8; P < 0.001]. Different letters denote statistically significant differences among groups (P < 0.01). [Magnification: ×10 (A and D) and ×40 (C).]
siRNA in vivo did not elicit a nonspecific systemic immune response. Levels of 8 of 14 cytokines were within normal range in all treatment groups (Table 1). Treatment with siRNA resulted in a greater variability within the group than that seen with long dsRNA treatment (Table 1); however, the differences between the groups were not statistically significant. Effects of siRNA on inhibition of protein were shorter in duration as compared with long dsRNA (maximum of 4 days vs. 7–8 days, respectively). Because effects of long dsRNA persisted for a longer duration, we used long dsRNA for the studies reported here.

**Inhibition of Basal CRF but Not UcnII Ablates TxA-Induced Acute Tissue Damage.** To determine the precise role of peripherally synthesized CRF and UcnII in TxA-induced inflammation, dsRNA for CRF, UcnII, or GFP was injected into the ileal wall 4–6 days before TxA exposure. As anticipated, TxA induced CRF peptide expression in the nerve fibers of the submucosa, lamina propria, and the myenteric plexus in all treatment groups (Fig. 1A). As predicted, CRF dsRNA treatment effectively prevented this induction of CRF expression in TxA-treated rats (Fig. 1A and C). Interestingly, not only was CRF expression inhibited by dsCRF, but also the degree of ileal inflammation was markedly decreased in these animals. Blind numerical scoring of the degree of epithelial damage, tissue edema, and neutrophil infiltration of hematoxylin/eosin-stained sections demonstrated that dsCRF-treated animals resembled buffer controls (Fig. 1 D and E). In contrast to dsCRF, dsUcnII did not reduce inflammation (Fig. 1D). The severity of tissue damage paralleled ileal loop fluid secretion (expressed as a loop ratio). dsCRF animals had significantly decreased fluid secretion as compared with dsUcnII and dsGFP control (Fig. 1F).

**Inhibition of CRF and UcnII Increases Ileal Motility Under Basal Conditions.** Peripheral CRF- and UcnII-related mechanisms are known to contribute to intestinal motility (29, 30). Silencing of CRF, but not UcnII, increased the total number of fecal pellets excreted over 24 h as measured daily for 4 days after dsRNA injections [ANOVA: F (2, 13) = 4.7; P < 0.03; Fig. 2A]. Interestingly, the total dry weight of fecal pellets collected in a 24-h period was similar in all treatment groups (data not shown). The increase in the number of pellets was more predominant in the lights-on period and before the diurnal rise of endogenous corticosterone concentrations. To further elucidate diurnal effects of corticosterone on ileal motility, animals were preinjected with dsRNA and implanted with ileal catheters (design 2). On the morning of days 3 and 4, Carmine red dye was injected into the catheter, and the intestinal transit time and fecal frequency were monitored for 5 h. In contrast to cumulative fecal output, both dsCRF and dsUcnII groups displayed an increased number of red mucus/pelets excreted in a 5-h period as compared with dsGFP controls [ANOVA: F (2, 8) = 5.2; P < 0.04; Fig. 2B]. No abnormal luminal fluid accumulation was discernable at the time of death, and hematoxylin/eosin staining displayed normal ileal histology in all groups. Enhanced expulsion of red dye was attributed to increased propulsion activity of ileum due to inhibition of basal CRF or UcnII.

**CGR Receptors Are Differentially Regulated by CRF and UcnII.** We investigated whether specific inhibition of either CRF (that acts predominantly through CRF-R1) or UcnII (acting specifically through CRF-R2 (8)) would lead to increased expression of CRF-R1 or CRF-R2 as a means of compensation for the loss of ligand after TxA treatment. Western blot analysis of ileal tissue from buffer- or TxA-treated animals shows specific up-regulation of CRF-R1 receptor in rats treated with dsCRF (Fig. 3). Expression of CRF-R2 was similar in all treatment groups.

**Plasma Corticosterone (B) Response to TxA Is Blunted in dsCRF-Treated Animals.** To examine the effects of TxA-mediated inflammation on B release, we measured plasma B levels before and after exposure to TxA. Basal or pre-TxA concentrations of B were within normal range in all treatment groups. TxA exposure increased plasma B levels in dsUcnII and control animals, but not in dsCRF-treated animals, as compared with buffer-injected controls (Fig. 4).

**Discussion**

Our results provide direct evidence that (i) both long dsRNA and siRNA are effective in silencing gene expression in peripheral tissues and neither evokes a nonspecific immune response in vivo, (ii) both CRF and UcnII are synthesized locally in the ileum under basal conditions, (iii) ileal-specific CRF is essential for propagation of proinflammatory effects of TxA, and (iv) inhibition of CRF and UcnII modulate intestinal motility.

**Table 1. Cytokine profile after long dsRNA or siRNA treatment**

<table>
<thead>
<tr>
<th>Treatment</th>
<th>GM-CSF*</th>
<th>IL-1α</th>
<th>MCP-1</th>
<th>IL-2</th>
<th>IL-6</th>
<th>IL-12p70</th>
<th>IFN-γ</th>
<th>IL-18</th>
<th>GRO/KC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Naive</td>
<td>24.5</td>
<td>24.5</td>
<td>173.3</td>
<td>63.8</td>
<td>24.5</td>
<td>24.5</td>
<td>24.5</td>
<td>186.3</td>
<td>88.4</td>
</tr>
<tr>
<td>dsControl</td>
<td>24.5</td>
<td>24.5</td>
<td>212.0</td>
<td>37.8</td>
<td>75.0</td>
<td>67.3</td>
<td>32.5</td>
<td>61.3</td>
<td>44.6</td>
</tr>
<tr>
<td>dsCRF</td>
<td>24.5</td>
<td>24.5</td>
<td>58.7</td>
<td>45.6</td>
<td>171.3</td>
<td>41.2</td>
<td>42.5</td>
<td>300.8</td>
<td>184.2</td>
</tr>
<tr>
<td>dsUcnII</td>
<td>24.5</td>
<td>24.5</td>
<td>43.3</td>
<td>25.1</td>
<td>155.2</td>
<td>26.1</td>
<td>&lt;24.5</td>
<td>46.2</td>
<td>28.9</td>
</tr>
<tr>
<td>siCRF</td>
<td>24.5</td>
<td>24.5</td>
<td>259.4</td>
<td>236.8</td>
<td>284.5</td>
<td>61.0</td>
<td>&lt;24.5</td>
<td>439.3</td>
<td>375.8</td>
</tr>
<tr>
<td>siControl</td>
<td>24.5</td>
<td>24.5</td>
<td>212.0</td>
<td>28.7</td>
<td>28.8</td>
<td>0.4</td>
<td>&lt;24.5</td>
<td>32.5</td>
<td>10.7</td>
</tr>
</tbody>
</table>

Cytokine concentrations in basal plasma sample were measured (pg/ml). Long dsRNA or siRNA treatment had either no effect or a similar effect on cytokine levels (±SEM). si, small interfering; MCP, monocyte chemotactic protein 1; GRO/KC is the murine homologue.

*Levels of 8 of 14 cytokines (GM-CSF, IL-1α, IL-4, IL-5, IL-10, IL-12p70, and TNF-α) were within normal (basal) range (<24.5 pg/ml).
myenteric plexus, and lamina propria under basal conditions. UcnII are expressed in the nerve fibers of the ileal submucosa, duodenum has been reported (7). We now show that CRF and immunoreactive peptide within nerve fibers of the myenteric area of the intestine.

La Fleur et al. CRF or UcnII dsRNA differentially alters TxA-mediated rise in corticosterone (B) concentrations. Basal prefasting or pre-TxA B concentrations were not significantly different among the groups. TxA treatment, however, increased B concentrations significantly in UcnI and control dsRNA-treated rats [ANOVA: F(3, 14) = 7.58; P < 0.01]. B levels in CRF dsRNA-treated rats were not significantly elevated or different from naïve buffer-treated controls. Different letters denotes statistically significant differences.

Interestingly, dsCRF also confirmed a potent effect of local CRF inhibition on acute inflammation. Even when dsCRF injection preceded TxA exposure by up to 7 days, there was no histologic evidence of inflammation, and ileal fluid secretion was reduced 3 h after TxA exposure (the time of killing) in the dsCRF-treated animals. Other reports show that intestinal inflammation is associated with changes in the local expression and signaling of CRF and its receptors (16). Macrophages and lymphocytes are known to synthesize CRF in response to stress (29, 30). Thus, the possibility exists that the local inflammatory cells were also exposed to dsCRF during the ileal muscle layer injection, contributing to the absence of an inflammatory reaction. Thus, CRF appears to be pivotal for the initiation and/or propagation of acute inflammation after ileal TxA exposure. In contrast to CRF, specific UcnII down-regulation did not alter the course of TxA-induced inflammation. It has been reported that peripheral administration of Ucn is more effective in suppressing inflammation and cytokine release than CRF (31). Thus, the final outcome of activation of CRF/CRF-R1 or UcnI/CRF-R2 paracrine circuit on inflammation appears to be highly organ/tissue-specific and context-dependent.

Expression of CRF-R1 and CRF-R2 mRNA levels is increased (as assessed by RT-PCR) in response to TxA-mediated ileitis in CRF-intact animals (16). By using RNAi, it was possible to directly address the response of these receptors in TxA-challenged tissue in the absence of ileal CRF or UcnII. TxA treatment induced CRF-R1 receptor expression specifically in the dsCRF-preinjected animals. Surprisingly, CRF-R2 receptor levels were not significantly different in any treatment group, although they tended to be low in dsCRF and high in dsUcnII animals. These results suggest that the differential up-regulation of CRF R1/R2 may also depend on their ligand rather than TxA treatment as suggested (16). In this regard, CRF and UcnII have been shown to regulate transcription of CRF-R1 in a positive-feedback loop in both central and peripheral cells (32, 33). It is also possible that the message up-regulation for CRF-R2 precedes increase in protein levels. Further investigation is required to understand the regulation of receptor expression during inflammation.

Corticosterone (B) is released in response to stress and pain and has been suggested to reduce GI inflammatory response to TxA treatment (34). In contrast, Ctrh−/− mice, which are unable to hypersecrete B in response to stress, also have decreased inflammatory response to TxA (17). Before initiation of TxA treatment, rats were fasted overnight, and fasting is known to increase plasma B levels (35). In our study, ileal-specific CRF or UcnII down-regulation did not alter this B response to fasting (data not shown); thus, it seems unlikely that B levels at the time of TxA administration were different. After TxA, B levels correlated with the extent of tissue damage over time in all animals, most likely because the decreased severity of inflammation in the dsCRF animals decreased their abdominal pain and systemic stress. Moreover, peripheral release of proinflammatory cytokines can also directly or indirectly (by means of the hypothalamic–pituitary–adrenal axis) increase B release from the adrenals, and release of these cytokines may be altered after TxA treatment.

Ileal-specific inhibition of CRF resulted in increased number of fecal pellets over 24 h. The difference in number of pellets excreted between controls and CRF or UcnII dsRNA-treated animals was more pronounced during the morning phase, corresponding to the period when circulating B levels are at their nadir. This finding is not surprising because CRF, Ucn I, and UcnII expression is regulated by B (36, 37). It appears that CRF or CRF-dependent circuits in the ileum act to decrease local motility, which is in contrast to its effect on colonic motility (11, 13). Thus, CRF and UcnII effects on GI motility can be either...
inhibitory or stimulatory, depending on the region of GI tract and the specific pathways (signals to musculature vs. neurons) that are activated or repressed (17, 38, 39). Further study of the signals that interact with CRF and Ucn will help resolve this issue.

In summary, we show that both CRF and UcnII are synthesized locally in the terminal ileum, and inhibition of CRF by RNAi protects ileal tissue from tissue damage against TxA. Local release of CRF may also be critical for activation of stress response in the local immune cells. This finding suggests that the local release of CRF after stress may trigger, as well as exacerbate, an inflammatory response, perhaps contributing to the pathophysiology of inflammatory bowel disease.

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