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RESEARCH ARTICLE

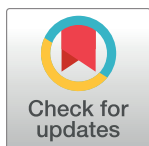
Maternal exposure to low levels of corticosterone during lactation protects adult rat progeny against TNBS-induced colitis: A study on GR-mediated anti-inflammatory effect and prokineticin system

Manuela Zinni¹, Anna Rita Zuena¹, Veronica Marconi¹, Carla Petrella², Ilaria Fusco¹, Chiara Giuli¹, Nadia Canu^{2,3}, Cinzia Severini², Maria Broccardo¹, Vassilia Theodorou⁴, Roberta Lattanzi¹, Paola Casolini^{1*}

1 Department of Physiology and Pharmacology "Vittorio Erspamer", Sapienza University of Rome, Rome, Italy, **2** Institute of Cell Biology and Neurobiology, CNR, Rome, Italy, **3** Department of System Medicine, University of Rome "Tor Vergata", Rome, Italy, **4** INRA, EI-Purpan, UMR 1331 TOXALIM Neuro-Gastroenterology and Nutrition Team, Toulouse, France

☞ These authors contributed equally to this work.

* paola.casolini@uniroma1.it



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Abstract

The early phase of life represents a critical period for the development of an organism. Interestingly, early life experiences are able to influence the development of the gastrointestinal tract and the reactivity to colonic inflammatory stress. We recently demonstrated that adult male rats exposed to low doses of corticosterone during lactation (CORT-nursed rats) are protected against experimental colitis induced by the intracolonic infusion of 2,4,6-trinitrobenzenesulfonic acid (TNBS). Based on these interesting results, we wanted to better investigate which cellular actors could be involved in the protection of CORT-nursed rats from TNBS-induced experimental colitis. Therefore, in the present work, we focused our attention on different factors implicated in GR-mediated anti-inflammatory effect. To address this issue, colonic tissues, collected from control and CORT-nursed healthy animals and from control and CORT-nursed colitic rats, were processed and the following inflammatory factors were evaluated: the expression of (i) glucocorticoid receptors (GR), (ii) glucocorticoid-induced leucine zipper (GILZ), (iii) phospho-p65NF-κB, (iv) the pro-inflammatory cytokines IL-1β and TNF-α, (v) the prokineticins PK2 and PK2L and (vi) their receptors PKR1 and PKR2. We found that adult CORT-nursed rats, in comparison to controls, showed increased expression of colonic GR and reduced expression of pro-inflammatory molecules (IL-1β, TNF-α, PK2 and PK2L) in response to inflammatory colitis. The observed changes were associated with an increase in GILZ colonic expression and with a reduction in phospho-p65NF-κB colonic expression.

Introduction

The inflammatory bowel diseases (IBDs) are chronic relapsing-remitting or progressive inflammatory conditions of the gastrointestinal tract (GI) [1]. Their aetiology implicates a complex interplay between genes and environmental factors [2,3], such as stress [4]. Some human studies have reported a positive association between exacerbation and relapse of IBD and stress [5–13], while others have shown a null association or a minor role of stress [14–19]. This literature disagreement can be mainly due to the great difficulty in establishing the kind of stress and how stress is perceived by the subject. On the contrary, animal studies that allow the standardisation of the experimental procedure (type, intensity and duration of stress) have shown a direct association between stress and susceptibility to experimental colitis [20–25].

The early phase of life is a critical period for neuroendocrine and behavioural development and any stress in this period can hesitate in long-term outcomes [26–29]. Interestingly, maternal separation occurring during the early postnatal period, influences the homeostasis of the GI tract and the vulnerability to stress-induced colitis, as demonstrated by higher susceptibility to dextran sulphate sodium (DSS)-induced colitis in adult rodents [24]. On the other hand, our previous studies [27,30] conducted in rats showed that offspring nursed by mothers with mild hypercorticozonaemia develop the ability to better cope with different situations later in life. In this animal model, the drinking water of mother rats during lactation was supplemented with corticosterone (0.2 mg/ml) [27,31]. Maternal corticosterone is in equilibrium between blood and milk in rodents [30,31], and the hormone is easily absorbed by the GI tract of the pups, as the glucocorticoid permeability of the gut is very high in early postnatal life, up to 17–18 days of age [32]. Indeed, the progeny of these mothers (CORT-nursed rats), once adults, showed improved learning capabilities, reduced fearfulness in anxiogenic situations, a persistent hyporeactivity of the hypothalamus-pituitary-adrenal axis due to an increased number of glucocorticoid receptors (GR) in the hippocampus, and resistance to ischemic neuronal damage [27,30]. Moreover, we have recently demonstrated that adult CORT-nursed rats are protected against 2,4,6-trinitrobenzenesulfonic acid (TNBS)-induced experimental colitis [33]. In fact, colitic CORT-nursed rats showed an improvement in some indices of the pathology (loss of body weight and food intake, increased colonic myeloperoxidase (MPO) activity, and mast cell degranulation) with respect to colitic control animals (adult male rats whose mothers drank water without corticosterone during lactation).

Based on these interesting results, we wanted to better investigate which cellular actors could be involved in the protection of CORT-nursed rats from TNBS-induced experimental colitis. Therefore, in the present work, we focused our attention on some molecular factors linked to GR activation.

Glucocorticoids (GCs) activate the cytosolic glucocorticoid receptors. The ligand-receptor complex then translocates into the nucleus, where it interacts with the promoter regions of different genes. One of the earliest genes induced by GCs is the glucocorticoid-induced leucine zipper (GILZ), a dexamethasone-inducible gene that is rapidly and ubiquitously induced by GCs [34]. GILZ belongs to the transforming growth factor- β (TGF- β) stimulated clone 22 (TSC-22) family of proteins and mediates some of the immunomodulatory effects of GCs [34–36]. The immunomodulatory activity of GILZ involves a physical interaction (homo- and heterodimerisation) between GILZ and the transcription factors that regulate the expression of pro-inflammatory genes [34]. One of the targets of GILZ is the transcription nuclear factor κ B (NF- κ B) [34], a cytosolic protein implicated in the regulation of the function of immune system and of inflammatory processes. After activation, NF- κ B translocates into the nucleus, where it induces the expression of genes encoding for adhesion molecules and pro-inflammatory cytokines [37] involved in the development of the inflammatory response. Studies have

shown that GILZ physically interacts with the p65 subunit of NF- κ B, preventing its activation and nuclear translocation [38,39]. Thus, through a direct interaction with the p65 subunit, GILZ inhibits NF- κ B activation and reduces the expression of pro-inflammatory cytokines such as IL-1 β and TNF- α . Several chemokines are among the mediators of innate immunity regulated by NF- κ B during inflammation [40]. The Bv8/Prokineticins belong to a new family of chemokines comprehending the amphibian Bv8 and Bo8 (from the skin secretion of *Bombina variegata* and *Bombina orientalis*) and the mammalian prokineticin 2 (PK2) [41]. PK2 exerts its biological functions by binding two G-protein-coupled receptors called prokineticin receptor 1 (PKR1) and prokineticin receptor 2 (PKR2), with comparable affinity for both receptors. PK2 displays a major role in inflammatory pain and results strongly up-regulated in inflammatory processes associated with infiltrating cells [42–44]. A splice variant of PK2, containing 21 amino acid insert has been also identified and named PK2L. Interestingly, Marsano and collaborators demonstrated that NF- κ B is involved in the regulation of Bo8 expression [45].

Overall, also considering that it has been demonstrated that GILZ transgenic mice (overexpressing GILZ) are less susceptible to DNBS-induced colitis as compared to wild-type animals [35], in the present investigation, we studied the effects of TNBS-induced colitis in adult CORT-nursed rats, focusing on the following different factors involved in inflammation: (i) GR, (ii) GILZ, (iii) phospho-p65NF- κ B, (iv) the pro-inflammatory cytokines IL-1 β and TNF- α , (v) the prokineticins PK2 and PK2L and (vi) their receptors PKR1 and PKR2.

Materials and methods

Ethics statement

As indicated in [33], all experimental procedures were carried out according to EU Directive 86/609/EEC and to Italian legislation on animal experimentation. The experimental protocols were also approved by the Local Animal Care and Use Committee of Institut National de la Recherche Agronomique (authorisation number MP/02/46/11/08).

CORT-nursed animal model

Female Wistar rats (Charles River, Calco, Italy) weighing 280–320 g were mated and then housed individually. The day after birth, litters were culled to eight pups (four males and four females). Mothers of control rats were maintained on tap water, whereas mothers of CORT-nursed rats had *ad libitum* access to a solution of 0.2 mg/ml corticosterone 21-hemisuccinate from postnatal day 1 to postnatal day 21. Weaning was performed at 21 days of age, and animals were then housed three per cage. Three-month-old male CORT-nursed rats and their controls were used in this study. They were kept in a temperature-controlled room (21 °C) and were allowed free access to water and food.

Induction of experimental colitis and tissue collection

The first day of experiment (day 0), both controls and CORT-nursed male animals were housed one per cage until the day of sacrifice (day 4) and divided in two groups: 1) healthy rats, intracolonicly infused with saline and 2) colitic rats, intracolonicly infused with TNBS (2,4,6-trinitrobenzenesulphonic acid). Overnight fasted control and CORT-nursed rats were anaesthetised by intraperitoneal injection of xylazine (0.6 mg/kg) and ketamine (120 mg/kg), and colitis was induced by an intra-colonic (IC) administration of TNBS at a dose of 30 mg/kg in 0.3 ml of 50% ethanol. Healthy rats were intracolonicly infused with 0.3 ml of saline. TNBS and/or saline were infused through a silicone catheter introduced into the distal colon, 6

cm into the anus, as previously described [46]. On the fourth day after a single TNBS and/or saline instillation, all animals were euthanized by CO₂ inhalation and, for each experimental group, colonic tissue was collected and stored at -80°C.

The current research is a progression of the previous study [33] in which we had $n = 12$ rats for each experimental group and, in order to avoid the litter effect, each litter contributed one or maximum two offspring per group. In the current study, we have analysed a total number of 17 samples coming from different litters, and divided as follows: control healthy ($n = 4$), control colitic ($n = 5$), CORT-nursed healthy ($n = 4$) and CORT-nursed colitic ($n = 4$). Of note, all samples were tested for an inflammatory state by myeloperoxidase (MPO) activity (see ref. 33 for technical details and results).

RNA extraction and real-time PCR

Total RNA was extracted from whole distal colonic samples using the Trizol reagent (Invitrogen, Carlsbad, CA) according to the manufacturer's instructions. RNA yield and purity were determined by spectrophotometric absorption at 260 and 280 nm. mRNA was quantified and 1 µg of mRNA was used to perform reverse transcription (Reverse Transcriptase, Promega) to obtain cDNA. cDNA was used as a template in real-time PCR (iCycler; Bio-Rad) using iQ SYBR Green Supermix (Bio-line). All the measures were performed in triplicate. The reaction conditions were as follows: 95°C for 10 min (polymerase activation), followed by 40 cycles at 95°C for 15 min, 55–50°C (temperature depends on the T_m of the primers) for 15 s and 72°C for 15 s. The reaction mixture without cDNA was used as the control. The results were quantified using the comparative threshold method. The Ct value of the specific gene of interest was normalised to the Ct value of the endogenous control, β-actin, and the comparative Ct method ($2^{-\Delta\Delta C_t}$) was then applied using control healthy rats as the reference samples. The primer sequences are available in [S1 Table](#).

Western blotting analysis

Tissue from whole distal colonic samples from control or CORT-nursed rats was homogenised at 4°C in 200 µl of Ripa buffer (Sigma) containing protease inhibitors (1% v/v) (Sigma). Protein concentrations were determined using the Bradford protein assay. Forty micrograms of protein were re-suspended in sodium dodecyl sulphate (SDS)-bromophenol blue loading buffer with 0.7 M 2-mercaptoethanol. The samples were boiled for five minutes and separated on 8% (for GR), 10% (for PKR1 and PKR2) or 15% (for PK2) SDS-polyacrylamide gels. After electrophoresis (Bio-Rad, Mini Protean Tetra-Cell), the proteins were transferred to nitrocellulose membranes (Bio-Rad) using a system of mini transblot cell (Bio-Rad) at 4°C. After transfer, blots were incubated in a blocking solution containing Tris-buffered saline (TBS), 10% (w/v) Tween-20, 1% (w/v) non-fat milk and 1% (w/v) bovine serum albumin for GR, PKR1, PKR2, β-actin and in a solution containing Tris-buffered saline (TBS), 10% (w/v) Tween-20, 5% (w/v) non-fat milk for PK2. Subsequently, blots were incubated overnight with rabbit anti-GR (1:1000; sc-1004, Santa Cruz Biotechnology Inc.), goat anti-PKR1 (1:500, sc-54313, Santa Cruz Biotechnology Inc.), goat anti-PKR2 (1:500, sc-54313, Santa Cruz Biotechnology Inc.) and rabbit anti-PK2 (1:500; 87360, Abcam) in blocking solution at 4°C. After incubation with the primary antibody, the blots were incubated with horseradish peroxidase-conjugated goat anti-rabbit (1:2000; 7054, Cell Signaling), horseradish peroxidase-conjugated goat anti-mouse (1:2000; 7056, Cell Signaling) or with horseradish peroxidase-conjugated rabbit anti-goat (1:10000; 6741, Abcam) for 1 h at room temperature (21°C±2). To ensure that each lane was loaded with an equivalent amount of protein, the blots were probed with an anti-β actin serum (1:1000; Sigma) overnight at 4°C. Subsequently, blots were incubated with

horseradish peroxidase-conjugated goat anti-mouse antibodies (1:2000; 7056, Cell Signaling) for 1 h at room temperature. Bands were visualised with an enhanced chemiluminescence system (Aurogene). After immunoblotting, digitised images of bands immunoreactive for target and control (actin) molecules were acquired and the area of immunoreactivity corresponding to each band was measured using the NIH ImageJ medical imaging software. A ratio of target to actin was then determined, and these values were compared for statistical significance.

Immunofluorescence assay

Distal colonic samples were collected and fixed in 4% paraformaldehyde and immersed for 24 h in 30% of sucrose at 4°C. Samples were embedded Tissue-Tek OCT (Sakura Finetek Europe B. V) and frozen in isopentane at -45°C. Sections (8- μ m thickness) were cut at -25°C and mounted directly on glass slides. Sections were then processed using fluorescent methods. Slides were pre-incubated with 3% normal goat serum in PBS + 0.3% Triton X-100 (PBST) for 2 h and then incubated overnight at 4°C with rabbit anti-GILZ (1:200, sc-33780, Santa Cruz) or rabbit anti-phospho-p65NF- κ B (1:200, 3033S, Cell Signaling) dissolved in PBST + 1% goat serum. After washing in PBS, slides were incubated with a goat anti-rabbit rhodamine-conjugated secondary antibody (Sigma) for 30 min at room temperature. For nuclei visualisation, slides were incubated with Hoechst 33258 (0.25 μ g/ml) for 5 min at room temperature. Sections were visualised by a confocal laser scanning microscope (Leica SP5, Leica Microsystems, Wetzlar, Germany), and the images captured using 40X oil objective. Pinhole size, image resolution, colour depth, scan speed and averaging (number of scans per image) were the same for all samples analysed. Immunofluorescence was quantified in three fields for every section in at least three sections for each sample by converting pixels in brightness values using the RGB (red, green and blue) [47]. The number of red cells is expressed as percentage (%) of positive labelled with respect to the total area (1024x1024 inches) with comparable blue Integrated Density Value (IDV).

Statistical analysis

Results were analysed by two-way ANOVA (CORT treatment x colitis). The ANOVA analyses were followed by Fisher's LSD *post-hoc* comparisons, where appropriate. Statistical significance was set at $p < 0.05$. All data are expressed as sample mean \pm SEM.

Results

Colonic GR expression

As reported in Fig 1, two-way ANOVA (colitis x CORT treatment) showed a main effect of colitis ($F_{1,13} = 59.72$, $p < 0.0001$) and a significant effect of CORT treatment ($F_{1,13} = 5.73$, $p < 0.05$) on colonic GR expression. Moreover, the ANOVA analysis revealed a significant interaction between colitis and CORT treatment ($F_{1,13} = 6.27$, $p < 0.05$). On this basis, we performed Fisher's LSD *post-hoc* analysis to show that TNBS instillation significantly increased the colonic GR expression in control ($p < 0.01$) and in CORT-nursed colitic animals ($p < 0.001$), with respect to the corresponding healthy rats. Of note, the inflammatory colitis-induced increase in GR was greater in CORT-nursed than in control colitic animals (Fisher's LSD *post-hoc*, $p < 0.05$). No differences were observed between control and CORT-nursed healthy animals.

Colonic GILZ immunofluorescent staining

As reported in Fig 2, two-way ANOVA (colitis x CORT treatment) showed a significant effect of colitis ($F_{1,13} = 65.96$, $p < 0.0001$) and of CORT treatment ($F_{1,13} = 147.6$, $p < 0.0001$) on

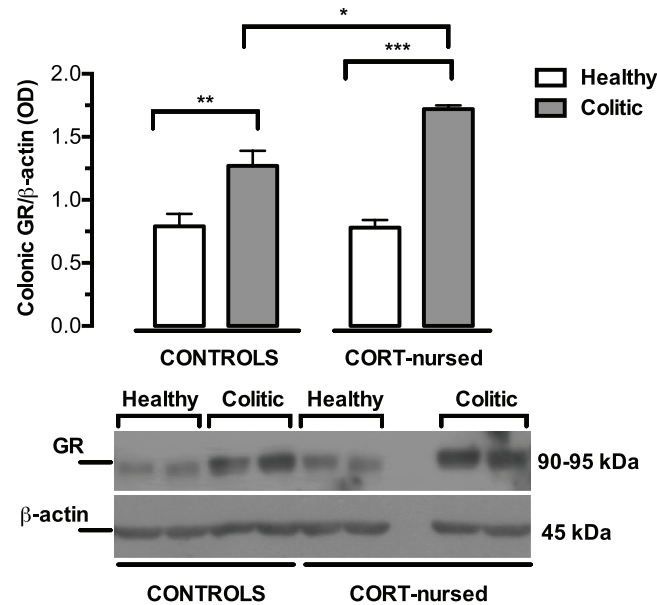


Fig 1. Effects of TNBS-induced colitis on colonic GR expression. Four days after TNBS instillation, increased colonic GR expression was observed in both control and CORT-nursed colitic animals (Fisher's LSD *post-hoc*, ** $p < 0.01$, *** $p < 0.001$ vs. own healthy animals). The CORT-nursed colitic group showed GR expression levels greater than those of the control colitic group (Fisher's LSD *post-hoc*, * $p < 0.05$). The results are expressed as the ratio of optical density (OD) of the GR to that of the β -actin band (control healthy $n = 4$, control colitic $n = 5$, CORT-nursed healthy $n = 4$, CORT-nursed colitic $n = 4$). A representative western blot image was reported.

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colonic GILZ expression. Furthermore, a significant interaction between colitis and CORT treatment was observed ($F_{1,13} = 80.48$, $p < 0.0001$). Fisher's LSD *post-hoc* analysis showed no difference between control colitic and control healthy animals and a significant increase in GILZ expression in the CORT-nursed group (CORT-nursed colitic vs. CORT-nursed healthy) ($p < 0.001$). Interestingly, CORT-nursed colitic animals showed a greater colonic GILZ expression in comparison with control colitic group (Fisher's LSD *post-hoc*, $p < 0.001$). No differences were observed between control and CORT-nursed healthy animals.

Colonic phospho-p65NF- κ B levels

We next investigated the effect of GILZ overexpression showed by CORT-nursed rats, on NF- κ B activation, through the expression of colonic levels of phosphorylated p65 (Fig 3). TNBS instillation significantly increased colonic phospho-p65NF- κ B expression in colitic animals with respect to healthy rats (colitis effect, $F_{1,13} = 32.30$, $p < 0.001$). Moreover, a significant reduction in NF- κ B activation was observed in CORT-nursed rats with respect to control group (CORT treatment effect, $F_{1,13} = 10.94$, $p < 0.01$). We did not find a clear evidence of non-linear interaction between CORT treatment and colitis ($F_{1,13} = 4.39$, $p = 0.056$) while a pure additive effect was observed.

Colonic mRNA expression of the pro-inflammatory cytokines IL-1 β and TNF- α

Two-way ANOVA (colitis x CORT treatment) showed a significant effect of colitis on IL-1 β ($F_{1,13} = 335.4$, $p < 0.0001$) and TNF- α ($F_{1,13} = 422.2$, $p < 0.0001$) and a significant effect of CORT treatment on both cytokines expression (for IL-1 β $F_{1,13} = 257.9$, $p < 0.0001$; for TNF- α

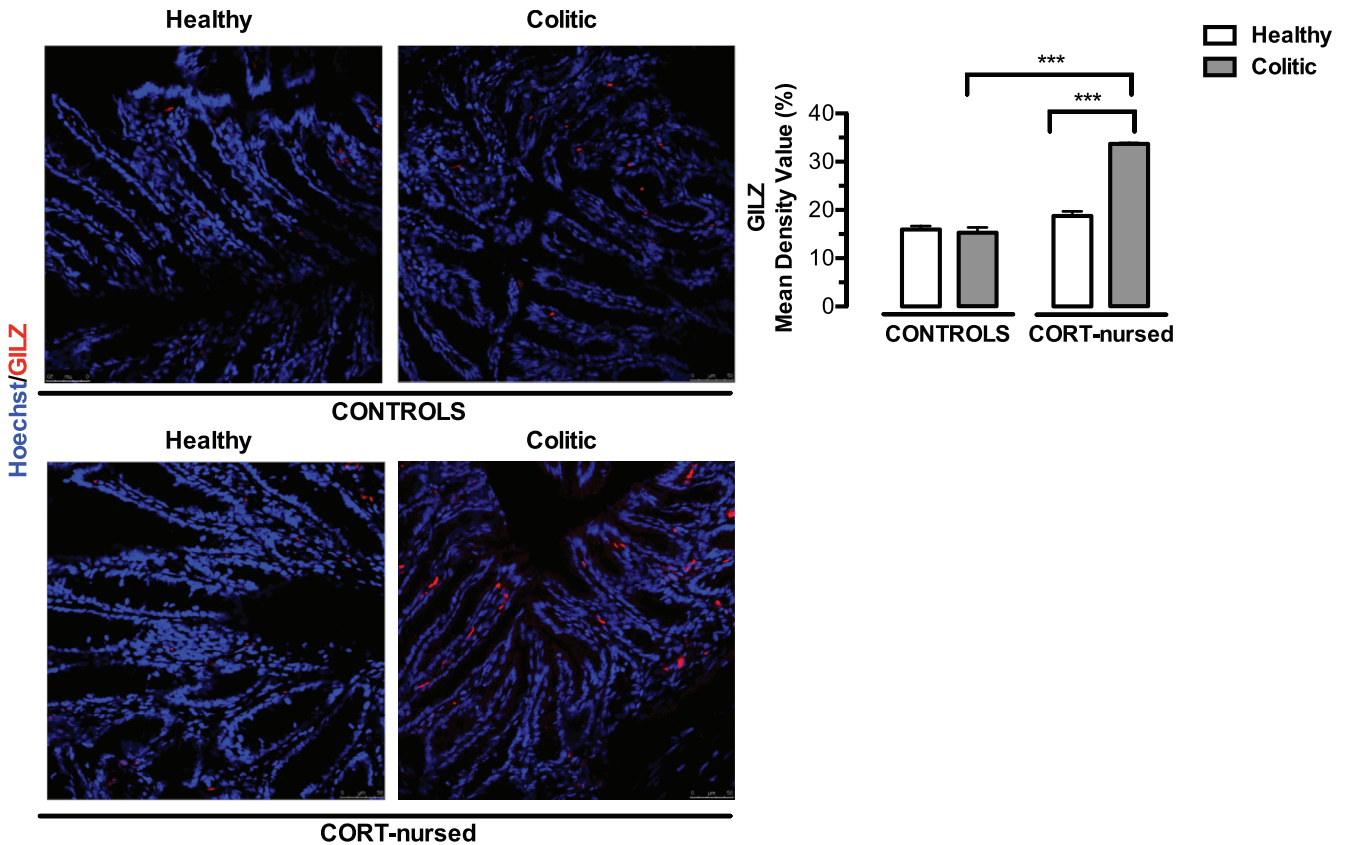


Fig 2. Effects of TNBS-induced colitis on colonic GILZ expression. Four days after TNBS instillation, an increase in colonic GILZ expression was observed in CORT-nursed colitic animals (Fisher's LSD *post-hoc*, *** $p < 0.001$ vs. CORT-nursed healthy). GILZ expression was significantly greater in CORT-nursed colitic rats with respect to control colitic rats (Fisher's LSD *post-hoc*, *** $p < 0.001$). Tissue sections were incubated with a primary antibody against GILZ (red signal). Nuclei were visualised by Hoechst 33258 (blue signal). Results are expressed as percentage (%) of positive labelled with respect to the total area (control healthy $n = 4$, control colitic $n = 5$, CORT-nursed healthy $n = 4$, CORT-nursed colitic $n = 4$). Representative immunofluorescence images are shown.

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$F_{1,13} = 362.3$, $p < 0.0001$) (Fig 4A and 4B). A significant interaction between CORT treatment and colitis was observed (for IL-1 β $F_{1,13} = 271.6$, $p < 0.0001$; for TNF- α $F_{1,13} = 372.4$, $p < 0.0001$). Fisher's LSD *post-hoc* analysis of the four experimental groups showed that cytokines expression followed the same trend as NF- κ B activation. Indeed, inflammatory colitis induced a significant increase in the mRNA levels of IL-1 β (Fig 4A) ($p < 0.001$) and TNF- α ($p < 0.001$) (Fig 4B) in control colitis rats, with respect to control healthy animals, while no modification of these pro-inflammatory cytokines was observed in CORT-nursed colitic rats, in comparison with CORT-nursed healthy rats. Moreover, CORT-nursed colitic rats had lower levels of IL-1 β (Fisher's LSD *post-hoc*, $p < 0.001$) and TNF- α (Fisher's LSD *post-hoc*, $p < 0.001$) mRNA, with respect to control colitic animals. No differences were observed between control and CORT-nursed healthy animals.

mRNA and protein expression of the colonic prokineticins PK2 and PK2L

Fig 5A shows the effect of colitis and CORT treatment on PK2 mRNA colonic expression. Two-way ANOVA (colitis x CORT treatment) showed a significant effect of colitis ($F_{1,13} = 736.0$, $p < 0.0001$) and a significant effect of CORT treatment ($F_{1,13} = 611.4$, $p < 0.0001$). An interaction between colitis and CORT treatment was observed ($F_{1,13} = 626.5$, $p < 0.0001$).

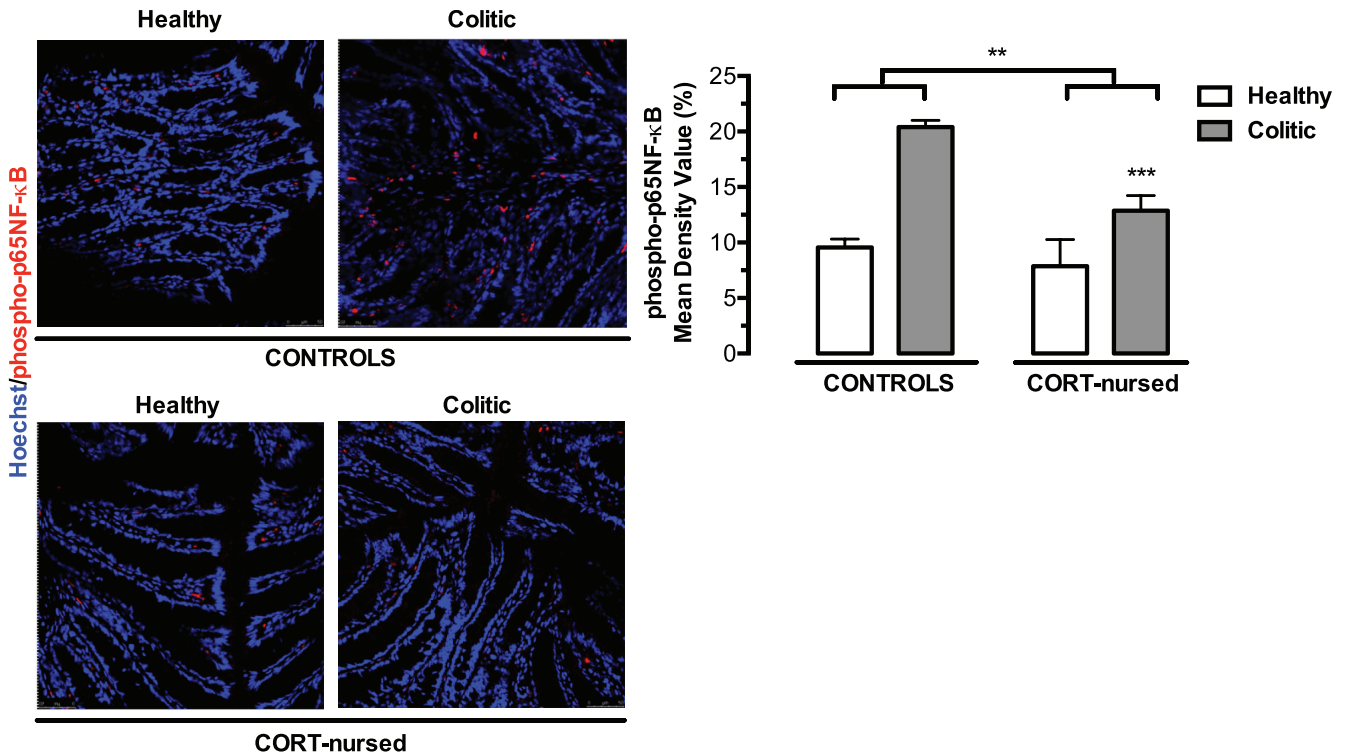


Fig 3. Effects of TNBS-induced colitis on colonic phospho-p65NF-κB expression. Four days after TNBS instillation, an increase in colonic phospho-p65NF-κB expression was observed in colitic animals (** $p < 0.001$ colitic vs. healthy group). A reduction in phospho-p65NF-κB was observed in CORT-nursed rats (** $p < 0.01$ CORT-nursed vs. control group). Tissue sections were incubated with a primary antibody against phospho-p65NF-κB (red signal). Nuclei were visualised by Hoechst 33258 (blue signal). Results are expressed as percentage (%) of positive labelled with respect to the total area (control healthy $n = 4$, control colitic $n = 5$, CORT-nursed healthy $n = 4$, CORT-nursed colitic $n = 4$). Representative immunofluorescence images are shown.

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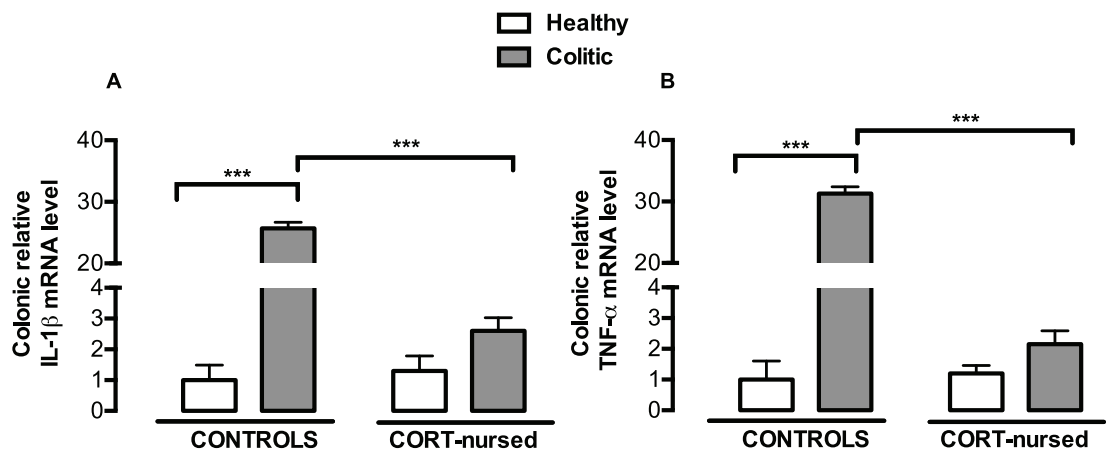


Fig 4. Effects of TNBS-induced colitis on colonic IL-1β and TNF-α mRNA expression. (A) An increase in colonic IL-1β mRNA expression was observed in control colitic rats (Fisher's LSD *post-hoc*, *** $p < 0.001$ vs. control healthy rats). A reduction in IL-1β mRNA expression was observed in CORT-nursed colitic rats (Fisher's LSD *post-hoc*, *** $p < 0.001$ vs. control colitic rats). (B) An increase in colonic TNF-α mRNA expression was observed in control colitic rats (Fisher's LSD *post-hoc*, *** $p < 0.001$ vs. control healthy rats). A reduction in TNF-α mRNA expression was observed in CORT-nursed colitic rats (Fisher's LSD *post-hoc*, *** $p < 0.001$ vs. control healthy rats). The colonic relative mRNA expression levels are expressed in relation to β-actin and presented as fold increases relative to control rats. (Control healthy $n = 4$, control colitic $n = 5$, CORT-nursed healthy $n = 4$, CORT-nursed colitic $n = 4$).

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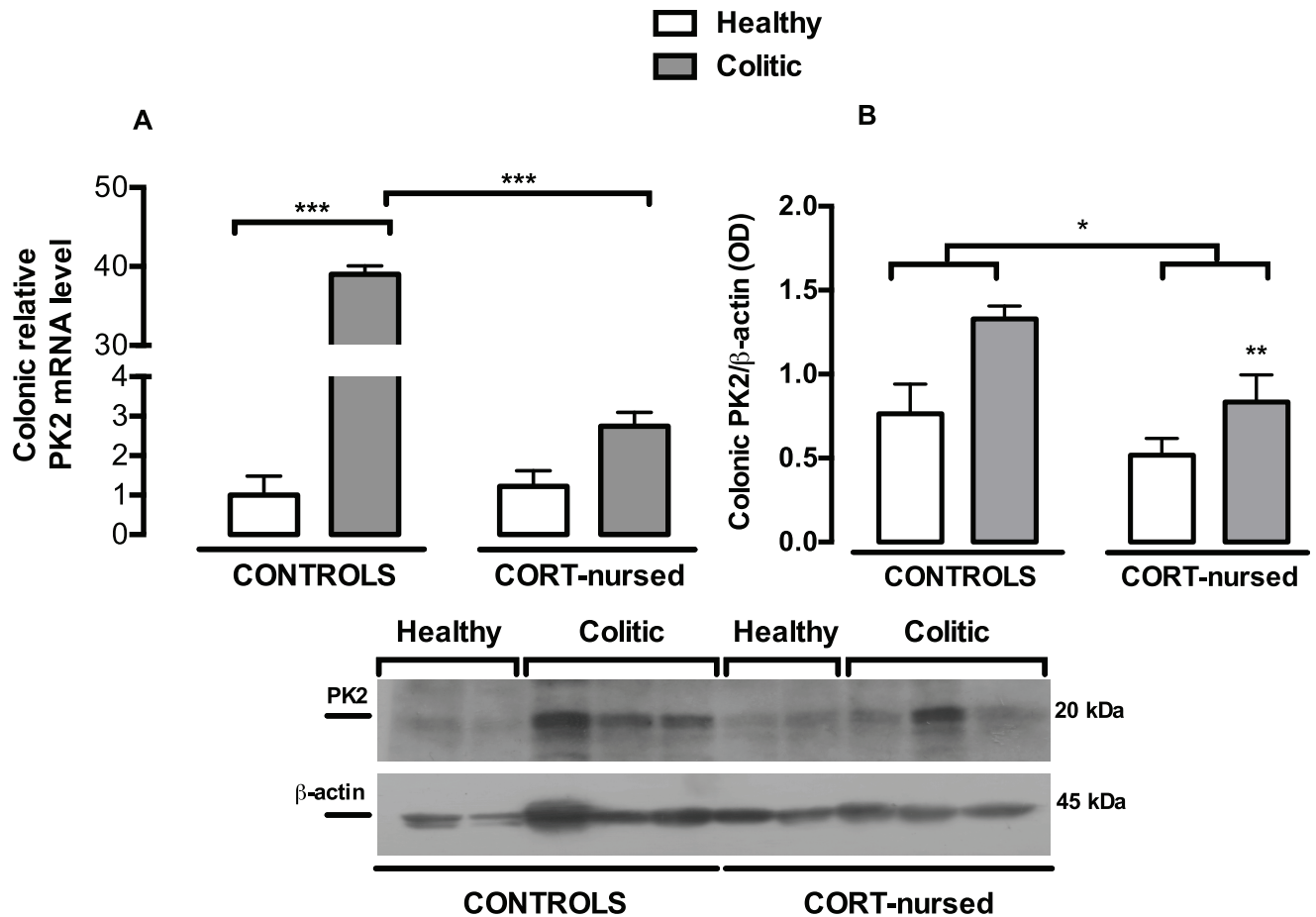


Fig 5. Effects of TNBS-induced colitis on colonic PK2 mRNA and protein expression. (A) Four days after TNBS instillation, an increase in colonic PK2 mRNA expression was found in control colitic rats (Fisher's LSD *post-hoc*, *** $p < 0.001$ vs. control healthy rats). A reduction in PK2 mRNA levels was observed in CORT-nursed colitic animals (Fisher's LSD *post-hoc*, *** $p < 0.001$ vs. control colitic rats). The colonic relative mRNA expression levels are expressed in relation to β -actin and presented as fold increase relative to control rats. (B) Western blot analysis showed an increase in PK2 in colitic rats (** $p < 0.01$ colitic vs. healthy group). A reduction in PK2 was observed in CORT-nursed rats (* $p < 0.05$ CORT-nursed vs. control group). Results are expressed as the ratio of the optical density (OD) of the PK2 and the β -actin band. (control healthy $n = 4$, control colitic $n = 5$, CORT-nursed healthy $n = 4$, CORT-nursed colitic $n = 4$). A representative western blot image is shown.

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Fisher's LSD *post-hoc* analysis showed a significant increase in PK2 mRNA in control colitic rats with respect to control healthy group ($p < 0.001$), whereas no substantial difference was observed in CORT-nursed colitic animals, in comparison with CORT-nursed healthy rats. Interestingly, a significant reduction in PK2 mRNA was observed in CORT-nursed colitic rats, with respect to control colitic animals ($p < 0.001$). No differences were found between control and CORT-nursed healthy animals. Concerning the PK2 protein expression (Fig 5B), TNBS instillation significantly increased colonic PK2 in colitic animals with respect to healthy rats (colitis effect, $F_{1,13} = 11.50$, $p < 0.01$) and CORT treatment significantly reduced PK2 expression in CORT-nursed rats, with respect to control group (CORT treatment effect, $F_{1,13} = 8.16$, $p < 0.05$). No significant interaction between CORT treatment and colitis was observed ($F_{1,13} = 0.92$, $p = 0.35$). Regarding the PK2 long isoform (PK2L) mRNA (Fig 6), two-way ANOVA showed a main effect of colitis ($F_{1,13} = 594.2$, $p < 0.0001$), of CORT treatment ($F_{1,13} = 505.6$, $p < 0.0001$) and an interaction between the two factors ($F_{1,13} = 479.7$, $p < 0.0001$). In agreement with the PK2 mRNA results, we observed an increase in PK2L mRNA in control colitic rats

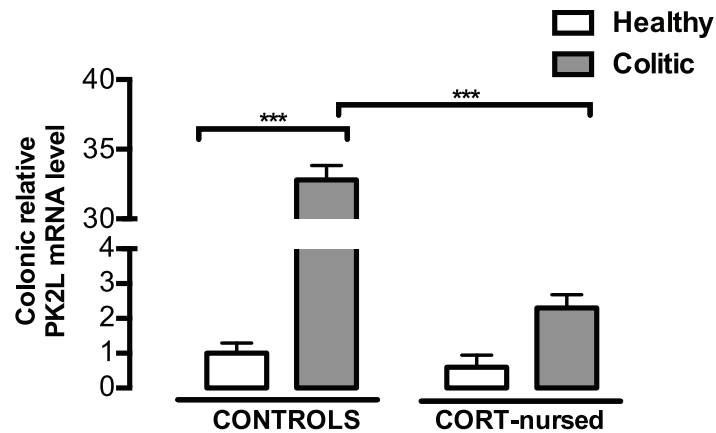


Fig 6. Effects of TNBS-induced colitis on colonic PK2L mRNA expression. Four days after TNBS instillation, an increase in colonic PK2L mRNA expression was revealed in control colitic rats (Fisher's LSD *post-hoc* *** $p < 0.001$ vs. control healthy rats). PK2L mRNA levels in CORT-nursed colitic animals were significantly lower (Fisher's LSD *post-hoc* *** $p < 0.001$ vs. control colitic). The colonic relative mRNA expression levels are expressed in relation to β -actin and presented as fold increase relative to control rats. (control healthy $n = 4$, control colitic $n = 5$, CORT-nursed healthy $n = 4$, CORT-nursed colitic $n = 4$).

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with respect to control healthy animals (Fisher's LSD *post-hoc*, $p < 0.001$), while no difference was showed in CORT-nursed colitic animals with respect to CORT-nursed healthy rats. Interestingly, a significant reduction in PK2L mRNA was also observed in CORT-nursed colitic rats with respect to control colitic rats (Fisher's LSD *post-hoc*, $p < 0.001$). No differences were observed between control and CORT-nursed healthy animals. We could not evaluate PK2L protein levels because there are no antibodies available at the moment.

mRNA and protein expression of the colonic prokineticin receptors PKR1 and PKR2

PKR1 mRNA and protein expression were not affected by colitis and of CORT treatment (Fig 7A and 7B). TNBS instillation significantly increased colonic PKR2 mRNA colonic expression in colitic animals with respect to healthy rats (Fig 8A), (colitis effect $F_{1,13} = 18.77$, $p < 0.001$). On the contrary, no significant effect of CORT treatment and no significant interaction between the two factors were observed. No differences were found between control and CORT-nursed healthy animals and between control colitic and CORT-nursed colitic animals. PKR2 protein expression (Fig 8B) did not show differences between groups.

Discussion

We recently demonstrated that CORT-nursed animals were protected against a TNBS-induced experimental colitis [33]. In fact, adult male offspring reared by dams that drank water supplemented with low dose of corticosterone during lactation (CORT-nursed rats) showed an improvement in some indices of pathology (loss of body weight and food intake, increased colonic MPO activity and mast cell degranulation) with respect to colitic control animals (adult male offspring whose mothers drank water during lactation). The results presented here indicate that CORT-nursed colitic rats showed an increase in colonic GR expression that in turn, triggers modifications of molecular factors implicated in GR intracellular anti-inflammatory response: GILZ over-expression, reduction in colonic NF- κ B activation and reduced expression of the pro-inflammatory cytokines IL-1 β and TNF- α and of the prokineticin PK2.

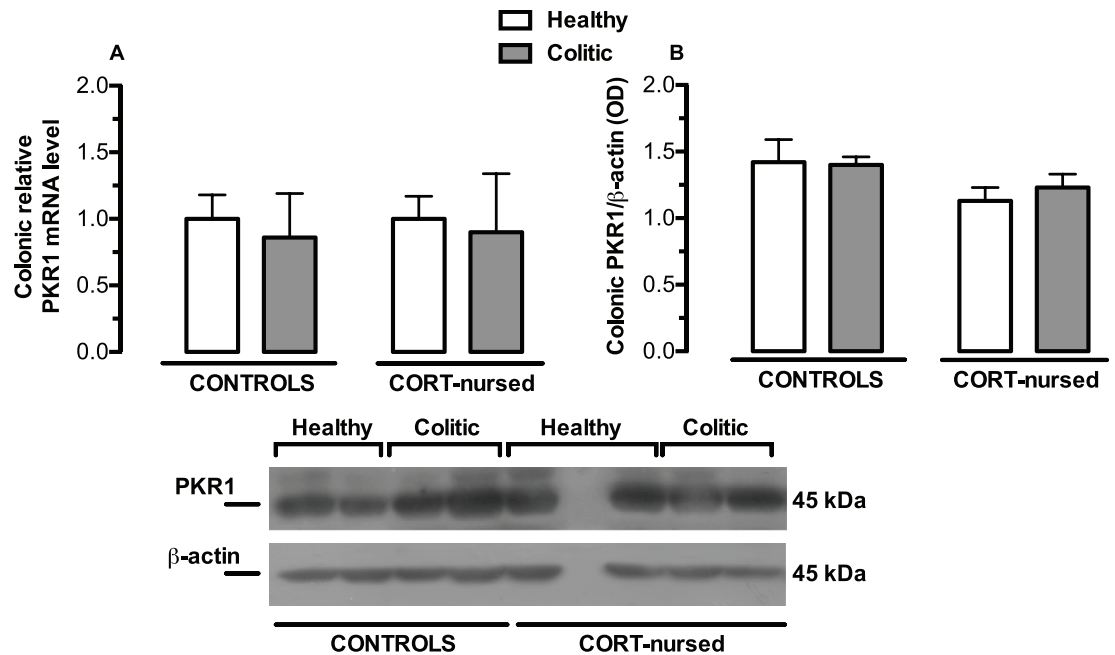


Fig 7. Effects of TNBS-induced colitis on colonic PKR1 mRNA and protein expression. (A) Four days after TNBS instillation, no significant differences were observed in colonic PKR1 mRNA expression between groups. The colonic relative mRNA expression levels are expressed in relation to β -actin and presented as fold increase relative to control rats. (B) Analysis of PKR1 protein expression showed no differences between experimental groups. Results are expressed as the ratio of the optical density (OD) of the PKR1 and the β -actin band. (control healthy $n = 4$, control colitic $n = 5$, CORT-nursed healthy $n = 4$, CORT-nursed colitic $n = 4$). A representative western blot image is shown.

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GR are expressed in different tissues, including the intestine, where they exert a role in controlling the local inflammatory state [48–50]. The results of present research show that TNBS-induced colitis increased colonic GR expression both in control and CORT-nursed rats. Interestingly, this increase was more evident in the CORT-nursed group. In our previous study [33], on the contrary, we observed no change in colonic mucosa GR expression in colitic rats. The disagreement could be explained considering that, in the previous study, we analysed only the mucosal layer while in the present study we have analysed the whole colonic tissue, including both mucosal layers and all other colonic layers (submucosa, muscular layer and serosa). Therefore, the increased GR expression we observed may be accounted for colonic layers rather than by mucosal layer.

In inflammatory colitis, GCs exert some of their immunomodulatory effects by inducing the early gene GILZ [34]. In the present study, we demonstrated that CORT-nursed colitic rats, with respect to the control colitic group, showed higher colonic GILZ expression, which could contribute to lower susceptibility to the TNBS-induced colitis insult. This observation is well-supported by a study conducted in mice by Cannarile and collaborators, showing that GILZ transgenic mice (overexpressing GILZ) were less susceptible to a DNBS-induced colitis, as compared to wild-type animals [35]. Moreover, it has recently been demonstrated that a deficiency in GILZ in knock-out mice exacerbates the effects of chemically-induced colitis [51].

One way by which GILZ could mediate its immunomodulatory effects is through binding to the p65 subunit of the transcription factor NF- κ B [38,39]. This physical interaction prevents NF- κ B activation and its nuclear translocation [38,39]. Our results demonstrate that protected CORT-nursed colitic rats showed reduced activation of NF- κ B, expressed as a reduced level of

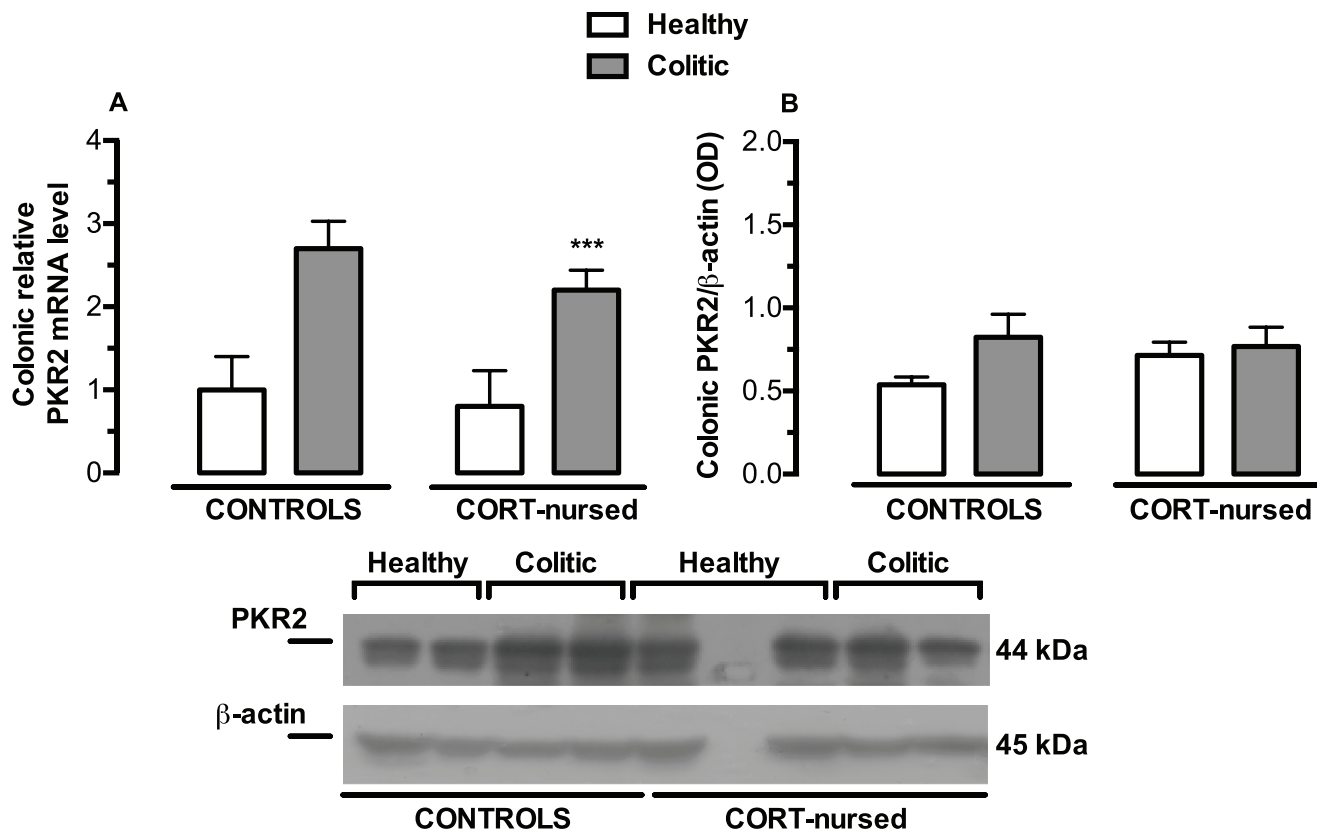


Fig 8. Effects of TNBS-induced colitis on colonic PKR2 mRNA and protein expression. (A) Four days after TNBS instillation, an increase in colonic PKR2 mRNA expression was observed in colitic rats (** $p < 0.001$ colitic vs. healthy group). The colonic relative mRNA expression levels are expressed in relation to β -actin and presented as fold increase relative to control rats. (B) Western blot analysis showed no differences between experimental groups. Results are expressed as the ratio of the optical density (OD) of the PKR2 and the β -actin band. (control healthy $n = 4$, control colitic $n = 5$, CORT-nursed healthy $n = 4$, CORT-nursed colitic $n = 4$). A representative western blot image is shown.

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colonic phospho-p65NF- κ B. Therefore, sustained by the study conducted by Cannarile and collaborators [35], we can hypothesise that GILZ overexpression reduces NF- κ B activation induced by intracolonic TNBS administration and that GILZ-dependent inhibition of the NF- κ B pathway could contribute to the lower susceptibility to TNBS-induced colitis observed in CORT-nursed rats. However, the induction of GILZ expression is not the only mechanism by which GCs can exert their anti-inflammatory action. Indeed, activated GR are also able to interact directly with NF- κ B with a consequent inhibition of NF- κ B translocation and activity [52]. Thus, we cannot exclude that the protection observed in CORT-nursed rats could also be ascribed to the direct inhibition of NF- κ B promoted by the GR itself.

After translocation into the nucleus, NF- κ B induces the expression of pro-inflammatory genes such as those encoding for adhesion molecules and pro-inflammatory cytokines [37]. The results presented here indicate that, in CORT-nursed rats, in comparison with control colitic animals, the colonic pro-inflammatory cytokine levels are in line with the NF- κ B activation profile: a reduction in NF- κ B activation corresponded to a reduction in IL-1 β and TNF- α mRNA levels. Since NF- κ B-mediated cytokine release plays a critical role in the development of experimental colitis [53], the present data suggest that the inhibitory role of GILZ on NF- κ B activity in colonic cells, mediated by GR activation, could contribute to the prevention of disease development in CORT-nursed rats, through the inhibition of pro-inflammatory cytokine

release. These results are again sustained by those of Cannarile and collaborators who demonstrated in GILZ overexpressing transgenic mice that the reduced susceptibility to DNBS colitis is associated with the inhibition of NF- κ B activation and the resulting reduction in IL-1 β and TNF- α expression [35].

During inflammatory processes, cytokines are released together with other molecules such as chemokines belonging to the Bv8/prokineticin family [41]. Here, we have shown that inflammatory colitis induced a significant increase in colonic PK2 mRNA levels and protein expression in control colitic animals, with respect to control healthy rats. The increase in PK2 under inflammatory conditions is well-demonstrated in different animal models of inflammation [42–44]. In particular, in agreement with our results, Watson and collaborators also showed that TNBS instillation increases colonic PK2 levels in rats [54]. Interestingly, in the less vulnerable CORT-nursed rats, PK2 (both mRNA and protein) levels were significantly reduced with respect to the control colitic rats. This protection could be ascribed to the further involvement of NF- κ B with the prokineticin system. Indeed, it has been demonstrated that NF- κ B is able to regulate the expression of the Bo8 gene and that the consensus sequence for NF- κ B identified in the Bo8 promoter is similar to those present in the promoter regions of human and mouse homologs [45]. As reported for PK2, we have also observed an increase in PK2L (the PK2 long form) mRNA levels in colitic rats, confirming that PK2L strongly contributes to inflammation. Indeed, our previous work [43] demonstrated that, in an animal model of inflammation induced by the injection of complete Freund's adjuvant in the hind paw, stronger inflammatory responses observed in rats correlate with higher PK2L mRNA expression in the rat paw. We suppose that PK2L strongly contributes to inflammation through the smaller peptide, PK2 β , derived by its proteolytic cleavage. Indeed, in functional and receptor binding assays, the recombinant PK2L protein demonstrated very poor activity, whereas PK2 β retained its activity and selective affinity for PKR1 [55]. As we observed a significant decrease in PK2L levels in CORT-nursed colitic rats, with respect to control colitic animals, we can assume that PK2L expression, similar to PK2 expression, could undergo regulation by NF- κ B.

Regarding the prokineticin receptors, we found no differences in PKR1 levels (at both the mRNA and protein levels) between control and CORT-nursed colitis rats, while PKR2 mRNA was increased after TNBS-induced colitis; this is in accordance with previous evidence suggesting that only PKR2 is the inducible receptor [42,56]. Conversely, PKR2 protein expression levels were unchanged. Since we analysed PKR2 mRNA and protein expression at only one time point (four days after TNBS instillation), without performing a time-course, we cannot exclude the possibility that PKR2 mRNA up-regulation had just started at this time point and that increased protein expression occurred at a later time.

In summary, in the current research, we identified different factors implicated in GR-mediated anti-inflammatory effect that could be involved in the protection from colitis in CORT-nursed rats. However, we are aware of the limited number of samples we had at our disposal and of the limits of immunofluorescence analysis for the quantification of GILZ and NF- κ B protein expression, so further studies are needed in order to better clarify the role of these factors in the lower susceptibility of CORT-nursed rats to colitis.

In conclusion, a question still arises: how does maternal exposure to a low dose of corticosterone during lactation protect the adult offspring from intestinal inflammation? Increased maternal plasma levels of corticosterone may be responsible for the effects in the offspring through two different hypothetical mechanisms: (i) by acting directly on the pups or (ii) by influencing maternal behaviour. Regarding the first point, maternal corticosterone reaching the pups through milk [31] might contribute to the induction of long-lasting changes in the gastrointestinal tract, thus modifying (in a protective way) parameters involved in the inflammatory gastrointestinal response. On the other hand, we demonstrated that hypercorticosteronaemic

dams showed an increase in maternal care (arched-back nursing and licking/grooming behaviour) [30]. Thus, we cannot exclude that the positive modifications of inflammatory parameters could have been induced by maternal behaviour that, as demonstrated by Meaney, is able to affect gene expression through epigenetic mechanisms [57]. Certainly, the two mechanisms may coexist; thus, the protection of CORT-nursed rats against inflammatory colitis could be promoted by both maternal behaviour and corticosterone action.

Supporting information

S1 Table. Real-time PCR primer sequences.
(DOCX)

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Author Contributions

Conceptualization: MZ PC RL MB ARZ.

Formal analysis: MZ ARZ VM IF CP.

Funding acquisition: PC.

Investigation: MZ VM IF CP NC CS.

Methodology: PC ARZ MZ CG RL CP VT.

Resources: PC RL MB.

Writing – original draft: MZ ARZ PC RL.

Writing – review & editing: CP MB VT IF VM.

References

1. Khor B, Gardet A, Xavier RJ. Genetics and pathogenesis of inflammatory bowel disease. *Nature*. 2011 Jun 15; 474(7351):307–17. doi: [10.1038/nature10209](https://doi.org/10.1038/nature10209) PMID: [21677747](https://pubmed.ncbi.nlm.nih.gov/21677747/)
2. Ahmad T, Tamboli CP, Jewell D, Colombel J-F. Clinical relevance of advances in genetics and pharmacogenetics of IBD. *Gastroenterology*. 2004 May; 126(6):1533–49. PMID: [15168365](https://pubmed.ncbi.nlm.nih.gov/15168365/)
3. Shanahan F. Inflammatory bowel disease: Immunodiagnostics, immunotherapeutics, and ecotherapeutics. *Gastroenterology*. 2001 Feb; 120(3):622–35. PMID: [11179240](https://pubmed.ncbi.nlm.nih.gov/11179240/)
4. Collins SM. Stress and the Gastrointestinal Tract IV. Modulation of intestinal inflammation by stress: basic mechanisms and clinical relevance. *Am J Physiol Gastrointest Liver Physiol*. 2001 Mar; 280(3):G315–318. PMID: [11171612](https://pubmed.ncbi.nlm.nih.gov/11171612/)
5. Fava GA, Pavan L. Large bowel disorders. I. Illness configuration and life events. *Psychother Psychosom*. 1976 1977; 27(2):93–9.
6. Paar GH. [Psychosomatic aspects in patients with Crohn disease—attempt at a status determination]. *Psychother Psychosom Med Psychol*. 1988 Nov; 38(11):376–89. PMID: [3070623](https://pubmed.ncbi.nlm.nih.gov/3070623/)
7. Bach O, Wild HJ. [Life stress events preceding illness episodes in multiple sclerosis and ulcerative colitis—a comparison]. *Z Gesamte Hyg*. 1990 Aug; 36(8):442–3. PMID: [2267849](https://pubmed.ncbi.nlm.nih.gov/2267849/)
8. Duffy LC, Zielezny MA, Marshall JR, Byers TE, Weiser MM, Phillips JF, et al. Relevance of major stress events as an indicator of disease activity prevalence in inflammatory bowel disease. *Behav Med Wash DC*. 1991; 17(3):101–10.

9. Bitton A, Sewitch MJ, Peppercorn MA, deB Edwardes MD, Shah S, Ransil B, et al. Psychosocial determinants of relapse in ulcerative colitis: a longitudinal study. *Am J Gastroenterol*. 2003 Oct; 98(10):2203–8. doi: [10.1111/j.1572-0241.2003.07717.x](https://doi.org/10.1111/j.1572-0241.2003.07717.x) PMID: [14572569](https://pubmed.ncbi.nlm.nih.gov/14572569/)
10. Garrett VD, Brantley PJ, Jones GN, McKnight GT. The relation between daily stress and Crohn's disease. *J Behav Med*. 1991 Feb; 14(1):87–96. PMID: [2038047](https://pubmed.ncbi.nlm.nih.gov/2038047/)
11. Bitton A, Dobkin PL, Edwardes MD, Sewitch MJ, Meddings JB, Rawal S, et al. Predicting relapse in Crohn's disease: a biopsychosocial model. *Gut*. 2008 Oct; 57(10):1386–92. doi: [10.1136/gut.2007.134817](https://doi.org/10.1136/gut.2007.134817) PMID: [18390994](https://pubmed.ncbi.nlm.nih.gov/18390994/)
12. Greene BR, Blanchard EB, Wan CK. Long-term monitoring of psychosocial stress and symptomatology in inflammatory bowel disease. *Behav Res Ther*. 1994 Feb; 32(2):217–26. PMID: [8155059](https://pubmed.ncbi.nlm.nih.gov/8155059/)
13. Levenstein S, Prantera C, Varvo V, Scribano ML, Andreoli A, Luzi C, et al. Stress and exacerbation in ulcerative colitis: a prospective study of patients enrolled in remission. *Am J Gastroenterol*. 2000 May; 95(5):1213–20. doi: [10.1111/j.1572-0241.2000.02012.x](https://doi.org/10.1111/j.1572-0241.2000.02012.x) PMID: [10811330](https://pubmed.ncbi.nlm.nih.gov/10811330/)
14. North CS, Alpers DH, Helzer JE, Spitznagel EL, Clouse RE. Do life events or depression exacerbate inflammatory bowel disease? A prospective study. *Ann Intern Med*. 1991 Mar 1; 114(5):381–6. PMID: [1992880](https://pubmed.ncbi.nlm.nih.gov/1992880/)
15. von Wietersheim J, Köhler T, Feiereis H. Relapse-precipitating life events and feelings in patients with inflammatory bowel disease. *Psychother Psychosom*. 1992; 58(2):103–12. PMID: [1484919](https://pubmed.ncbi.nlm.nih.gov/1484919/)
16. Riley SA, Mani V, Goodman MJ, Lucas S. Why do patients with ulcerative colitis relapse? *Gut*. 1990 Feb 1; 31(2):179–83. PMID: [2311975](https://pubmed.ncbi.nlm.nih.gov/2311975/)
17. Vidal A, Gómez-Gil E, Sans M, Portella MJ, Salamero M, Piqué JM, et al. Life events and inflammatory bowel disease relapse: a prospective study of patients enrolled in remission. *Am J Gastroenterol*. 2006 Apr; 101(4):775–81. doi: [10.1111/j.1572-0241.2006.00476.x](https://doi.org/10.1111/j.1572-0241.2006.00476.x) PMID: [16494590](https://pubmed.ncbi.nlm.nih.gov/16494590/)
18. Targownik LE, Sexton KA, Bernstein MT, Beatie B, Sargent M, Walker JR, et al. The Relationship Among Perceived Stress, Symptoms, and Inflammation in Persons With Inflammatory Bowel Disease. *Am J Gastroenterol*. 2015 Jul; 110(7):1001–1012; quiz 1013. doi: [10.1038/ajg.2015.147](https://doi.org/10.1038/ajg.2015.147) PMID: [26077178](https://pubmed.ncbi.nlm.nih.gov/26077178/)
19. Li J, Nørgård B, Precht DH, Olsen J. Psychological stress and inflammatory bowel disease: a follow-up study in parents who lost a child in Denmark. *Am J Gastroenterol*. 2004 Jun; 99(6):1129–33. doi: [10.1111/j.1572-0241.2004.04155.x](https://doi.org/10.1111/j.1572-0241.2004.04155.x) PMID: [15180736](https://pubmed.ncbi.nlm.nih.gov/15180736/)
20. Reber SO, Obermeier F, Straub HR, Falk W, Neumann ID. Chronic Intermittent Psychosocial Stress (Social Defeat/Overcrowding) in Mice Increases the Severity of an Acute DSS-Induced Colitis and Impairs Regeneration. *Endocrinology*. 2006 Oct 1; 147(10):4968–76. doi: [10.1210/en.2006-0347](https://doi.org/10.1210/en.2006-0347) PMID: [16794011](https://pubmed.ncbi.nlm.nih.gov/16794011/)
21. Veenema AH, Reber SO, Selch S, Obermeier F, Neumann ID. Early Life Stress Enhances the Vulnerability to Chronic Psychosocial Stress and Experimental Colitis in Adult Mice. *Endocrinology*. 2008 Jun 1; 149(6):2727–36. doi: [10.1210/en.2007-1469](https://doi.org/10.1210/en.2007-1469) PMID: [18308845](https://pubmed.ncbi.nlm.nih.gov/18308845/)
22. Gué M, Bonbonne C, Fioramonti J, Moré J, Del Rio-Lachèze C, Coméra C, et al. Stress-induced enhancement of colitis in rats: CRF and arginine vasopressin are not involved. *Am J Physiol*. 1997 Jan; 272(1 Pt 1):G84–91. PMID: [9038880](https://pubmed.ncbi.nlm.nih.gov/9038880/)
23. Qiu BS, Vallance BA, Blennerhassett PA, Collins SM. The role of CD4+ lymphocytes in the susceptibility of mice to stress-induced reactivation of experimental colitis. *Nat Med*. 1999 Oct; 5(10):1178–82. doi: [10.1038/13503](https://doi.org/10.1038/13503) PMID: [10502822](https://pubmed.ncbi.nlm.nih.gov/10502822/)
24. Milde AM, Enger Ø, Murison R. The effects of postnatal maternal separation on stress responsivity and experimentally induced colitis in adult rats. *Physiol Behav*. 2004 Mar; 81(1):71–84. doi: [10.1016/j.physbeh.2004.01.002](https://doi.org/10.1016/j.physbeh.2004.01.002) PMID: [15059686](https://pubmed.ncbi.nlm.nih.gov/15059686/)
25. Barreau F, Ferrier L, Fioramonti J, Bueno L. Neonatal maternal deprivation triggers long term alterations in colonic epithelial barrier and mucosal immunity in rats. *Gut*. 2004 Apr; 53(4):501–6. doi: [10.1136/gut.2003.024174](https://doi.org/10.1136/gut.2003.024174) PMID: [15016743](https://pubmed.ncbi.nlm.nih.gov/15016743/)
26. Maccari S, Krugers HJ, Morley-Fletcher S, Szyf M, Brunton PJ. The consequences of early-life adversity: neurobiological, behavioural and epigenetic adaptations. *J Neuroendocrinol*. 2014 Oct; 26(10):707–23. doi: [10.1111/jne.12175](https://doi.org/10.1111/jne.12175) PMID: [25039443](https://pubmed.ncbi.nlm.nih.gov/25039443/)
27. Catalani A, Alemà GS, Cinque C, Zuena AR, Casolini P. Maternal corticosterone effects on hypothalamus-pituitary-adrenal axis regulation and behavior of the offspring in rodents. *Neurosci Biobehav Rev*. 2011 Jun; 35(7):1502–17. doi: [10.1016/j.neubiorev.2010.10.017](https://doi.org/10.1016/j.neubiorev.2010.10.017) PMID: [21056056](https://pubmed.ncbi.nlm.nih.gov/21056056/)
28. Felitti VJ, Anda RF, Nordenberg D, Williamson DF, Spitz AM, Edwards V, et al. Relationship of childhood abuse and household dysfunction to many of the leading causes of death in adults. The Adverse Childhood Experiences (ACE) Study. *Am J Prev Med*. 1998 May; 14(4):245–58. PMID: [9635069](https://pubmed.ncbi.nlm.nih.gov/9635069/)

29. Bifulco A, Brown GW, Adler Z. Early sexual abuse and clinical depression in adult life. *Br J Psychiatry J Ment Sci.* 1991 Jul; 159:115–22.
30. Casolini P, Domenici MR, Cinque C, Alemà GS, Chiodi V, Galluzzo M, et al. Maternal exposure to low levels of corticosterone during lactation protects the adult offspring against ischemic brain damage. *J Neurosci Off J Soc Neurosci.* 2007 Jun 27; 27(26):7041–6.
31. Angelucci L, Patacchioli FR, Scaccianoce S, Di Sciullo A, Cardillo A, Maccari S. A model for later-life effects of perinatal drug exposure: maternal hormone mediation. *Neurobehav Toxicol Teratol.* 1985 Oct; 7(5):511–7. PMID: [4080068](#)
32. Yorty JL, Bonneau RH. Impact of maternal stress on the transmammary transfer and protective capacity of herpes simplex virus-specific immunity. *Am J Physiol Regul Integr Comp Physiol.* 2004 Dec; 287(6): R1316–1324. doi: [10.1152/ajpregu.00685.2003](#) PMID: [15308492](#)
33. Petrella C, Giuli C, Agostini S, Bacquie V, Zinni M, Theodorou V, et al. Maternal exposure to low levels of corticosterone during lactation protects against experimental inflammatory colitis-induced damage in adult rat offspring. *PLoS One.* 2014; 9(11):e113389. doi: [10.1371/journal.pone.0113389](#) PMID: [25405993](#)
34. Ronchetti S, Migliorati G, Riccardi C. GILZ as a Mediator of the Anti-Inflammatory Effects of Glucocorticoids. *Front Endocrinol.* 2015; 6:170.
35. Cannarile L, Cuzzocrea S, Santucci L, Agostini M, Mazzon E, Esposito E, et al. Glucocorticoid-Induced Leucine Zipper Is Protective in Th1-Mediated Models of Colitis. *Gastroenterology.* 2009 Feb; 136(2):530–41. doi: [10.1053/j.gastro.2008.09.024](#) PMID: [18996377](#)
36. Ayroldi E, Riccardi C. Glucocorticoid-induced leucine zipper (GILZ): a new important mediator of glucocorticoid action. *FASEB J Off Publ Fed Am Soc Exp Biol.* 2009 Nov; 23(11):3649–58.
37. Tak PP, Firestein GS. NF-kappaB: a key role in inflammatory diseases. *J Clin Invest.* 2001 Jan; 107(1):7–11. doi: [10.1172/JCI11830](#) PMID: [11134171](#)
38. Ayroldi E, Migliorati G, Bruscoli S, Marchetti C, Zollo O, Cannarile L, et al. Modulation of T-cell activation by the glucocorticoid-induced leucine zipper factor via inhibition of nuclear factor kappaB. *Blood.* 2001 Aug 1; 98(3):743–53. PMID: [11468175](#)
39. Riccardi C, Bruscoli S, Ayroldi E, Agostini M, Migliorati G. GILZ, a glucocorticoid hormone induced gene, modulates T lymphocytes activation and death through interaction with NF-kB. *Adv Exp Med Biol.* 2001; 495:31–9. PMID: [11774584](#)
40. Saklatvala J, Dean J, Clark A. Control of the expression of inflammatory response genes. *Biochem Soc Symp.* 2003;(70):95–106.
41. Negri L, Lattanzi R. Bv8-prokineticins and their receptors: modulators of pain. *Curr Pharm Biotechnol.* 2011 Oct; 12(10):1720–7. PMID: [21466441](#)
42. Maftei D, Marconi V, Florenzano F, Giancotti LA, Castelli M, Moretti S, et al. Controlling the activation of the Bv8/prokineticin system reduces neuroinflammation and abolishes thermal and tactile hyperalgesia in neuropathic animals. *Br J Pharmacol.* 2014 Nov; 171(21):4850–65. doi: [10.1111/bph.12793](#) PMID: [24902717](#)
43. Giannini E, Lattanzi R, Nicotra A, Campese AF, Grazioli P, Screpanti I, et al. The chemokine Bv8/prokineticin 2 is up-regulated in inflammatory granulocytes and modulates inflammatory pain. *Proc Natl Acad Sci U S A.* 2009 Aug 25; 106(34):14646–51. doi: [10.1073/pnas.0903720106](#) PMID: [19667192](#)
44. Lattanzi R, Maftei D, Marconi V, Florenzano F, Franchi S, Borsani E, et al. Prokineticin 2 upregulation in the peripheral nervous system has a major role in triggering and maintaining neuropathic pain in the chronic constriction injury model. *BioMed Res Int.* 2015; 2015:301292. doi: [10.1155/2015/301292](#) PMID: [25685780](#)
45. Marsango S, di Patti MCB, Barra D, Miele R. The Bv8 gene from *Bombina orientalis*: molecular cloning, genomic organization and functional characterization of the promoter. *Peptides.* 2009 Dec; 30(12):2182–90. doi: [10.1016/j.peptides.2009.09.007](#) PMID: [19747954](#)
46. Morteau O, Hachet T, Caussette M, Bueno L. Experimental colitis alters visceromotor response to colorectal distension in awake rats. *Dig Dis Sci.* 1994 Jun; 39(6):1239–48. PMID: [8200256](#)
47. Inman CF, Rees LEN, Barker E, Haverson K, Stokes CR, Bailey M. Validation of computer-assisted, pixel-based analysis of multiple-colour immunofluorescence histology. *J Immunol Methods.* 2005 Jul; 302(1–2):156–67. doi: [10.1016/j.jim.2005.05.005](#) PMID: [15992812](#)
48. Ergang P, Leden P, Bryndová J, Zbáňková S, Miksík I, Kment M, et al. Glucocorticoid availability in colonic inflammation of rat. *Dig Dis Sci.* 2008 Aug; 53(8):2160–7. doi: [10.1007/s10620-007-0125-6](#) PMID: [18095161](#)
49. van Tol EA, Petrusz P, Lund PK, Yamauchi M, Sartor RB. Local production of corticotropin releasing hormone is increased in experimental intestinal inflammation in rats. *Gut.* 1996 Sep; 39(3):385–92. PMID: [8949642](#)

50. Schulman G, Robertson NM, Eifenbein IB, Eneanya D, Litwack G, Bastl CP. Mineralocorticoid and glucocorticoid receptor steroid binding and localization in colonic cells. *Am J Physiol.* 1994 Mar; 266(3 Pt 1):C729–740. PMID: [8166236](#)
51. Bereshchenko O, Coppo M, Bruscoli S, Biagioli M, Cimino M, Frammartino T, et al. GILZ promotes production of peripherally induced Treg cells and mediates the crosstalk between glucocorticoids and TGF- β signaling. *Cell Rep.* 2014 Apr 24; 7(2):464–75. doi: [10.1016/j.celrep.2014.03.004](#) PMID: [24703841](#)
52. Rivest S. Regulation of innate immune responses in the brain. *Nat Rev Immunol.* 2009 Jun; 9(6):429–39. doi: [10.1038/nri2565](#) PMID: [19461673](#)
53. Atreya I, Atreya R, Neurath MF. NF-kappaB in inflammatory bowel disease. *J Intern Med.* 2008 Jun; 263(6):591–6. doi: [10.1111/j.1365-2796.2008.01953.x](#) PMID: [18479258](#)
54. Watson RP, Lilley E, Panesar M, Bhalay G, Langridge S, Tian S-S, et al. Increased prokineticin 2 expression in gut inflammation: role in visceral pain and intestinal ion transport. *Neurogastroenterol Motil Off J Eur Gastrointest Motil Soc.* 2012 Jan; 24(1):65–75, e12.
55. Chen J, Kuei C, Sutton S, Wilson S, Yu J, Kamme F, et al. Identification and pharmacological characterization of prokineticin 2 beta as a selective ligand for prokineticin receptor 1. *Mol Pharmacol.* 2005 Jun; 67(6):2070–6. doi: [10.1124/mol.105.011619](#) PMID: [15772293](#)
56. Kisiouk T, Podlovni H, Meidan R. Unique expression and regulatory mechanisms of EG-VEGF/prokineticin-1 and its receptors in the corpus luteum. *Ann Anat Anat Anz Off Organ Anat Ges.* 2005 Nov; 187(5–6):529–37.
57. Weaver ICG, Cervoni N, Champagne FA, D'Alessio AC, Sharma S, Seckl JR, et al. Epigenetic programming by maternal behavior. *Nat Neurosci.* 2004 Aug; 7(8):847–54. doi: [10.1038/nn1276](#) PMID: [15220929](#)