Dysregulation of Toll-Like Receptors in Dogs with Chronic Enteropathies

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Abstract

**Background:** Toll-like receptors (TLRs) recognize microbe-associated molecular patterns (MAMPs) and play an important role in the regulation of inflammation in the gastrointestinal tract.

**Hypothesis/objectives:** There is a dysregulation of TLRs recognizing bacterial MAMPs at the protein level in canine chronic enteropathies (CCE).

**Animals:** 20 healthy control dogs (HCD), 20 dogs with steroid-responsive (SR) and 20 dogs with food-responsive (FR) diarrhea.

**Methods:** Prospective study. Biopsies from duodenum and colon were taken before and after standard therapy in 40 dogs with SR and FR and at necropsy in 20 healthy beagles. Immunohistochemistry was performed to determine the expression of TLR2, 4, 5 and 9 using immunohistochemistry.

**Results:** TLR2 positive cells were downregulated in the duodenum and colon in FR and SR compared to HCD before and after therapy (all p<0.0001). In regard to the epithelium, the expression of TLR2 was higher in FR before therapy (p=0.008) and after (p=0.02) therapy compared to HCD. TLR4 positive cells were significantly upregulated in duodenum in FR before (p=0.005) and after therapy (p=0.002) therapy compared to HCD. In the colon, TLR4 positive cells were significantly upregulated in FR and SR before (p=0.0005) and after (p=0.020) therapy compared to HCD. TLR5 positive cells were significantly decreased in the duodenum and colon of dogs with SR and FR before (p=0.0009) and after (p=0.0002) therapy compared to HCD. In the colon, TLR5 positive cells were significantly decreased in the duodenum and colon in FR and SR compared to HCD before therapy (p<0.0001). In regard to the epithelium, the expression of TLR9 was significantly lower in FR before and after therapy compared to SR group.

**Conclusion and clinical importance:** In this study, TLR4 and TLR5 expression at the protein level was significantly upregulated in inflammatory cells in CCE leading to increased inflammation. The downregulation of TLR9 in cells and epithelium seen in CCE compared to healthy dogs points towards the important role that TLR9 also plays in gut homeostasis in mice and men. Further studies are needed to evaluate the potential use of new therapeutic approaches to impair TLR4 and 5 expression or induce TLR9 activation with corresponding ligands (e.g., synthetic lipopeptides or synthetic CpGs, respectively) in CCE.

Keywords Inflammatory bowel disease; Food-responsive diarrhea; Commensal bacteria; Diarrhea

Introduction

Canine chronic enteropathies (CCE) are a common cause for vomiting and diarrhea in dogs and include inflammatory bowel disease (IBD; sometimes also called steroid-responsive diarrhea (SR)) and food-responsive diarrhea (FR). IBD is characterized by persistent or recurrent clinical signs of gastrointestinal disease of unknown cause associated with histologic evidence of inflammation in the small and/or large intestinal mucosa [1]. Canine IBD is still a clinical exclusion diagnosis, and the therapy is mostly based on experience and includes a combination of elimination diet, antibiotics and immunosuppressive drugs and is often frustrating [2]. The pathogenesis of IBD is incompletely understood, but environmental, immunologic and genetic factors may play an important role. The presence of the intestinal microbiome as well as the loss of tolerance to commensal microorganisms is important for the development of intestinal inflammation in mouse models of IBD [1,3,4]. There is good evidence that excessive mucosal immune response to components of the microbiome, due to abnormal or impaired effector or regulatory cell activity, is important in the pathogenesis of human IBD in genetically predisposed individuals [4]. Several studies about human and canine IBD have indicated significant differences in the intestinal bacterial microbiota between IBD patients and healthy controls [4,5]. Suchodolski et al. [5] demonstrated a significant difference in the...
The purpose of this study was to determine the expression of TLR2, TLR4, TLR5 and TLR9 using these antibodies [21] for immunohistochemistry in dogs with chronic enteropathies (FR or SR) compared to healthy control dogs (HCD) before and after standard therapy and to test the hypothesis that there is a dysregulation of expression of TLRs on the protein level in dogs suffering of CCE.

Material and Methods

Healthy control dogs

Control biopsies were taken at necropsy from 20 healthy Beagle dogs, which served as placebo controls in unrelated pharmacologic studies. This group included 8 males and 12 females (all intact), 11-168 months old (median 87 months), with body weights from 5.4 to 15.5 kg (median 10 kg). These dogs did not receive any drugs, were clinically healthy with no signs of diarrhea or vomiting, and showed no abnormalities in complete blood count, serum biochemical profile, and urinalysis. Furthermore, parasitic and bacterial analyses of fecal samples were performed and the samples were free of Giardia sp., Salmonella sp., Campylobacter sp. and other potential causes of diarrhea. Full thickness biopsies of mesenteric lymph nodes and at least 6 biopsies with an endoscopic biopsy forceps as well as full thickness biopsies from duodenum and colon were taken at necropsy directly after euthanasia. The biopsies were kept in 4% neutral-buffered formalin. The samples were examined by a board-certified pathologist and were histologically unremarkable [17].

Dogs with chronic enteropathies

Forty dogs with signs of chronic gastrointestinal disease referred to the Small Animal Teaching Hospital of the University of Bern between November 2006 and February 2009 were included in this prospective study. Some of the dogs described here were used in another study and information about these dogs and the study protocol is available elsewhere [17]. Selection criteria included a history of chronic diarrhea with or without vomiting that lasted for at least 3 weeks, exclusion of identifiable underlying disorders, and histopathological evidence of intestinal inflammatory cellular infiltrate. Owners of dogs signed a letter of consent in which they agreed to participate in initial and follow-up diagnostic evaluation, including endoscopic exams before and after therapy. None of the dogs had been treated with antibiotics, corticosteroids, or antacids at least 7 days before entering the study. All experimental procedures were approved by the Cantonal Committee for Animal Experimentation, Bern, Switzerland (118-05).

All dogs were given a clinical score using the canine IBDD activity index (CIBDAI) as established by Jergens et al. [22] and were classified as clinically insignificant (score 0–3), mild (4–5), moderate (6–8), or severe (9–18) before and after therapy. Furthermore, the dogs were classified according to their predominant clinical signs as having upper or lower gastrointestinal disease or both. Duodenoscopy and colonoscopy were performed in all dogs except those with severe hypobulinemia (n=7; all <18 g/L), where a 36 h fasting period was used for colonoscopy was considered to be detrimental. After endoscopy, all dogs were treated initially with an elimination diet for 14 days. Recommendations usually state that dogs should be fed an appropriate formulation for at least 4–6 weeks [23]. The elimination diet trial in our study was accomplished for 14 days based on recently obtained evidence, that diet trials of 10 days are adequate to show complete remission in dogs with food responsive diarrhea [24].

Dogs that responded to the elimination diet in the first 14 days (clinical signs improved or resolved) were assigned to the FR group. This group included 20 dogs, 12 males (10 intact, 2 neutered) and 8 females (6 intact, 2 neutered), 6-76 months old (median 21.5 months), with body weights from 7.5 to 61 kg (median 27.9 kg). Breeds in this group included: Bernese Mountain Dog (3), German Shepherd Dog (10), Labrador Retriever (2), Mixed breed (3) and one of each belonging to Whippet, West Highland White Terrier, Leonberger, Great Dane, Malinois, Labrador Retriever, Schwyzer Laufhund, Shi-Tzu, Alaskan Malamute, English Setter and Weimaraner breed.

The dogs that did not respond in the first 14 days of treatment (clinical signs persisted while on the elimination diet) were assigned to the SR group and were given oral prednisolone (1 mg/kg PO q12 h) for 14 days followed by a tapering dosage (50% reduction every 2 weeks if possible). This group included 20 dogs, 12 males (5 intact, 7 neutered)
and 8 females (1 intact, 7 neutered), 32-154 months old (median 61 months), with body weights from 2.9 to 71.7 kg (median 20.5 kg). Breeds in this group included: mixed breed (3), Dachshund (2), Shar Pei (2), Yorkshire Terrier (2) and one of each belonging to Rottweiler, Bull Mastiff, Cavalier King Charles Spaniel, Papillon, Beauceron, Bernese Mountain Dog, Pug, German Shepherd Dog, Golden Retriever, West Highland White Terrier and Malinois breed.

The protocol that was used in this study was described before [17]. Seven of 20 dogs in the SR group were clinically classified as having protein-losing enteropathy (PLE). These dogs showed a panhypoproteinemia and a severe hypoalbuminemia (all<18 g/L). Because of the severe hypoalbuminemia and the need of a quick response, the elimination diet and the treatment with prednisolone in these dogs were started immediately after the first endoscopy.

Tissue samples

Dogs were prepared for endoscopy by withholding food for 36 h and administering a colonic lavage solution by gastric intubation (2 doses of 30 mL/kg of body weight 6–8 h apart). At least six adequate mucosal biopsy specimens were obtained from the duodenum (~10 cm below the caudal duodenal flexure), and middle portion of the descending colon, or from where lesions were visible. An endoscopic score [25] was assigned based on mucosal appearance and on the severity of changes. Samples for subsequent histopathological evaluation were placed in 4% neutral-buffered formalin or paraffin. Biopsy specimens were examined by a board-certified pathologist who was blinded with regard to clinical diagnosis and treatments used.

Immunohistochemistry

The paraffin embedded tissue sections were deparaffinized and rehydrated with a xylol-alcohol series and washed three times with Tris-buffered saline (TBS). The epitope retrieval was performed with citrate buffer (pH 6) in a standardized microwave (92°C for 15 minutes). Sections were cooled to room temperature for 20 minutes. After another washing step with TBS, Triton-X 100 (1%) was utilized for 10 minutes to penetrate the cell membrane. To minimize background, the endogenous peroxidase was blocked with methanol and 3% hydrogen peroxide solution for 15 minutes and the samples were washed again with TBS. To reduce nonspecific binding of proteins, samples were incubated 30 minutes with 5% normal goat serum. The sections were incubated overnight at 4°C with the primary antibody (purified rabbit-anti canine TLR; TLR2 diluted 1:500, TLR4 diluted 1:50, TLR5 diluted 1:150 and TLR9 diluted 1:300 in 0.01 M Tris-NaCl pH 7.4, respectively) [21]. The secondary antibody was administered for 15 minutes by room temperature. After a washing step with TBS, sections were stained with biotin-streptavidin-horseradish-peroxidase for 15 minutes by room temperature and washed again. Amino-9-ethyl-carbazole was used as chromogen. After counterstaining with hematoxilin, sections were cover-slipped with an aqueous mounting medium and digital pictures were taken and analyzed. Positive (mesenterial lymph node) and negative (intestine and mesenterial lymph node without primary antibody) controls were run with every batch of samples.

Coloration of cells as well as the staining of the epithelium was evaluated separately. The degree of expression of TLRs in the biopsy specimens was blindly evaluated by two examiners (MS and AZ) as described previously [26]. In brief, the scoring scheme graded epithelium and cells as follows: epithelium: 0=the same as background; 0.5=close to background; 1=well-marked positivity; 1.5=strong positivity; 2=very strong positivity. Cells: 0=no positive cells visible; 0.5=sporadic single cells; 1=scattered single cells; 1.5= scattered cells with discrete clusters; 2=large groups or clusters of cells; 2.5=dense dissemination.

Statistical analysis

All statistical analyses were performed with NCSS8 (www.ncss.com). A Kruskal-Wallis One-way ANOVA and Kruskal-Wallis Multiple-Comparison z-Value Test (Dunn’s Test) was used to compare the three groups in regard to cells and staining of epithelium in colon and duodenum. Spearman Correlation test was used to show correlations between cell count and staining of epithelium in duodenum and colon. A repeated Measures ANOVA was used to compare the measurements within the three groups over time. For CIBDAI, a 2-way repeated measures routine with groups (FR, SR) and time (before, after) was utilized. A p-value <0.05 was considered significant.

Results

CIBDAI

The CIBDAI was significantly higher in the SR group compared to FR before and after therapy (p=0.022 and 0.011, respectively). Furthermore, there was a significant improvement over time within both groups (both p<0.001). * and † shown in Figure 1).

TLR 2

As a positive control, slides from the same lymph node from a healthy dog were used in all runs. The staining pattern revealed...
positive cells predominantly in the cortex (T-cell area and dendritic cells) and medulla (dendritic cells, macrophages and B-cell area).

In the duodenum, the epithelium was positive stained in the villus and in the crypt area. The staining was more intense in the apical epithelial cells and attenuated in the crypt area both in the duodenum and the colon. Miscellaneous TLR2 positive cell populations were found in the lamina propria and submucosa. Positive cells stained mainly cytoplasmatic and in some extent granular but only scattered membranous. The distribution of TLR2 positive cells in the lamina propria, likely lymphocytes, dendritic cells and possibly macrophages, was disseminated. In addition to these cell populations cells of the submucosal plexus (Meissner’s plexus), possibly neurons, were also positive for TLR2. Most positive cells were found in the healthy control dogs. TLR2 positive cells were significantly down-regulated in the duodenum and the colon in FR (cell scoring range before therapy 0,5-2,5 (median 1,5) and 0,5-2 (median 1) and cell scoring range after therapy 0,5-2 (median 1,5) and 0-2 (median 1), respectively) and SR (cell scoring range before therapy 0,5-2 (median 1,5) and 0,5-1,5 (median 1) and cell scoring range after therapy 0,5-2 (median 1) and 0,5-1,5 (median 1), respectively) group compared to HCD (cell scoring range 1,5-2,5 (median 2,5) and 1,5-2,5 (median 2)) before and after therapy (all p<0.0001). In regard to the epithelium, the expression of TLR2 in the duodenum was higher in FR before therapy (p=0.009, cell scoring range before therapy 0-2 (median 1)) compared to HCD (cell scoring range 0-1 (median 0,5)). No significance was shown in expression of TLR2 positive epithelium in FR group compared to HCD after therapy as well as SR group before and after therapy compared to HCD.

TLR 4

Expression of TLR4 positive cells in the mesenteric lymph node from a healthy dog, which served as positive control, was accentuated in the germinal centre (B- and T-lymphocytes, dendritic cells). The staining pattern was considerably membranous (Figure 2).

Intestinal epithelial cells were negative, as no expression of TLR4 was detected neither in the epithelium of the duodenum nor the colon in healthy and diseased dogs. However, in the lamina propria of the duodenum and the colon invading cells stained positive for TLR4. The staining pattern of positive cells was particularly membranous and infrequently granular-cytoplasmatic. Beside lymphocytes endothelial cells of intestinal capillaries stained positive. TLR4 positive cells were up-regulated in duodenum in FR before (p=0.008; cell scoring range 0,5-2,5 (median 1,5)) and after therapy (p=0.02; cell scoring range 0,5-2,5 (median1,5)) compared to HCD (cell scoring range 0-2 (median 0,75)). In the colon, TLR4 positive cells were significantly up-regulated in FR (cell scoring range before therapy 0,5-2 (median 1,5) and after therapy 0,5-2 (median 1,5) and SR (cell scoring range before therapy 0,5-2 (median 1) and after therapy 0,5-2,5 (median 1,5) group before (p=0.0009) and after therapy (p=0.0002) compared to HCD (cell scoring range 0-1,5 (median 0,5).

TLR 5

The staining pattern of the mesenteric lymph node (positive control) revealed positive lymphocytes and macrophages in the perivascular sinus as well as lymphocytes in the parafollicular space (T-cell area) and the border area of the lymph follicle (B-cell area) as shown in Figure 2.

In the duodenum, the epithelium was stained continuously in both the villus and crypt area. The staining in the villus was accentuated in the apical epithelial cell domain. Furthermore, an apical accentuated coloration was found in epithelial cells in the colon. In the duodenum and colon, different cell populations were positive for TLR5 with a disseminated distribution. There was a basal granular staining of intraepithelial cells in the crypts, supposedly lymphocytes. The number of these cells decreased from duodenum to colon. In the lamina propria, lymphocytes, macrophages and endothelial cells stained positive. The staining pattern seemed to be granular and more cytoplasmatic than membranous. In full thickness biopsies of healthy dogs, the apical epithelium was more intensely stained than the crypts. In addition, the Meissner’s plexus was positively staining for TLR5. In addition to the positive lymphocytes in lymph follicles in full thickness biopsies of the duodenum, also dendritic cells in the middle of lymph follicles were positively stained. However, cells were not precisely defined with other surface markers.

Figure 2: Representative immunohistochemistry results of TLR4, TLR5 and TLR9 antibodies compared with CD79, MAC and CD3 staining at a comparable site of a canine reactive mesenterial lymph node. Lysozyme staining and a negative control section are added for comparison Bar=100 µm.
The FR and SR groups expressed significantly more TLR5 positive cells in the colon in dogs with food responsive (FR) diarrhea, healthy control dogs (HCD) and steroid responsive (SR) diarrhea before therapy compared to the HCD (p=0.005). After therapy, only the FR group still yielded significantly more positive cells in the colon (p=0.020; cell scoring range 0-2; median 1,5; Figure 3B). In the epithelium, there was no significant difference in staining patterns between the groups neither in the duodenum nor in the colon before and after therapy.

The expression of TLR5 in cells and epithelium correlated before and after therapy in duodenum (r=0.506, p<0.001; and r=0.584, p<0.001, respectively) and colon (r=0.501, p<0.001; and r=0.513, p<0.001, respectively). Therefore, a positive correlation exists between the staining of the cells from the lamina propria mucosae and the cells of the epithelium mucosae in FR and SR.

**TLR 9**

The same lymph node as used for TLR2, 4 and 5 was also taken as a positive control for TLR9 in all runs. TLR9 positive cells were accentuated in the lymph follicles (B-cell area), but also some lymphocytes in the precortex and cortex were positively stained (Figure 2). The cellular staining was granular and cytoplasmatic.

Most positive cells were found in healthy dogs, and in decreasing tendency in dogs suffering from FR, SR or PLE. The positive cells in HCD were equally distributed in the crypts and the villi. In general, there were more positive cells in the duodenum, but staining of the epithelium was stronger in the colon than in the duodenum. Cells positive for TLR9 were composed of a homologous group of small cells, supposedly lymphocytes. Meissner’s plexus yielded also positive staining.

Expression of TLR9 positive cells in duodenum and colon were significantly lower in the diseased dogs compared to HCD before and after therapy (all p<0.001). The epithelium was stained significantly weaker in the FR group in duodenum (p=0.009) and colon (p=0.036) before therapy (epithelium scoring range duodenum 0-1 (median 0) and colon 0-1 (median 0.5)) compared to HCD (epithelium scoring range duodenum 0-1 (median 0.75) and colon 0-1 (median 1)) (Figures 5A and C). After therapy, the epithelial staining was significantly weaker in FR (epithelium scoring range duodenum 0-1 (median 0.25) and colon 0-1,5 (median 0,5) for both duodenum (p=0.023) and colon (p=0.013) (Figures 5B and 5D).

The expression of TLR9 correlated within cells or epithelium in the 2 segments before (cells r=0.611, p<0.001; epithelium r=0.519, p<0.001) and after therapy (cells r=0.649, p<0.001; epithelium r=0.667, p<0.001). As there is a positive correlation between the staining of the cells of the lamina propria mucosae and the cells of the epithelium mucosae a strong staining of cells is attended by a strong staining of the epithelium.
The epithelium was stained significantly weaker in the FR group in duodenum (p=0.009) and colon (p=0.036) before therapy (epithelium scoring range 0-1 (median 0) and 0-1 (median 0.5) compared to HCD (epithelium scoring range 0-1 (median 0.75) and 0-1.5 (median 1) (Figures 5A and 5C). After therapy, the epithelial staining was significantly weaker in FR (epithelium scoring range duodenum 0-1 (median 0.25) and colon 0-1.5 (median 0.5) and SR group (epithelium scoring range duodenum 0-1 (median 0) and colon 0-1 (median 0.5) for both duodenum (p=0.023) and colon (p=0.013) (Figure 5B, D). The expression of TLR9 correlated within cells or epithelium in the 2 segments before (cells r=0.611, p<0.001; epithelium r=0.519, p<0.001) and after therapy (cells r=0.649, p<0.001; epithelium r=0.667, p<0.001). As there is a positive correlation between the staining of the cells of the lamina propria mucosae and the cells of the epithelium mucosae a strong staining of cells is attended by a strong staining of the epithelium.

**Figure 5:** TLR9-positive epithelium in the duodenum (ED9) and colon (EC9) in dogs with food responsive (FR) diarrhea, healthy control dogs (HCD) and steroid responsive (SR) diarrhea before (A, C) and after therapy (B, D) the epithelium was stained significantly weaker in the FR group in duodenum and colon before therapy (A, C) compared to HCD (p=0.009 and 0.036, respectively). After therapy, the epithelium was stained significantly weaker in FR and SR group compared to HCD for both duodenum (p=0.023) and colon (p=0.013).

**Discussion**

Pattern recognition receptors like TLRs are responsible for the recognition of MAMPs [6,7] present on diverse microbes including Gram-positive and Gram-negative bacteria, fungi, viruses and parasites. Binding of a TLR ligand results in a downstream signaling cascade with activation of several kinases and ultimately in the activation of NF-kB and the production of proinflammatory cytokines, chemokines, costimulatory molecules and reactive oxygen and nitrogen intermediates [6,11,12]. To the author’s knowledge, this is the first study evaluating protein expression of TLR2, TLR4, TLR5 and TLR9 in the canine intestine. Furthermore, we could prove our hypothesis that there is a dysregulation in TLR expression on the protein level in dogs suffering of CCE.

TLR2 expression was significantly higher in the intestinal epithelium of FR before therapy compared to HCD, but not in SR. Furthermore, a significant down-regulation of TLR2 positive invading cells in duodenum and colon was found in FR and SR before and after therapy. These results are partially discrepant to findings from previously performed studies in dogs [17,18] and men [26,27] which showed an up-regulation of TLR2 mRNA expression in IBD. After mRNA expression, several regulatory mechanisms are involved like post-transcriptional and translational modifications and protein degradation, which could regulate and modify the abundance of protein expression [28,29]. Furthermore, a few studies in men provided evidence of up-regulation of TLR2 in individuals suffering from IBD, not only on the level of mRNA but also protein expression [26,27,30]. In regard to the bacterial population, there are some alterations of the microbiome towards a more gram negative flora and less diversity in canine IBD [5,31]. A reduced TLR2 expression as a result of diminished gram positive bacterial population may be a reason for TLR2 downregulation in dogs with CE, as TLR2 expression is influenced by LTA and PGN, two principal components of the gram positive bacterial wall [10]. An enhanced expression of TLR2 (mRNA and protein) was demonstrated by Volta and coworkers in colonic mucosa and IEC of mice after either supplementation or incubation of *Lactobacillus crispatus* M247 [32]. In dogs, stimulation of duodenal biopsies with TLR ligands revealed anti-inflammatory effects, whereas the impact of Enterococcus faecium was limited [33]. Furthermore, TLR2 appears to mediate some host-protective qualities in mice by restoring intestinal epithelial barrier integrity and reducing colonic mucosal inflammation [34,35]. Last but not least, regulatory T cells (Tregs) play a crucial role in modulating immune reaction, in particular maintaining tolerance to self-antigen. In duodenum of dogs suffering from IBD, diminished numbers of Tregs were found compared to healthy controls [36]. In vitro and in vivo studies in mice revealed regulatory mechanisms of TLR2 signaling in T cell (effector and regulatory T cell) homeostasis [37,38]. On one hand TLR2 ligands enhance the proliferation and IL-2 production of effector T cells while transiently suppressing Treg function, which is regained with declining quantity of TLR2 ligands. On the other hand, TLR2 agonists increase Treg numbers. This mechanism could enable the host to combat pathogens or inducing tolerance against commensals and controlling inflammation accordingly. Diminished TLR2 expression in dogs with CE may therefore point toward a defective T cell homeostasis. Another theory for our findings may be the downregulation of TLR2 as a consequence of severe inflammation in order to protect the host of overwhelming inflammatory response.

In contrast to TLR2 and TLR9, TLR4 expression was significantly up-regulated in invading cells in the duodenum in FR and in the colon in FR and SR group before and after therapy. A trend towards an increased duodenal expression of TLR4 positive cells in SR group before and after therapy was recorded, though significance was not reached. Hence, a larger study population may be needed to achieve statistical significance. Similar findings were found on mRNA [17,19] and protein level in canine and human IBD, respectively [39]. Igarashi and colleagues indicated an upregulation of mRNA of TLR4, 2 and 4 in miniature dachshunds with inflammatory colorectal polyps compared to healthy control dogs [40]. In addition, enhanced transcriptional and protein level of TLR4 positive intestinal LP macrophages were reported in human IBD [30]. Pronounced TLR4 expression of invading cells of diseased dogs may suggest increased inflammation and added contact to LPS, possible through translocated bacteria in response to disturbed intestinal permeability. There are conflicting results in regard to TLR4...
protein expression in intestinal epithelial cells [30,39]. We did not detect any TLR4 positive IEC, neither in dogs suffering from CE nor in controls. In agreement to our findings, human IEC remained negative for TLR4 protein expression in IB and control samples [30]. Furthermore, our findings indicate altered TLR4 expression in CCE compared to healthy controls. Dysregulation of TLR4 may therefore play a role in the pathogenesis of canine IBD as already indicated in German Shepherd Dogs [20].

Similar to findings of TLR4, dogs suffering from CCE showed a significant upregulation of TLR5 positive cells in the lamina propria mucosae (lymphocytes, dendritic cells and endothelial cells) in the colon. Furthermore, TLR5 expression correlated in cells and epithelium within the intestinal segment in duodenum and colon. These results point towards a dysregulated expression and may indicate a pro-inflammatory function of TLR5 in SR and FRD. After therapy, only the FR group exhibited an upregulated expression of TLR5. Since the SR group was treated with prednisolone, it is possible that the gastrointestinal inflammation in these dogs was suppressed through suppression of proinflammatory transcription factors (for example NF-kB) [41]. Furthermore, Amsterdam and coworkers pointed out that steroid hormones can induce apoptosis in hematopoietic cells like monocytes, macrophages and T-lymphocytes which are involved in inflammatory processes [42]. In accordance with the literature, our study suggests a potential influence of glucocorticoids on the expression of TLR5 by induction of apoptosis in cells belonging to the innate immune system, but further studies are needed to investigate this relationship. In Jewish people, the heterozygous carriage of TLR5-stop, a dominant-negative TLR5 polymorphism, is negatively influencing the gene expression [42]. In accordance with the literature, our findings indicate altered TLR4 expression in CCE showing a significant decrease of CD11c+ cells (a marker of human and murine dendritic cells) in the duodenum, ileum and colon of dogs with inflammatory bowel disease compared to healthy control dogs [49].

In conclusion, this is the first study evaluating protein expression of TLR2, TLR4, TLR5 and TLR9 in the canine intestine. TLR4 and TLR5 expression measured by immunohistochemistry was found to be upregulated in inflammatory cells of the lamina propria in the duodenum of FR (TLR4) and colon in dogs with FR and SR (TLR4 and TLR5). However, TLR2 and TLR9 expression was downregulated in lamina propria cells and in the epithelium (TLR9) in dogs with FR and SR. The downregulation of TLR9 in cells and epithelium was found to be significantly decreased in healthy Beagles [19]. Another limitation of the study is the missing double staining of TLR-positive cells to better define the cells staining positive for TLR2, TLR4, TLR5 or TLR9. Possible alteration in the cell population would be possible to detect before and after therapy. Furthermore, it may be interesting to state which cells express TLR2, 4, 5 and 9 before and after therapy, as this may give some hints which kind of cytokine profile is presented and possible may be further investigated. Uematsu and coworkers [48] found a high expression of TLR 5 mRNA in CD11c+ lamina propria cells in the intestine of mice. In addition, they elucidated a flagellin induced stimulation of proinflammatory cytokines like IL-6 and IL-12 through CD11c+ intestinal lamina propria cells in TLR5+/+ but not in TLR5-/- knock out mice [48]. Their data suggested that CD11c+ lamina propria cells induce an innate immune response via TLR 5 by detecting pathogenic flagellated bacteria. However, Kathrani et al. [49] revealed a significant decrease of CD11c+ cells (a marker of human and murine dendritic cells) in the duodenum, ileum and colon of dogs with inflammatory bowel disease compared to healthy control dogs [49].
References


