Interleukin 31 mediates MAP kinase and STAT1/3 activation in intestinal epithelial cells and its expression is upregulated in inflammatory bowel disease

Julia Dambacher, Florian Beigel, Julia Seiderer, Dirk Haller, Burkhard Göke, Christoph J Auernhammer, Stephan Brand

Background/aim: Interleukin 31 (IL31), primarily expressed in activated lymphocytes, signals through a heterodimeric receptor complex consisting of the IL31 receptor alpha (IL31Rα) and the oncostatin M receptor (OSMR). The aim of this study was to analyse IL31 receptor expression, signal transduction, and specific biological functions of this cytokine system in intestinal inflammation.

Methods: Expression studies were performed by RT-PCR, quantitative PCR, western blotting, and immunohistochemistry. Signal transduction was analysed by western blotting. Cell proliferation was measured by MTS assays, cell migration by restitution assays.

Results: Colorectal cancer derived intestinal epithelial cell (IEC) lines express both IL31 receptor subunits, while their expression in unstimulated primary murine IEC was low. LPS and the proinflammatory cytokines TNF-α, IL1β, IFN-γ, and sodium butyrate stimulation increased IL31, IL31Rα, and OSMR mRNA expression, while IL31 itself enhanced IL8 expression in IEC. IL31 mediates ERK-1/2, Akt, STAT1, and STAT3 activation in IEC resulting in enhanced IEC migration. However, at low cell density, IL31 had significant antiproliferative capacities (p<0.005). IL31 mRNA expression was not increased in the TNFARE mouse model of ileitis but in inflamed colonic lesions compared to non-inflamed tissue in patients with Crohn’s disease (CD; average 2.4-fold increase) and in patients with ulcerative colitis (UC; average 2.6-fold increase) and correlated with the IL-8 expression in these lesions (r = 0.564 for CD; r = 0.650 for UC; total number of biopsies analysed: n = 88).

Conclusion: IEC express the functional IL31 receptor complex. IL31 modulates cell proliferation and migration suggesting a role in the regulation of intestinal barrier function particularly in intestinal inflammation.

Abbreviations: CD, Crohn’s disease; ERK, extracellular signal regulated kinase; ELISA, enzyme linked immunosorbent assay; FCS, fetal calf serum; GAPDH, glyceraldehyde-3-phosphate dehydrogenase; GPL, gp130-like receptor; IBD, inflammatory bowel disease; IEC, intestinal epithelial cell; IFN, interferon; IL, interleukin; IL31Rα, interleukin 31 receptor alpha; IFR, leukemia inhibitory factor receptor; LPS, lipopolysaccharide; MAP kinase, mitogen activated protein kinase; MCMV, murine cytomegalovirus; MEK, mitogen activated protein kinase kinase; MTS, 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium; OSM, oncostatin M; OSMR, oncostatin M receptor; PCR, polymerase chain reaction; PI, phosphatidylinositol; RT-PCR, reverse transcriptase polymerase chain reaction; SOCS, suppressor of cytokine signalling; STAT, signal transducer and activator of transcription; TNF, tumour necrosis factor alpha; UC, ulcerative colitis
receptor complex has not been demonstrated in intestinal epithelial cells (IEC) so far. Based on the expression of IL31Rx in epithelial tissues such as skin and lung epithelium and similarities between skin and gut demonstrated by us for other STAT inducing cytokines such as IL22,12 we analysed the IL31 receptor expression in the intestinal epithelium and characterised its signal transduction and its specific biological functions in IEC, particularly in intestinal inflammation.

MATERIALS AND METHODS

Reagents

Polyclonal antibodies to phosphorylated extracellular signal regulated kinase (ERK) 1/2 (Thr183/Tyr185) and phospho-Akt (Ser473) were purchased from Cell Signaling (Beverly, MA, USA). Anti-ERK-1/2 and anti-Akt antibodies were also from Cell Signaling. Phospho-STAT1 antibody was from BD Transduction Laboratories (Franklin Lakes, NY, USA), phospho-STAT3 antibody from Upstate Biotechnology (Lake Placid, NY, USA), and antibodies against STAT1 and STAT3 were from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Horseradish peroxidase conjugated secondary antibodies to mouse or rabbit IgG and chemiluminescent substrate (SuperSignal West Dura Extended Duration Substrate) were from Pierce (Rockford, IL, USA). Human IL31Rx antibody was from R&D Systems (Minneapolis, MN, USA). Recombinant human IL31, tumour necrosis factor (TNF)-α, IL1β, and interferon (IFN)-γ were obtained from R&D Systems (Minneapolis, MN, USA). Lipopolysaccharide (LPS) from Escherichia coli (O26:B6) prepared by phenol extraction was purchased from Sigma (St Louis, MO, USA) and prepared as dispersed sonicate in endotoxin-free water (Life Technologies, Rockville, MD, USA) before diluting to final concentration in supplemented media.

Cell culture

The human colorectal cancer derived IEC lines HT-29, SW480, HCT116, T84, CaCo-2, and DLD-1 were obtained from American Type Culture Collection (Rockville, MD, USA). Cells were grown in Dulbecco’s modified Eagle medium (Gibco BRL/Life Technologies, Gaithersburg, MD, USA) with 100 IU/ml penicillin, 100 μM streptomycin, and 10% heat inactivated fetal calf serum (PAA, Pasching, Austria) in a humidified 5% CO2 atmosphere at 37˚C. For signal transduction experiments with IL31, cells were starved overnight in serum-free medium.

Reverse transcriptase polymerase chain reaction

Reverse transcriptase polymerase chain reaction (RT-PCR) was performed as previously described.12 Briefly, total RNA was isolated using Trizol reagent (Gibco BRL/Life Technologies, Gaithersburg, MD, USA). For RT-PCR, RNA was treated with ribonuclease (RNase)-free deoxyribonuclease (DNA-free-Kit, Ambion) to remove potential genomic DNA contaminants. A volume of 2 μg of total RNA was reverse transcribed using Qiagen Omniscript RT Kit. To control for genomic contamination, an identical parallel PCR reaction was performed containing starting material that had not been reverse transcribed. The following conditions were used for semi-quantitative PCRs: 40 cycles denaturing at 95˚C for 30 seconds, annealing at 60˚C for 30 seconds, extension at 72˚C for 30 seconds. The primers (MWG Biotech, Ebersberg, Germany) for the PCR reactions are shown in table 1. The PCR products were subcloned into pcR 2.1 vector (Invitrogen, Carlsbad, CA, USA) and sequenced.

Quantitative PCR

Real time PCR was performed with an ABI Prism 7700 Sequence Detection System (Applied Biosystems, Darmstadt, Germany) using the Quantitect SYBR Green PCR Kit from Qiagen (Hilden, Germany) following the manufacturer’s guidelines. Oligonucleotide primers (table 1) were designed according to the published sequences. All mRNA expression levels were normalised to β-actin or GAPDH expression in the respective cDNA preparation.

Signal transduction experiments, gel electrophoresis, and immunoblotting

The signal transduction experiments were performed in overnight serum starved IEC lines as indicated. Cells were stimulated with 100 ng/ml IL31, unless indicated otherwise. This concentration was based on pilot experiments demonstrating a significantly higher effect of 100 ng/ml for the activation of certain kinases and cell migration than lower concentrations. Cells were solubilised in lysis buffer containing 1% Nonidet P-40, 20 mM TRIS-HCl (pH 7.4), 150 mM NaCl, 2 mM EDTA, 2 mM EGTA, 10 μg/ml aprotinin, 2 mM phenylmethylsulfonyl fluoride, 10 μg/ml leupeptin and phosphatase inhibitors (400 mM sodium orthovanadate, and 4 mM NaF) and were passed six times through a 21 gauge needle. After 30 minutes on ice, lysates were cleared by centrifugation at 10 000 g for 20 minutes. The protein concentration of each sample was quantified by the Bradford method. Immunoblotting was performed as previously described.13

Immunofluorescence staining

Immunofluorescence analysis of IL31Rx expression in the IEC cell line HCT116 was performed as follows. Cells were fixed with 3.2% paraformaldehyde/PBS for 20 minutes and were then permeabilised with 0.5% Triton X-100/PBS for 5 minutes. Cells were blocked with 10% rabbit serum/PBS for 1 hour and then incubated with the primary antibody (anti-IL31Rx, R&D Systems) for 1 hour. After washing with PBS, cells were incubated with a FITC conjugated anti-goat secondary antibody (Sigma, Taufkirchen, Germany) and Hoechst 33342 (Sigma) for 1 hour. Cells were washed with PBS, mounted with Mowiol and were analysed under a fluorescence microscope. In negative controls, cells were stained omitting the primary antibody.

Enzyme linked immunosorbent assay

For the quantification of IL-8 release, BD OptEIA Human IL-8 ELISA Kit II (BD Biosciences, Bedford, MA, USA) was used according to the manufacturer’s instructions.

Cell proliferation and cell restitution assays

HCT116 cells were seeded onto 96 well plates at densities of cells per well as indicated and were allowed to attach overnight.

Table 1 Primers used for PCR analysis

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<th>hIL31Rx forward: 5'-gaagacggaatcctgagcag-3'</th>
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Cells were then stimulated with 10, 50, and 100 ng/ml IL31, or with cytokine-free medium (negative control) for 48 hours. The cell proliferation rate was determined by MTS assay on day 2 using the CellTiter 96 Aqueous Non-Radioactive Cell Proliferation Assay (Promega, Madison, WI, USA) according to the manufacturer’s instructions.

Cell restitution assays were performed as wounding assays as previously described. Briefly, SW480 cells, which were the most suitable IEC line in pilot experiments, were grown in six well plates to complete confluence. Using a sterile razor blade, eight standardised wounds were created in each plate. Detached cells were removed by three washes with PBS, and the cell medium was changed from 10% FCS containing medium to 0.1% FCS containing medium. The cells were stimulated with IL31 (100 ng/ml) or PBS. The cells were washed with PBS after 24 hours and the number of migrated cells (over the wounding edge) was counted under a microscope. Five dishes were analysed for each group (IL31 stimulated and PBS stimulated), whereas for each dish eight separate fields were counted.

**Isolation of primary ileal epithelial cells from heterozygous TNFΔARE mice**

Heterozygous TNFΔARE mice (a generous gift from Dr G Kollias; Hellenic Pasteur Institute, Athens, Greece), which gradually develop chronic inflammation in the ileum from moderate to severe levels at 8 weeks and 18 weeks of age, and C57BL/6 wild type (WT) mice were killed at the age of 18 weeks. Primary IEC from the ileal and colonic epithelium of WT and TNFΔARE/WT mice were purified as previously described. Briefly, the intestinal tissue was cut into pieces and incubated at 37°C in DMEM containing 5% FCS and 1 mM dithiothreitol (DTT) for 30 minutes. The remaining tissue was incubated in 30 ml PBS containing 1.5 mM EDTA for additional 10 minutes. The supernatants were filtered, centrifuged for 5 minutes at 400 g, and the cell pellet was resuspended in DMEM containing 5% FCS. Finally, the primary IEC suspension was purified by centrifugation through a 25%/40% discontinuous Percoll gradient at 600 g for 30 minutes. Cell purity was assessed by determining the absence of CD3+ T cell contamination. Trypan blue exclusion confirmed the presence of at least 80% viable cells after the 2 hour isolation procedure. Primary IEC from the ileum were collected in sample buffer for subsequent RNA isolation.

**Murine cytomegalovirus infection in vivo**

C57BL/6 mice were infected intravenously with 1×10⁶ pfu murine cytomegalovirus (MCMV) of the Smith strain in PBS as previously described. Control mice received an injection of PBS only. After 45 hours, mice were killed by CO₂ asphyxiation. Total RNA of the colon was isolated using Trizol reagent. The mice in vivo studies were approved by the animal care and use committee of the State of Bavaria (Regierung von Oberbayern) following the National Institutes of Health Guide for the Care and Use of Laboratory Animals.
Colonic biopsies

Biopsies were taken from patients with CD and UC undergoing diagnostic colonoscopy. The study was approved by the ethics committee of the Medical Faculty of the University of Munich. All participating subjects gave written, informed consent before biopsy sampling. Four biopsies were collected from each patient: two from macroscopically non-inflamed sites and two from macroscopically inflamed mucosa. IL31, IL31R, and OSMR mRNA expression in HCT116 cells after stimulation with recombinant IL31. Recent studies in other cell lines reported activation of STAT and MAP kinase signalling by IL31, which we therefore analysed in IEC. These experiments demonstrated moderate activation of both STAT1 and STAT3 in IEC after stimulation with IL31 (100 ng/ml). Moreover, IL31 (100 ng/ml) increased phosphorylation levels of ERK-1/2 and Akt proteins.

Statistical analysis

Statistical analysis was performed using two tailed Student’s t test. p Values < 0.05 were considered as significant.

RESULTS

The IL31 receptor complex is expressed in IEC lines

Firstly, we determined if the IL31 receptor complex consisting of IL31Rα and OSMR is expressed in IEC using RT-PCR and cDNA derived from the human IEC lines HT-29, SW480, HCT116, T84, CaCo-2, and DLD-1. In addition, we examined IL31 expression in these cell lines. RT-PCR analysis demonstrated weak IL31 mRNA expression in all cell lines tested (fig 1A). Using primers specific for the long isoform of this receptor, IL31Rα expression was found in all IEC lines analysed with the exception of T84 and DLD-1 cells (fig 1A). OSMR was expressed in all cell lines although only at low levels in HT-29 and T84 cells (fig 1A). The long isoform of IL31Rα was also expressed on protein level in IEC as demonstrated by western blot (fig 1B) and immunofluorescence staining (fig 1C) with an IL31Rα specific antibody. The strongest expression of both receptor subunits was demonstrated in SW480 and HCT116 cells (fig 1A and 1B), which we therefore used in the following signal transduction experiments.

IL31 induces STAT1/3, ERK-1/2, and Akt phosphorylation

After confirming IL31Rα and OSMR expression in IEC, we next analysed if this complex is functional in IEC investigating various signalling pathways after stimulation of HCT116 and SW480 cells with recombinant IL31. Recent studies in other cell lines reported activation of STAT and MAP kinase signalling by IL31, which we therefore analysed in IEC. These experiments demonstrated moderate activation of both STAT1 and STAT3 in IEC after stimulation with IL31 (100 ng/ml). Moreover, IL31 (100 ng/ml) increased phosphorylation levels of ERK-1/2 and Akt proteins.

IL31 modulates IEC proliferation and migration

The activation of ERK MAP kinases and Akt has been linked to increased cell proliferation, while a recent study demonstrated antiproliferative effects for IL31 in lung epithelial cells mediated via STAT3 activation. Therefore, we analysed the effect of IL31 on IEC proliferation using MTS assays and the IEC line HCT116. At cell densities of <3000 cells/well, IL31 strongly decreased cell proliferation (p = 0.001 for 500 cells/well stimulated with 50 ng/ml of IL31, respectively, vs medium stimulated cells), while this effect was lost at higher cell densities and even an opposite, but much less pronounced proliferation-stimulating effect of IL31 (100 ng/ml) was observed with 10 000 cells/well and 15 000 cells/well compared to medium stimulated controls (p = 0.046 and p = 0.013, respectively; fig 3A).

Recently, we demonstrated that activation of Akt kinases and STAT proteins is also involved in IEC migration. Therefore, we next analysed the influence of IL31 on IEC migration. As demonstrated in figure 3B, 100 ng/ml IL31 increased the migration of HCT116 cells significantly (p = 0.005).

IL31, IL31Rα, and OSMR mRNA expression is upregulated in an in vitro model of cell differentiation

Given the influence of cell density on the IL31 mediated effects on cell proliferation, we next analysed potential effects of IEC differentiation on IL31 and IL31 receptor expression in sodium butyrate treated IEC, which is an established in vitro model of cell differentiation. We measured IL31, IL31Rα, and OSMR mRNA expression by quantitative PCR in HCT116 cells after sodium butyrate treatment. There was an increased expression of all three genes that was most pronounced for IL31Rα mRNA (270-fold upregulation after 48 hours, fig 3C), while IL31 and OSMR were upregulated up to 50-fold and 12-fold, respectively (fig 3C).
LPS and the proinflammatory cytokines TNF-α, IL1β, and IFN-γ. The mRNA expression of all three genes was upregulated 24 hours after stimulation with these proinflammatory cytokines (fig 4). The strongest effect on the transcriptional regulation of all three IL31 related genes was seen after stimulation with IFN-γ, which increased OSMR expression 5.4-fold, IL31R mRNA 3.5-fold, and IL31 mRNA 12.1-fold after 24 hours of stimulation.

IL31 increases the expression of proinflammatory cytokines in IEC

Based on the transcriptional upregulation of IL31 and its receptor complex by proinflammatory cytokines, we next analysed if IL31 itself regulates the expression of proinflammatory proteins. In this experiment, we studied in the IEC line HCT116 the effect of IL31 on protein expression of IL8, a prototypic inflammatory cytokine produced by IEC.24 As shown in figure 5, IL31 increased IL8 expression 3.1-fold after 24 hours of stimulation.

IL31 mRNA expression does not correlate with TNF-α mRNA expression in a Th1 mediated model of murine ileitis

Having shown that IL31, IL31Rα, and OSMR mRNAs are upregulated under inflammatory conditions in vitro, we next examined their expression in murine models of intestinal inflammation in vivo. Using heterozygous TNFΔARE mice as a model of chronic ileitis,13 we demonstrated no significant differences in IL31 mRNA expression in these mice compared to wild type mice of the same age and genetic background (fig 6A); however, there was a trend towards higher IL31 expression in colonic epithelial cells compared to ileal epithelial cells (p = 0.088). Similarly, OSMR expression was not significantly higher in TNFΔARE mice in comparison with wild type mice (fig 6B). In contrast, TNFΔARE mice had a significantly higher TNF-α expression than wild type mice (p<0.01; fig 6C).

IL31 mRNA expression is not upregulated in MCMV infection in vivo

In another set of experiments, we investigated colonic IL31, IL31Rα, and OSMR mRNA expression in viral infection using the murine in vivo model of MCMV infection. C57BL/6 mice were infected with 10⁶ pfu MCMV of the Smith strain17 as previously described.18 However, 45 hours after infection, IL31 and IL31Rα mRNA levels were not significantly different compared to non-infected mice (p = 0.73; fig 6D) while there was a trend towards a higher OSMR expression in infected mice (p = 0.06; fig 6D). In contrast, in TNFΔARE mice and corresponding wild type mice, IL31Rα mRNA levels were lower and just above the detection threshold in both mice strains without significant differences between the two strains (data not shown). However, in comparison with the cDNA isolation of the MCMV infected mice (fig 6D), the isolation procedure of ileal and colonic epithelial cells of TNFΔARE mice yielded much lower RNA and cDNA amounts, respectively. Using similar reaction volumes, there was GAPDH detection already after 18 PCR cycles in the MCMV infected mice, while GAPDH was detected in TNFΔARE mice only after 25 PCR cycles, explaining the different levels of IL31Rα expression in two mouse experiments.

IL31 mRNA expression is increased in the inflamed colonic mucosa of patients with inflammatory bowel disease

We next analysed IL31 and its receptor expression in human intestinal inflammation in vivo using real time PCR. In this analysis, we compared IL31, IL31Rα, and OSMR mRNA expression levels in a total of 88 biopsy samples taken from 22 different patients with inflammatory bowel disease (IBD)
including 12 patients with CD and 10 patients with UC. The biopsies were sampled from sites with endoscopically (macroscopic) inflamed colonic mucosa and compared to those of endoscopically non-inflamed colonic mucosa taken from the same 22 patients. We used the IL8 mRNA expression in these biopsies as a control marker of inflammation. IL8 mRNA expression was significantly increased ($p < 0.05$) in the inflamed biopsy samples in CD (fig 7A). Moreover, in CD IL31 mRNA expression levels were higher in inflamed colonic biopsy samples than in non-inflamed colonic lesions (mean increase 2.4-fold; fig 7A) and correlated with IL8 mRNA expression in these lesions ($r = 0.605$; fig 7B, D). In addition, IL31Rα and OSMR mRNA expression was slightly increased in the inflamed tissue of the UC patients (1.5-fold and 1.7-fold, fig 7B).

**DISCUSSION**

Crohn’s disease and ulcerative colitis, the two main forms of inflammatory bowel disease, are defined as idiopathic, chronic relapsing, inflammatory conditions that are immunologically mediated.25 Despite intensive research their aetiology is only incompletely understood.25 26 However, several cytokines such as TNF-α, IL1β, and IL23 play a key part in the aggravation of intestinal inflammation.25–30 Given its important role in inflammatory disorders of the skin,1 6–10 we therefore analysed in this study the novel cytokine IL31 focusing on expression, signalling, and its role in intestinal inflammation.

While activated T cells are the main source of IL31,1 we here demonstrate that IEC lines express the functional IL31 receptor complex consisting of IL31Rα and OSMR, which is consistent with the expression pattern shown for other epithelial cell types such as keratinocytes1 and lung epithelial cells.4 After stimulation with IL31, ERK, Akt, and STAT proteins are activated in IEC. Similar to our findings, activation of STAT1/3 and PI3 kinase signalling has been observed in glioblastoma and melanoma tumour cells,2 and in lung epithelial cells4 following IL31 stimulation. However, in agreement with a recent study in lung epithelial cells, in which strong activation was only obtained by IL31 receptor overexpression,4 the level of STAT activation in non-transfected IEC was only moderate.

Recently, we demonstrated that activation of Akt and STAT proteins mediates IEC proliferation and migration.11 12 19 20 Similarly, our experiments demonstrated that IL31 receptor activation increases IEC migration. Interestingly, IL31 is located on chromosome 12q24 in close proximity to chemokine-like factor 1, which has been very recently shown to be a functional ligand for CCR4.31 Given the results of our experiments, IL31 also has chemotactic properties influencing the migration of IEC. Moreover, an influence on cell migration has been demonstrated for the IL31 related cytokine IL-632 33 and several other IL6-like cytokines such as LIF.34 In addition, it has been shown that LIF may upregulate the gene expression of chemokines such as MCP-1.34 Similarly, we here show that IL31 increases the expression of the proinflammatory chemokine IL8. Particularly the activation of STAT3 is a common signalling pathway mediated by these migration stimulating cytokines.35 For example, we recently demonstrated that cell migration mediated by the IL-10-like cytokine IL22 is dependent on STAT3 activation.11 A very recent study
established that OSMR mediated signal transduction is negatively regulated by SOCS3 through a receptor tyrosine independent mechanism.36 Consistently, we demonstrated that SOCS3 can also inhibit signalling of other STAT activating cytokines.37–42

IL31Rα belongs to the IL6 cytokine receptor (IL6R) family. IL6Rs have divergent effects on cell proliferation in different epithelial cell types. While suppression of proliferation has been described for IL6R mediated signalling in some epithelial cells,43 it enhances cell growth in other epithelial cell lines.44 45 Here, we demonstrate that IL31 at high doses and low cell density significantly decreases cell proliferation in the colorectal cancer cell line HCT116. Similarly, a very recent study demonstrated that IL31 was highly effective in suppressing proliferation by altering expression of cell cycle proteins including upregulation of p27Kip1 and downregulation of cyclinB1, cdc2, cdk6, mcm4, and Rb in lung epithelial cells.4 Here, we demonstrate that the antiproliferative effect of IL31 in HCT116 cells was lost in dense cell cultures and under certain conditions even reversed into a proliferation stimulating effect. A similar bimodal effect on cell proliferation was previously described for IL31 in other epithelial cell types.46–48

Figure 6 IL31 mRNA expression is not increased in murine models of Th1 mediated intestinal inflammation. (A) IL31 expression was similar in ileal and colonic epithelial cells of TNFΔARE mice (n = 9) compared to C57BL/6 wild type mice (n = 9). Ileal mRNA expression in wild type mice was arbitrarily set to 1.0 and all other expression levels were calculated proportionally as fold increase. Data are presented as mean (SEM). (B) OSMR mRNA expression was not significantly different in TNFΔARE mice compared to wild type mice (p = 0.26). (C) TNF-α mRNA expression was significantly higher in TNFΔARE mice compared to wild type mice (p < 0.01). (D) IL31, IL31Rα, and OSMR mRNA expression were not significantly increased in C57BL/6 mice infected with murine cytomegalovirus (n = 10) compared to uninfected C57BL/6 mice (n = 4).

Figure 7 Colonic IL31 mRNA expression is increased in human inflammatory bowel disease. (A, C) IL31 expression is increased 2.4-fold in inflamed colonic biopsies from patients with Crohn’s disease compared to uninfamed biopsies while IL31Rα and OSMR expression were increased twofold. IL31 mRNA expression correlated with the IL8 expression (r = 0.56), which was measured as a marker for inflammation. (B, D) In patients with ulcerative colitis, a 2.6-fold increased IL31 mRNA expression was observed in inflamed biopsies in comparison to uninfamed tissue with a high correlation to IL8 mRNA levels (r = 0.65). IL31Rα and OSMR were increased 1.5-fold and 1.7-fold, respectively.
proliferation dependent on cell density has also been described for other cytokines such as TGF-β. However, a number of studies demonstrated that unstimulated cells in dense cultures undergoing contact inhibition are characterised by similar alterations of the expression of cell cycle proteins as caused by IL31 stimulation4 which includes downregulation of cyclins and upregulation of p27Kip1, which explains the relative loss of the antiproliferative effect of IL31 (in comparison with unstimulated cells) in dense cultures considering the overall low cell proliferation in (unstimulated) dense cultures. Similar to IL31, other proliferation inhibiting agents such as bone morphogenetic protein (BMP-7) and the MEK inhibitors PD98059 and U0126 have much higher effects on cell proliferation inhibition at low cell density compared to dense cultures. In contrast, cell proliferation stimulating cytokines such as hepatocyte growth factor (HGF) lose their proliferation stimulating properties in cell cultures with low cell density. It has been demonstrated that a single STAT3 recruitment site in the cytoplasmic domain of IL31Rα exerts a dominant function in the entire receptor complex and is critical for gene induction and growth inhibition. Similarly, we demonstrated upregulation of STAT3 following IL31 stimulation in our study. However, in the initial study, an IL31Rα overexpressing cell line was used explaining the stronger antiproliferative effects of IL31 observed in that study compared to our experiments. In addition, it has been demonstrated that high cell density itself is associated with ligand independent STAT3 activation mediating growth inhibition, which further explains the loss of the antiproliferative properties of IL31 in dense cell cultures. Moreover, we demonstrate that IL31 and IL31 receptor mRNA expression are cell differentiation dependent, which further supports the strong influence of cell density on the biological functions of IL31.

In this study, we demonstrate that proinflammatory cytokines increase the expression of IL31 and its receptor complex. The strongest upregulation of IL31, IL31Rα, and OSMR gene expression was seen after stimulation with IFN-γ. Similarly, in monocytes, gene upregulation of IL31Rα was stronger following IFN-γ compared to LPS stimulation. Given our quantitative PCR results of primary murine IEC, which demonstrated in comparison to colorectal cancer derived IEC only weak IL31Rα mRNA expression, it is therefore likely that proinflammatory cytokine activated monocytes are a major IL31Rα expressing cell type and target of IL31 in IBD.

Depending on the T cell source of cytokine production, cytokines have been differentiated in Th1 and Th2 cytokines. It has been proposed that CD represents a Th1 and Th17 mediated intestinal inflammation, while ulcerative colitis resembles more a Th2 mediated colitis. IL31 is mainly produced by activated Th2 cells1 and a very recent study implicates IL31R signalling as a novel negative regulatory pathway that specifically limits Th2 mediated inflammation while most colorectal cancer derived IEC expressed considerable levels of IL31Rα, its mRNA expression levels in unstimulated murine primary IEC were low suggesting that cancerogenesis might be another process in which this novel cytokine is involved. This is supported by IL31Rα expression in other malignancies such as glioblastoma and melanoma, which also secrete a soluble form of OSMR which may bind IL31 when combined with soluble IL31Rα, but further studies are required to elucidate potential functions of IL31 in cancerogenesis.

In summary, we demonstrated that IEC express the functional IL31 receptor complex. Binding of IL31 to its surface receptor complex in IEC leads to phosphorylation of STAT1/3, Akt, and ERK MAP kinases. IL31 inhibited cell proliferation in low density cell cultures and increased IEC migration. Moreover, the mRNA expression of IL31 was upregulated in inflamed colonic lesions of IBD patients but not in a Th1 mediated model of murine ileitis. Taken together, our data indicate a role for this cytokine in promoting proinflammatory gene expression and modulating IEC barrier function suggesting a role in the pathogenesis of IBD.

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