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## Anti-inflammatory effects of antibacterials on human bronchial epithelial cells

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### Abstract

**Background:** Human Bronchial epithelial cells (hu-BEC) have been claimed to play a significant role in the pathogenesis of chronic inflammatory airway diseases like COPD. In this context IL-8 and GM-CSF have been shown to be key cytokines. Some antibiotics which are routinely used to treat lower respiratory tract infections have been shown to exert additional immunomodulatory or anti-inflammatory effects. We investigated whether these effects can also be detected in hu-BEC.

**Methods:** Hu-BEC obtained from patients undergoing lung resections were transferred to air-liquid-interface (ALI) culture. These cultures were incubated with cefuroxime (CXM, 10-62.5 mg/l), azithromycin (AZM, 0.1-1.5 mg/l), levofloxacin (LVX, 1-8 mg/l) and moxifloxacin (MXF, 1-16 mg/l). The spontaneous and TNF- $\alpha$  (10 ng/ml) induced expression and release of IL-8 and GM-CSF were measured using PCR and ELISA in the absence or presence of these antibiotics.

**Results:** The spontaneous IL-8 and GM-CSF release was significantly reduced with MXF (8 mg/l) by  $37 \pm 20\%$  and  $45 \pm 31\%$ , respectively (both  $p < 0.01$ ). IL-8 release in TNF- $\alpha$  stimulated hu-BEC decreased by  $16 \pm 8\%$  ( $p < 0.05$ ) with AZM (1.5 mg/l). With MXF a concentration dependent decrease of IL-8 release was noted up to  $39 \pm 7\%$  ( $p < 0.05$ ). GM-CSF release from TNF- $\alpha$  stimulated hu-BEC was maximally decreased by  $35 \pm 24\%$  ( $p < 0.01$ ) with MXF (4 mg/l).

**Conclusion:** Using ALI cultures of hu-BEC we observed differential effects of antibiotics on spontaneous and TNF- $\alpha$  induced cytokine release. Our data suggest that MXF and AZM, beyond bactericidal effects, may attenuate the inflammatory process mediated by hu-BEC.

### Background

Antimicrobial agents of different classes - e.g. beta-lactams, quinolones, and macrolides - are standard of care in the treatment of respiratory tract infections. In addition to their antimicrobial activity some of these anti-

biotics, especially macrolides and fluoroquinolones, have immunomodulatory effects [1-3]. These anti-inflammatory or immunomodulatory capabilities have been demonstrated in human cells, cell lines, and in animal experiments [1,4-7].

Due to intracellular accumulation of macrolides and quinolones in lung cells and in alveolar macrophages a targeted modulation of the inflammatory reaction could be of additional therapeutic benefit by attenuation of the inflammatory process in lower respiratory tract infection (LRTI) as well as in chronic non-infectious airway diseases like COPD [8-10].

Airway epithelial cells have been shown to be of crucial importance in the pathogenesis of inflammatory airway diseases [11]. In addition to antimicrobial activities, macrolides directly affect pulmonary host defence like the neutrophil activation and the immune cell function. These effects are mediated by an alteration of cytokine and chemokine release, as has been demonstrated *in vitro* and *ex vivo* [2,12]. Moreover, macrolides like azithromycin are already clinically used in chronic respiratory diseases like diffuse panbronchiolitis (DPB), cystic fibrosis despite they have no antimicrobial activity against *Pseudomonas aeruginosa*. A beneficial effect on bacterial virulence factors by inhibiting quorum-sensing, a mechanism of bacterial communication, is described for macrolides and quinolones as well [13-17].

Additionally, immunomodulatory effects of macrolides are used in bronchiolitis obliterans syndrom after bone marrow transplantation and lung transplantation which are diseases without infectious background [12,18,19]. There are many studies, which elucidated the immunomodulatory effects of macrolides in human cells [20,21]. However, the underlying intracellular mechanisms of immunomodulation by macrolides are not completely understood yet [20,21].

Similarly to macrolides, immunomodulatory effects have been shown for fluorquinolones in a variety of cells of the immune system and in lung epithelial cells. These effects were especially pronounced in fluorquinolones with a cyclopropyl-moiety at position N1 like ciprofloxacin and moxifloxacin [1]. Moreover, expression of pro-inflammatory cytokines in human monocytes is suppressed by moxifloxacin *in vitro* and *in vivo* in an animal model of inflammation [4,7]. Beside the modulation of cytokine release from cells of the immune system it has been shown, that quinolones reduce pro-inflammatory activities of respiratory epithelial cell lines, thus potentially influencing pulmonary host defence [5,6].

Therefore, we investigated the modulation of cytokine release from primary human bronchial epithelial cells in air-liquid interface culture by different antibiotics.

## Methods

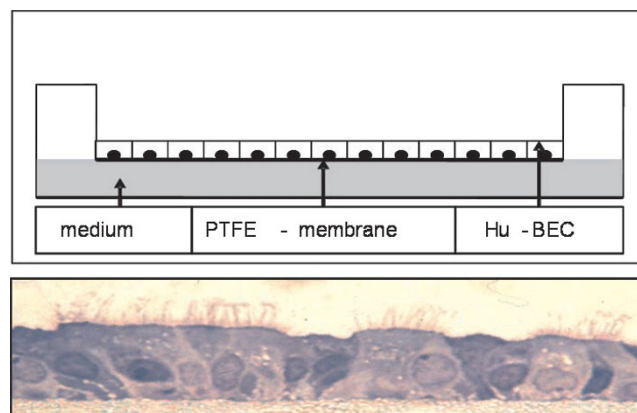
### Preparation of air-liquid interface cultures of human bronchial epithelial cells (hu-BEC)

The human bronchial epithelial cells were harvested from patients undergoing lung surgery for cancer resection or

transplantation [22,23]. Written informed consent was obtained from each patient according to the recommendations of the local ethic committee and there was an approval of our institutional review board. After preparation the resected bronchi were incubated for 24 h at 4 °C in DMEM (Dulbeccos Modified Eagle Medium, Invitrogen, USA) and DTT (Dithio-Threitol, Invitrogen, USA) containing penicillin G (Jenapharm, Germany), streptomycin (Rotexmedica, Germany), gernebcin (Infectopharm, Germany), imipenem (MSD, Germany) and amphotericin b (Bristol-Myer-Sqibb, Germany). Thereafter the bronchi were treated with protease Type XIV (Sigma, Germany) for 24 h at 4 °C and rinsed several times with DMEM to wash out the epithelial cells. Then the cells were grown to 80% confluence with airway epithelial cell growth medium (Promocell, Germany) and after treatment with trypsin (0.05%, Invitrogen, USA) the cells were transferred on a collagenised PTFE membrane (polytetrafluorethylen, Millipore, USA) of 6-well plates (Corning Costar, USA) at a concentration  $2 \times 10^6$  cells/ml and grown with DMEM containing HAM-12 (Invitrogen, USA), Ultrosor G (Pall Life Sciences, France) and antibiotics (penicillin 100 U/ml and streptomycin 100 µg/ml, Invitrogen) at 37 °C in 5% carbon dioxide/air. The supernatant was removed after 2 days and the cells were air-lifted. After another  $14.0 \pm 2.6$  days these air-liquid-interface cultured cells expressed their characteristic bronchial polarity (see Fig. 1). Cultures were considered confluent and differentiated if the Rt was stable and  $> 500 \Omega/\text{cm}^2$  measured by Ohmmeter (EVOM, World Precision Instruments, USA).

### Incubation experiments

To characterize spontaneous cytokine-expression and release of hu-BEC, we incubated air-liquid-interface (ALI) cultures with buffer or with cefuroxime (62.5 mg/l), azi-



**Figure 1**  
air-liquid-interface-culture (schematic); HE-stain of an air-liquid-interface-culture with characteristic polarity.

thromycin (1.5 mg/l), levofloxacin (8 mg/l), and moxifloxacin (8 mg/l) for 24 h. Thereafter, the basolateral medium of each well was collected and frozen at -20°C. The cells were lysed with Trizol (GIBCO, Germany) and the lysates were stored at -80°C.

To investigate cytokine-expression and release of hu-BEC under pro-inflammatory conditions the ALI cultures cells were pre-incubated for 24 hours with buffer or with various antibiotics at different concentrations (see Table 1) and stimulated with TNF-alpha (10 ng/ml) for another 24-h incubation. Thereafter, the basolateral medium of each well was collected and frozen at -20°C. The cells were lysed with Trizol reagent (GIBCO, Germany) and the lysates were frozen at -80°C.

To determine, whether there is a concentration-dependent effect of these antibiotics, we used a range of concentrations (see Table 1), which are reached in humans *in vivo* covering therapeutic levels in human serum, in bronchoalveolar lavage fluid, or in bronchial tissue [8,9].

Moxifloxacin and azithromycin was a generous gift from Bayer Healthcare Germany and Pfizer Germany. Cefuroxime and levofloxacin were purchased from Sanofi-Aventis (France) and DeltaSelect (Germany), respectively.

#### ELISA

IL-8 and GM-CSF were measured in basolateral medium using enzyme-linked immunosorbent assays (ELISA) (both R&D Systems, USA) as previously described [24].

#### RNA Extraction

RNA was extracted with Trizol according to the methods recommended by the manufacturer and frozen at -80°C. For analysis frozen epithelial cell lysates were re-dissolved in water. Total RNA yield was calculated by measuring the absorbance at 260 and 280 nm (assuming that  $A_{260}$  of 1 = 40 µg RNA). RNA integrity was judged by determining the ratio of  $A_{260}/A_{280}$ . Only samples with an  $A_{260}/A_{280}$  ratio from 1.6 to 2.0 were used for the subsequent measurements.

#### First-strand complementary deoxyribonucleic acid synthesis by reverse transcription

The RNA was transferred in cDNA with the cDNA synthesis kit (Fermentas, Germany) following the instruction of the manufacturer. The first-strand cDNA was stored at -80°C.

#### Semiquantitative polymerase chain reaction

A sample of 1 µl of cDNA was used for each 20 µl PCR reaction. Primer sets used for the amplification of cytokines and the housekeeping gene glyceraldehyde-3-phosphate dehydrogenase (GAPDH) were as follows:

GAPDH (MWG-Biotech, Germany): Forward: 5'-TGA AGG TCG GAG TCA ACG GAT TTG GT-3'; Reverse: 5'-CAT GTG GGC CAT GAG GTC CAC CAC-3', (size of PCR product: 900 base pair [bp]).

IL-8 (MWG-Biotech, Germany): Forward: 5'-ATT TCT GCA GCT CTG TGT GAA-3'; Reverse: 5'-TGA ATT CTC AGC CCT CTT CAA-3', (size of PCR product: 255 bp).

**Table 1: Concentrations of cefuroxime (CXM), azithromycin (AZM), levofloxacin (LVX) and moxifloxacin (MXF) used for incubation experiments**

	Concentration	TNF-α(10 ng/ml)	IL-8	GM-CSF	IL-8 PCR	GM-GSF PCR
CXM	10 mg/l	+	n = 11	n = 11	-	-
	25 mg/l	+	n = 17	n = 17	-	-
	62.5 mg/l	+	n = 11	n = 11	n = 11	n = 10
	62.5 mg/l	-	n = 11	n = 11	n = 11	n = 10
AZM	0.5 mg/l	+	n = 12	n = 10	-	-
	1.0 mg/l	+	n = 12	n = 10	-	-
	1.5 mg/l	+	n = 12	n = 10	n = 10	n = 11
	1.5 mg/l	-	n = 11	n = 8	n = 11	n = 11
LVX	1 mg/l	+	n = 12	n = 11	-	-
	4 mg/l	+	n = 16	n = 18	-	-
	8 mg/l	+	n = 11	n = 11	n = 11	n = 11
	8 mg/l	-	n = 10	n = 11	n = 11	n = 11
MXF	1 mg/l	+	n = 24	n = 22	-	-
	4 mg/l	+	n = 25	n = 23	-	-
	8 mg/l	+	n = 23	n = 23	-	-
	16 mg/l	+	n = 14	n = 14	n = 14	n = 14
	8 mg/l	-	n = 11	n = 11	n = 11	n = 11

GM-CSF (MWG-Biotech, Germany): Forward: 5'-ACA CTG CTG CTG AGA TGA ATG AAA CAG TAG-3', Reverse: 5'-TGG ACT GGC TCC CAG CAG TCA AAA GGG ATG-3', (size of PCR product: 286 bp).

Each 50-  $\mu$ l reaction mixture consisted of 5  $\mu$ l of 10 $\times$  PCR buffer, 1.5  $\mu$ l MgCl<sub>2</sub> (~1.5 mM), 1  $\mu$ l of 10 mM dNTP mix, 5  $\mu$ l of specific primer for GAPDH, the mediators (synthesized by MWG-Biotech, Germany) (~10  $\mu$ M), 0.25  $\mu$ l of *Taq* DNA Polymerase (GIBCO, Germany) (~2 U), and 37.25  $\mu$ l of H<sub>2</sub>O. The cycles (Peltier Thermal Cycler, MJ Research, USA) used were as follows: GAPDH: 94 °C for 3 min/94 °C for 45 sec/60 °C for 30 sec/72 °C for 90 sec for 25 cycles, followed by an extension step of 10 min at 72 °C. The same cycle conditions were used for the mediators. The annealing temperature and PCR cycles for the mediators were as follows: IL-8 58 °C for 35 cycles; GM-CSF 65 °C for 40 cycles.

Products of amplification were transferred on a 2% agarose gel and after electrophoresis viewed using a 300-nm ultraviolet transilluminator (Cybertech, Germany). Samples from RT reactions that did not contain RT served as negative controls. For quantification, PCR bands were stained with ethidium bromide (Sigma, Germany) and signal intensity was measured with an ultraviolet densitometer (Cybertech, Germany). Densitometric values are expressed as the ratio of IL-8/GAPDH and GM-CSF/GAPDH.

### Statistical analysis

All statistical analyses were performed with SPSS 11.5 (Chicago, USA). The results are expressed as mean values  $\pm$  SEM. We applied a non-parametric Wilcoxon-Test in our exploratory analysis. Conventionally,  $p < 0.05$  was considered significant. The correlations of the data obtained by ELISA and PCR were calculated using the Pearson's test.

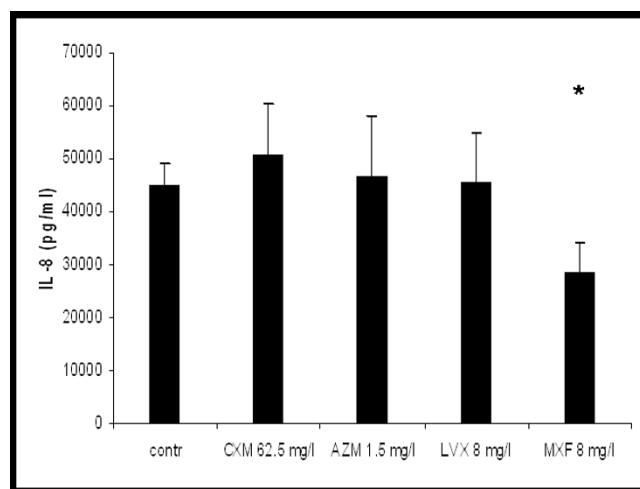
## Results

### Effects on spontaneous IL-8 release

Spontaneous IL-8-release of hu-BEC in ALI cultures was  $44.7 \pm 4.3$  ng/ml. No significant changes were observed with CXM (62.5 mg/l), AZM (1.5 mg/l), and LVX (8 mg/l). After 24 h incubation with MXF (8 mg/l) IL-8 release was reduced by  $37 \pm 20\%$  ( $p < 0.008$ ) (fig. 2).

### Effects on TNF- $\alpha$ stimulated IL-8 release

Stimulation with TNF- $\alpha$  resulted in a 3.4-fold increase of IL-8 release to  $160.2 \pm 6.4$  ng/ml ( $p < 0.001$ ). Incubation with cefuroxime at a concentration of 62.5 mg/l led to a significant further increase of IL-8 release in stimulated hu-BEC by  $33 \pm 6\%$  ( $p < 0.013$ ). Under stimulated conditions azithromycin showed a significant reduction of IL-8 production up to  $16 \pm 8\%$  at a concentration of 1.5 mg/l ( $p < 0.016$ ). No significant changes were observed with



**Figure 2**  
Effect of cefuroxime (CXM), azithomycin (AZM), levofloxacin (LVX) and moxifloxacin (MXF) on spontaneous IL-8 release from hu-BE, \* $p < 0.05$  vs. control.

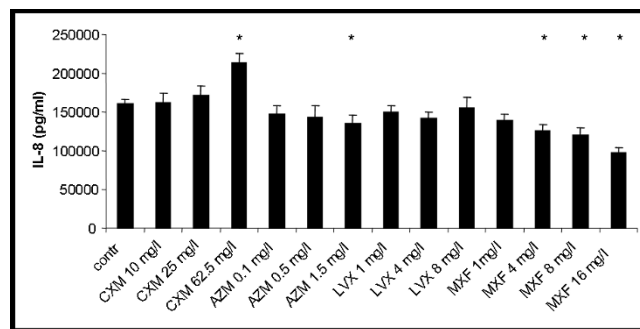
levofloxacin at concentrations of 1, 4, and 8 mg/l. Incubation with moxifloxacin led to a concentration dependent reduction of IL-8 release to a maximum of  $39 \pm 7\%$  ( $p < 0.001$ ) at a concentration 16 mg/l (see Fig. 3).

### Effects on spontaneous GM-CSF release

Spontaneous GM-CSF-release of hu-BEC in ALI cultures was  $654 \pm 108$  pg/ml. Incubation with CXM, AZM, or LVX did not show a significant effect on GM-CSF release with all concentrations tested. Only MXF reduced GM-CSF release by  $45 \pm 31\%$  ( $p < 0.004$ ) (see Fig. 4).

### Effects on TNF- $\alpha$ stimulated GM-CSF release

Stimulation with TNF- $\alpha$  did not significantly alter GM-CSF release from hu-BEC in ALI cultures (maximum effect  $+17 \pm 7\%$ , n.s.). GM-CSF release of TNF- $\alpha$  stimulated hu-



**Figure 3**  
Effect of cefuroxime (CXM), azithomycin (AZM), levofloxacin (LVX) and moxifloxacin (MXF) on TNF- $\alpha$ -stimulated IL-8-release; \*  $p < 0.05$  vs. control.

BEC in ALI cultures was also not significantly influenced by incubation with different concentrations of CXM, AZM, or LVX. Only MXF inhibited GM-CSF release in TNF- $\alpha$ -stimulated hu-BEC with an inverse concentration response characteristic (fig. 5). MXF concentration of 4 mg/l reduced GM-CSF release by  $35 \pm 24\%$  ( $p < 0.009$ ), MXF 8 mg/l reduced GM-CSF release by  $30 \pm 23\%$  ( $p < 0.013$ ), and MXF 16 mg/l reduced GM-CSF release by  $22 \pm 31\%$  ( $p < 0.019$ ) (fig. 5).

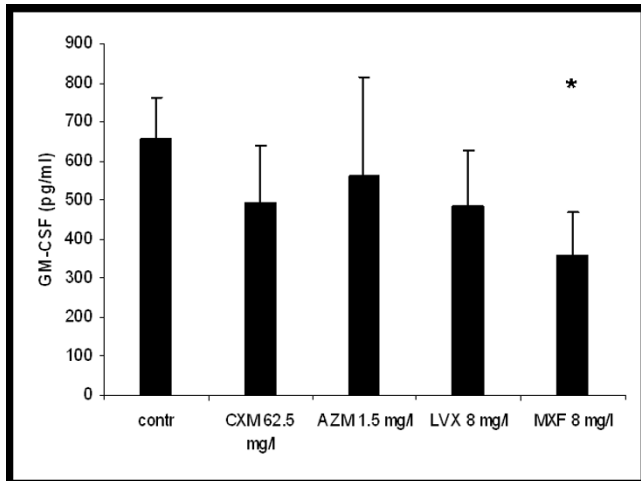
**PCR Analyses**

Spontaneous IL-8 mRNA/GAPDH ratio was  $1.2 \pm 0.06$  in the semi-quantitative PCR. The IL-8 mRNA/GAPDH ratio was reduced by  $21 \pm 8\%$  after incubation with 8 mg/l MXF in unstimulated cells. Smaller effects were observed with CXM, AZM, or LVX (all n.s.).

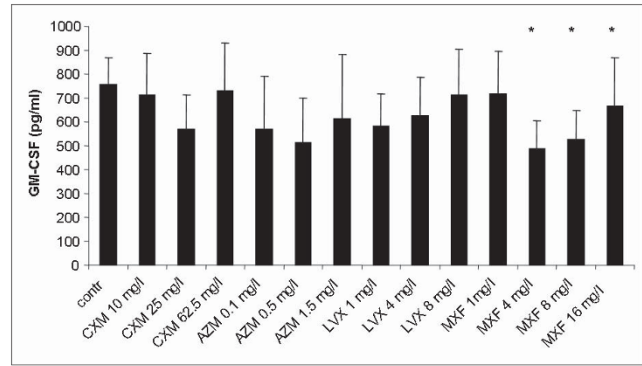
In TNF- $\alpha$  stimulated hu-BEC IL-8 mRNA/GAPDH ratio increased to  $1.54 \pm 0.06$  ( $p < 0,001$ ). Incubation with CXM, AZM, LVX, or MXF led to maximal changes of IL-8/GAPDAH ratio of  $-15 \pm 8\%$ ,  $+5 \pm 12\%$ ,  $-8 \pm 10\%$ , and  $-11 \pm 7\%$ , respectively (all n.s.).

The spontaneous and TNF- $\alpha$  stimulated GM-CSF/GAPDH-ratio of hu-BEC in ALI cultures was  $1.64 \pm 0.08$  and  $1.81 \pm 0.08$ , respectively. Incubation with CXM, AZM, LVX, or MXF did not significantly alter GM-CSF/GAPDH-ratio at all concentrations investigated (n.s.).

Correlation analysis revealed weak correlations between IL-8 protein as measured by ELISA and IL-8 m-RNA/GAPDH ratio ( $r = 0.373$ ,  $p < 0.001$ ) as well as between



**Figure 4**  
Effect of cefuroxime (CXM), azithomycin (AZM), levofloxacin (LVX) and moxifloxacin (MXF) on spontaneous GM-CSF release from hu-BE,\* $p < 0.05$  vs. control.



**Figure 5**  
Effect of cefuroxime (CXM), azithomycin (AZM), levofloxacin (LVX) and moxifloxacin (MXF) on TNF- $\alpha$ -stimulated GM-CSF-release; \*  $p < 0.05$  vs. control.

GM-CSF protein and GM-CSF mRNA/GAPDH ratio ( $r = 0.209$ ;  $p < 0.004$ ).

**Discussion**

The data presented here demonstrate that some antibiotics are capable of modifying the inflammatory activation of human bronchial epithelial cells. The differential effects observed between different groups of antibiotics suggest that the member of the cephalosporine group cefuroxime do not show this effect, whereas azithromycin and moxifloxacin exert anti-inflammatory effects on hu-BEC, with moxifloxacin suppressing IL-8 and GM-CSF with and without TNF- $\alpha$  stimulation in our experimental setting, whereas AZM decreased IL-8 only after stimulation with TNF- $\alpha$  and had no significant effect on GM-CSF.

Immunomodulatory effects of antibiotics have been described so far *in vivo* with animal models, *in vitro* with models of immune cells, NHBE cells (normal human bronchial epithelial cells) and immortalised respiratory cell lines [1,4-7]. In those experiments it could be demonstrated that MXF leads to a reduction of IL-8, TNF- $\alpha$ , IL-1 $\alpha$ , IL-1 $\beta$ , IL-4 and IFN- $\gamma$  in monocytes, lymphocytes and neutrophils after stimulation with different agents. The direct effect on GM-CSF has not been investigated yet in cells of the immune system. However, in a mouse model of bone-marrow ablation with cyclophosphamide MXF leads to an increase of WBC and GM-CSF was augmented in the lungs of these mice [25]. In contrast, our findings suggest that MXF reduces spontaneous and TNF- $\alpha$  stimulated GM-CSF production and release of hu-BEC. This could be related to the different models and the different stimuli used.

For lung cells, only data of A549 cells, an immortalized type II alveolar epithelial cell line and IB3 cells, a cystic fibrosis cell line, are published [5,6]. In IB 3 cells MXF

reduced the release of IL-8 and other cytokines [6]. In A549 cells MXF decreases NO production and NF- $\kappa$ B-activation [5]. Our study demonstrates anti-inflammatory effects of quinolones in a human *ex vivo* model of primary bronchial epithelial cells. The concentrations of the different antibiotics employed were comparable to concentrations reached by therapeutic medication in humans. Using primary hu-BEC in ALI cultures and therapeutically relevant concentrations of different antibiotics suggest that these findings may be also clinically relevant and may have implications for the treatment of human lung diseases.

In our study, we investigated the effect of different antibiotics on IL-8 and GM-CSF after application of TNF- $\alpha$  as an inflammatory stimulus. TNF- $\alpha$  is a proinflammatory cytokine with pro-fibrotic features which has a key role in lower respiratory tract infections as well as in chronic inflammatory lung disease like asthma, bronchiolitis obliterans, or COPD [11,26,27]. A blockade of TNF- $\alpha$  led to decrease of IL-8 after stimulation with LPS in lungs of patient with COPD [27].

Similarly, IL-8 and GM-CSF are key mediators not only in acute infectious inflammation but also in chronic inflammation as observed in COPD, bronchial asthma, and bronchiolitis obliterans [24,26,28]. IL-8 is rapidly induced by an inflammatory stimulus like TNF- $\alpha$  or LPS and is one of the most potent neutrophil chemoattractants in human tissue [27,29]. GM-CSF leads to an activation and increased survival of leukocytes and enhance oxidative burst in the lungs, thus maintaining and prolonging inflammatory reactions [28].

As we and other have shown, IL-8 and GM-CSF are secreted locally by the respiratory epithelium [24-26,28,30]. However, there is no specific treatment yet in humans to directly address and modify these cytokines to suppress the inflammatory cascade.

Our observations suggest that some antibiotics may have the capability to block or modulate this inflammation. In our experiments we employed concentrations of AZM, LVX and MXF which were comparable to therapeutic concentrations of these antibiotics and are reached in human lungs *in vivo* [8,9]. The serum level after therapeutic doses of MXF and LVX is 1-5 mg/l and after oral administration concentrations reached in the epithelial lining fluid (ELF) are 5 - 7 times higher than serum levels [9]. After oral administration with AZM serum level is 0.10 mg/l and the concentration in the ELF ranges from 0.94 mg/l to 1.2 mg/l after oral administration [8]. However, AZM accumulates intracellularly in alveolar macrophages with a concentration of 205.24 mg/l 24 hours after the last intake under steady state conditions [8,10]. The concentration of

cefuroxime used in our experiments covers a range of serum and intrapulmonary concentrations after oral and continuous *i.v.* administration in humans [31-34]. Additionally we used a concentration of cefuroxime (62.5 mg/L) above these therapeutic intrapulmonary concentrations.

So far AZM and other macrolides are the only antibiotics used for therapeutic modulation of the local immune system in the lung. A beneficial effect of AZM has been demonstrated in the management of cystic fibrosis lung disease and diffuse panbronchiolitis (DPB) [2,3,18]. DPB is a disease observed predominantly in Asia, which without medical intervention leads to a rapid decline of lung function and death [3]. AZM is also used for treatment of bronchiolitis obliterans after organ transplantation, a chronic inflammatory and fibroproliferative disease leading to bronchiolar obstruction and obliteration of distal airspaces after lung transplantation but also after haematopoietic stem cell transplantation [35,36]. In our experiments only AZM at a concentration of 1.5 mg/l was associated with a significant reduction of IL-8 release. These findings differ from results in NHBE cells, a human bronchial epithelial cell line, where AZM at a concentration of 1.0 mg/l did not show an effect, whereas at a concentration of 10 mg/l an increase in IL-8-secretion was observed [37]. However, *in vivo* a concentration of 10 mg/l cannot be found under steady state conditions in ELF of the normal lung and was, therefore, not investigated in our experiments with hu-BEC. Hence, the immunomodulatory effects mediated by macrolides may not only depend on a direct effect on lung epithelial cells, but also on a direct effect on alveolar macrophages because of the intracellular accumulation in alveolar macrophages.

We also investigated effects on IL-8 mRNA and GM-CSF mRNA expression. In general the mRNA expressions of both, IL-8 and GM-CSF, were correlated with IL-8 and GM-CSF protein release, thus supporting the view that changes in protein release were related to changes in gene expression. However, the differences in mRNA expression between different experimental groups were not statistical significant. This could be due to the fact that changes of gene expression may be transient and are less well detected after 24-hours of incubation, when the cells were lysed and the mRNA isolated. In this respect further studies are needed to quantify the effect on mRNA-levels at earlier time points.

Although our data suggests that quinolones exert anti-inflammatory effects on hu-BEC, these effects are not uniform for all quinolones. In our experiments, moxifloxacin, a quinolone with a cyclopropyl-moiety at N1 (like ciprofloxacin) had a more pronounced effect on cytokine release when compared to levofloxacin, a qui-

nolone lacking this cyclopropyl-moiety at N1 [1]. Despite the above-mentioned anti-inflammatory effects a careful use of quinolones is recommended due to risk of cross-resistance.

Several intracellular signal transduction pathways mechanisms are thought to be responsible for these anti-inflammatory effects [1,4-6]. Yet these mechanisms are not completely understood. Previous studies have shown that pre-treatment with MXF leads to an inhibition of the MAP-Kinases ERK 1/2 and JNK in monocytes [4,38]. MXF also inhibits the phosphorylation of these kinases in IB3 cells, C38 cells and A549 cells [5,6]. In contrast, the MAP-kinase p38 was not influenced by MXF [6]. Additionally, in monocytes and respiratory cell lines MXF inhibits NF- $\kappa$ B-activation due to reduced I $\kappa$ -B degradation [38]. This prevents NF- $\kappa$ B activation and translocation to the nucleus and thus inhibits the cytokine cascade.

## Conclusion

Our data confirm previous studies showing a significant inhibitory effect of quinolones with a cyclopropyl-moiety at N1 on cytokine release. Our study adds new aspects by using primary hu-BEC in ALI cultures and by employing therapeutically relevant concentrations of different antibiotics. When compared to MXF, AZM showed smaller effects on IL-8 release and did not affect GM-CSF release in concentration which can be reached in human ELF. In contrast, LVX showed no significant effects on cytokine release and CXM led to an increase in IL-8 release. Therefore, MXF appears to be more potent as an anti-inflammatory substance in bronchial epithelial cells. However, the clinical relevance of these findings has not been evaluated yet.

## Competing interests

GSZ has received a travel fee and a fund for speaking at symposium organized on behalf of Bayer Healthcare in 2007. The other authors have none to declare.

## Authors' contributions

GSZ and HVH have carried out the experimental work. GSZ carried out the data analysis and drafted the manuscript. GSZ, JB and RH initiated the study and designed the experiments. CN participated in the design of the study. RH provided the surgical specimens. All authors read and approved the final version of the manuscript.

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