EGFR plays an essential role in lipopolysaccharide (LPS)-induced MUC5AC hypersecretion in human bronchial epithelial cells.

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Abstract

Background: Excessive mucus secretion is an important characteristic of airway inflammatory diseases, such as asthma. Lipopolysaccharide (LPS) leads to mucus hypersecretion in asthma, and is associated with airway inflammation and remodeling. It is known little about the role of epidermal growth factor receptor (EGFR) in LPS-induced mucin 5AC (MUC5AC) hypersecretion in human bronchial epithelial cells. We aim to investigate the effects of EGFR in the process of MUC5AC hypersecretion.

Methods: The human bronchial epithelial (16HBE) cells were treated with LPS, and pre-treated with AG1478 (specific inhibitor of EGFR). The expression of MUC5AC and EGFR were detected and compared between the different groups. The expression of MUC5AC protein was measured by enzyme linked immunosorbent assay (ELISA). The expressions of MUC5AC and EGFR were measured by immunohistochemistry, and the protein expression of EGFR was monitored by Western blot. The mRNA expression of MUC5AC and EGFR were detected by reverse transcription-polymerase chain reaction (RT-PCR).

Results and conclusions: LPS could induce the high expression of MUC5AC and EGFR in 16HBE cells, and AG1478 decreased the expression of MUC5AC by inhibiting the expression of EGFR. These results showed that EGFR might play an important role in LPS-induced MUC5AC hypersecretion of airway epithelial cells, and AG1478 decreased MUC5AC secretion through inhibiting the expression of EGFR. AG1478 may be a new therapeutic approach for the airway inflammatory diseases.

Keywords: Bronchial epithelial cell, Mucus secretion, Epidermal growth factor receptor, AG1478, Lipopolysaccharide.

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Introduction

Asthma is one of the systemic inflammatory diseases, which is characterized by airway remodeling and airway inflammation, airway smooth muscle cell hyperplasia, mucus hypersecretion. The bronchial epithelial cells are affected firstly during the pathophysiologic process of asthma, which are associated with the airway inflammation, airway hyper-responsiveness, and airway remodeling. Mucus hypersecretion is one of the pathophysiological features of asthma, which can lead to airway obstruction.

A variety of inflammatory mediators, pathogenic microorganisms and their metabolites, cold air, proteases can promote the secretion of mucus, which are the main components of mucus [1,2]. Bacterial infection is one of the main factors which can induce mucus hypersecretion, and result in glandular secretion, airway damage and airway remodeling. LPS is a major component of gram-negative bacteria wall and can increase mucin5AC (MUC5AC) expression in a variety of tissues and cells [3,4]. LPS can damage the airway epithelial cells, destroy the defence barrier of airway and induce MUC5AC hypersecretion.

MUC5AC is the major mucin produced by bronchial epithelial cells. It is regulated by several pathways in different type cells. It has been reported that epidermal growth factor receptor (EGFR) activation plays an important role in regulation of MUC5AC secretion in a variety of cells, such as human intrahepatic biliary epithelial cells, gallbladder epithelial cells and human bronchial epithelial cells [4-6]. EGFR plays an important role in the injury and repair processes of airway epithelial cells. We have found that chronic exposure to Aspergillus fumigatus up-regulated the expression MUC5AC and EGFR in asthmatic rats, but whether EGFR activation is involved in MUC5AC hypersecretion in the human bronchial...
epithelial cells needs to be investigated further [7, 8]. Based on the above reports, we postulate that EGFR pathway may be involved in the regulation of MUC5AC expression in human bronchial epithelial cells. In this study, human bronchial epithelial (16HBE) cells were cultured in vitro with LPS and AG1478. The expression of MUC5AC and EGFR were examined in the 16HBE cells.

Materials and Methods

Cell culture and reagents

16HBE cells were purchased from Rochen Pharma Co. Ltd., (Shanghai, China). The 16HBE cells were cultured in the RPMI1640 cell culture medium supplemented with 10% fetal bovine serum (FBS), 100 IU/mL penicillin, 100 μg/mL streptomycin, at 37°C in a 5% CO₂ humidified atmosphere. This study designed 4 groups: 1) Group A (normal control group): the 16HBE cells were cultured in the serum-free RPMI1640 for 24 h; 2) Group B: The 16HBE cells were treated with 100 μg/mL LPS (Sigma) in serum-free RPMI1640 for 24 h; 3) Group C: the 16HBE cells were treated with 5 μmol/L of AG1478 (Sigma) in serum-free RPMI1640 for 24 h; 4) Group D: The 16HBE cells were pre-treated with 5μmol/L AG1478 for 10 min and then treated with 100 μg/mL of LPS in serum-free RPMI1640 for 24 h. After the intervention by LPS and AG1478, the expression of MUC5AC and EGFR were measured in each group cells.

Detection of MUC5AC in the cell culture supernatant

After 24 h, the cell culture supernatant were collected at different time points (6 h, 12 h, 24 h), and centrifuged for 20 min at 1000 × g at 2-8°C. The level of MUC5AC in the supernatant was measured by Enzyme-linked Immunosorbent Assay (ELISA) (the sensitivity was 0.188 pg/ml) according to the instructions provided by manufactures (Elabscience, Wuhan, China). The optical density (OD) was detected at a wavelength of 450 nm, and the OD value was proportional to the concentration of MUC5AC.

Immunohistochemistry (IHC)

The expression of MUC5AC and EGFR were detected with IHC according to the instructions of the immunohistochemical kit. The 16HBE cells were fixed with Paraformaldehyde. Antigen retrieval was performed by incubating with the 1% BSA for 1 h at 37°C in the incubator after the elimination of endogenous peroxidase activity with 3% hydrogen peroxide for 10 min. The 16HBE cells were subsequently incubated with monoclonal anti-MUC5AC and anti-EGFR antibody (1:200, purchased from Abcam Co., Shanghai, China) respectively at 4°C overnight. The horseradish peroxidase labelled goat anti-mouse antibody was used as the secondary antibody reagent and then stained by dianinobenzidin (DAB) kit. The anti-MUC5AC and anti-EGFR antibody were substituted with PBS buffer as the negative control. The integrated optical density of positive stain in the 16HBE cells were detected by Image-Pro plus 6.0 Software.

Western-blot

The 16HBE cells were lysed in RIPA lysis buffer for 2 h at ice and centrifuged for 10 min at 4°C. After removal of cell debris by centrifugation (12,000 g, 10 min), equal amounts (40 μg) of proteins were separated by 10% SDS-PAGE and transferred onto nitrocellulose membrane. Nonspecific reactivity was blocked in 5% non-fat milk in TB-ST for 1 h at room temperature. The membrane was then incubated with monoclonal antibody of mouse anti-EGFR (Santa Cruz) followed by reaction with anti-mouse IgG-HRP antibody. Specific protein bands were photographed and the immune-quantification of reactive proteins was performed by ECL chemiluminescence system (Santa Cruz).

Reverse transcription-PCR analysis of MUC5AC mRNA and EGFR mRNA

Total RNA was extracted with trizol and reverse transcribed. The cDNA synthesis were performed according to the manufacturer’s instructions (SYBRGreen PCR kit, TaKaRa, China).The PCR reactions were performed at 94°C, degeneration for 3 min, followed by 40 amplification cycles at 94°C for 20 s, 55°C for 20 s, and 70°C for 20 s. GAPDH mRNA was used as an internal control to standardize the RNA input. Primer sequences for target genes in the studies are shown in Table 1. Relative quantitation of the MUC5AC mRNA and EGFR mRNA were obtained by the 2^{-ΔΔCT} method.

<table>
<thead>
<tr>
<th>Sequences</th>
<th>MUC5AC</th>
<th>EGFR</th>
<th>GAPDH</th>
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<tr>
<td>Forward</td>
<td>5'-CCACACCAGTCACCAGAAAC-3'</td>
<td>5'-GGCACGGTGTATAAGGGACT-3'</td>
<td>5'-TGGTATCGTGGAAGGACTCA-3'</td>
</tr>
<tr>
<td>Reverse</td>
<td>5'-GGTTTCCTTGTCCTCACCAT-3'</td>
<td>5'-ATCGAGGATTTCCTTGTTGG-3'</td>
<td>5'-CCAGTAGAGGCAGGGATGAT-3'</td>
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Statistical analysis

Measurement data were expressed as mean ± SEM and analyzed with SPSS 16.0 statistical software. The comparisons between different groups were performed by t-test. Statistical significance was set at P<0.05.

Results

LPS up-regulated the expression of MUC5AC in 16HBE cells

The IHC analysis shown that MUC5AC were expressed in the cytoplasm of human bronchial epithelial cells, and stained brown. LPS induced the significantly increasing expression of MUC5AC than control group cells (P<0.05), and the
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expression were down-regulated after treatment with AG1478 (P<0.05). The level of MUC5AC in the cell culture supernatant was also higher than normal control group (P<0.05). AG1478 decreased the expression of MUC5AC compared with group B. There is no difference between group A and group C (Figure 1 and Table 2).

Figure 1. The expression of MUC5AC in 16HBE cells (stained brown): The expression of MUC5AC in group B was enhanced significantly compared with group A, and the effect of LPS on MUC5AC hypersecretion was impaired by AG1478 in group D. Original magnification, X400.

The expression of MUC5AC mRNA in different groups

The level of MUC5AC mRNA in the 16HBE cells significantly increased after treating with LPS compared with control group, and AG1478 decreased its higher expression induced by LPS. AG1478 did not decrease the expression of MUC5AC mRNA in group C versus group A (Figure 2).

Figure 2. The expression of MUC5AC mRNA levels in each group. Data are expressed as mean ± SEM. **P<0.01 versus Group A, dP<0.05 versus Group B.

LPS increased the expression of EGFR in 16HBE cells

The western blot analysis revealed that LPS up-regulated the expression of EGFR proteins in the 16HBE cells, and it could be inhibited by AG1478. The results of IHC also shown that EGFR were activated by LPS in the 16HBE cells, and its effect was inhibited by AG1478 (Figures 3-5).

The expression of EGFR mRNA increased by LPS

The mRNA level of EGFR increased significantly after intervention with LPS in the 16HBE cells, and its higher expression could be decreased by AG1478 (Figure 6).

Discussion

Mucus is the main defence barrier against the damage factors in the airway. Mucus hypersecretion is an important characteristic of airway inflammatory diseases, such as asthma, chronic obstructive pulmonary disease and cystic fibrosis. MUC5AC is a major component of mucus in airway epithelial cells and regulated by means of an epidermal growth factor receptor (EGFR)-signaling pathway [9]. A variety of inflammatory mediators, pathogenic microorganisms and their metabolites, cold air, proteases can promote the secretion of MUC5AC, and then induce glandular secretion and airway obstruction, which are the main factors of cough and sputum.
LPS is one of the initiators of inflammation, which is the main component of gram-negative bacteria cell walls. It can result in mucus hypersecretion, airway inflammation and oxidative stress etc. There were many reports that LPS increased the higher expression of MUC5AC in human intrahepatic biliary epithelial cells by up-regulating the expression of EGFR [3,4,9]. In the present study, we found that LPS up-regulated the MUC5AC expression in 16HBE cells, and promoted the expression of EGFR at the same time. We speculated that EGFR may play an important role in the process of mucus hypersecretion.

Table 2. The levels of MUC5AC in different groups (ng/ml) (ELISA).

<table>
<thead>
<tr>
<th>Group</th>
<th>6 h</th>
<th>12 h</th>
<th>24 h</th>
</tr>
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<tbody>
<tr>
<td>Group A</td>
<td>19.48 ± 0.36</td>
<td>18.41 ± 0.69</td>
<td>19.96 ± 0.92</td>
</tr>
<tr>
<td>Group B</td>
<td>31.41 ± 1.42*</td>
<td>66.78 ± 2.47*</td>
<td>80.68 ± 3.19*</td>
</tr>
<tr>
<td>Group C</td>
<td>17.93 ± 0.86</td>
<td>18.04 ± 0.61</td>
<td>18.92 ± 0.18</td>
</tr>
<tr>
<td>Group D</td>
<td>26.44 ± 0.36Δ</td>
<td>56.30 ± 1.17Δ</td>
<td>61.96 ± 0.94Δ</td>
</tr>
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Note: *P<0.05 versus group A; ΔP<0.05 versus group B.

EGFR is existed in the basal cells, secretory epithelium and interstitial cells of the lung with little expression at normal conditions. It is involved in the regulations of growth and differentiation of the cells. Studies have shown that EGFR expressed highly in some diseases, such as non-small cell lung cancer, asthma and chronic obstructive pulmonary disease. Asthma is a chronic inflammatory disease and involved in a variety of cells and cell components during its pathological process, which resulted in airway remodeling and airway inflammation including goblet cell and airway smooth muscle cell hyperplasia, mucus hypersecretion and so on. EGFR plays an essential role in injury and repair process of the tissue and cells during the pathophysiologic process of asthma. Studies have shown that EGFR was expressed higher in the airway epithelial cells and airway smooth muscle cells in patients with asthma than normal person [10]. The expression intensity of EGFR was positively correlated with the damage of airway epithelial cells, which may increase hormone resistance in patients with asthma. EGFR can be activated by several potent ligands and increase the MUC5AC expression in different cell types [4,11-16]. We found that LPS could activate EGFR and increase the MUC5AC expression in the 16HBE cells. These studies suggest that high expression of EGFR may be one reason for mucus hypersecretion in airway inflammatory diseases, which is accordance with our results.

AG1478 is one of tyrosinekinase inhibitors, which can inhibit the activity of EGFR tyrosine kinase with high selectivity. Studies have shown that AG1478 inhibited the activity of EGFR through binding the ATP competitively, and then blocked signal pathways mediated by EGFR, which can result in cell apoptosis at a dose-dependent manner [12]. AG1478 significantly inhibited the secretion of MUC5AC and IL-8 of NCI-H292 cells induced by LPS [13]. Shown in a study in vivo, intranasal instillation of AG1478 for an hour could significantly inhibit goblet cell metaplasia, mucus production, and neutrophil infiltration after intranasal LPS instillation in rat nasal epithelium. Our results showed that AG1478 could significantly decrease EGFR expression in the 16HBE cells and inhibit LPS-induced MUC5AC overexpression, indicating EGFR playing an important role in MUC5AC overexpression induced by LPS in the 16HBE cells. In the present study we found that blocking EGFR can inhibit LPS-induced MUC5AC hypersecretion in the 16HBE cells, indicating the potential target function of EGFR.

In summary, our results suggested that EGFR play an essential role in LPS-induced mucus hypersecretion and its inhibitor AG1478 could prevent the excessive mucus secretion, and may be used as an effective pharmacotherapy for airway inflammatory diseases, such as asthma.

Declaration of Interest

The authors report no conflicts of interest. The authors alone are responsible for the content and writing the paper.

References

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