Interleukin-6 promotes tumor progression in colitis-associated colorectal cancer through HIF-1α regulation

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Abstract. Interleukin-6 (IL-6) is a well-known etiological factor of colitis-associated colorectal cancer (CAC) and has a significant role in CAC progression. In addition, hypoxia-inducible factor 1α (HIF-1α) serves a primary role in the progression of CAC. However, the association between IL-6 and HIF-1α during the progression of CAC remains unclear. To investigate this association, the present study induced CAC in a mouse model using azoxymethane and dextran sulfate sodium. In addition, an anti-IL-6 receptor antibody was used to inhibit IL-6. In this model, anti-IL-6 receptor antibody treatment significantly inhibited the development of CAC and the expression of HIF-1α, in colorectal adenomas and adenocarcinomas. In patients with CAC, the HIF-1α gene was demonstrated to be overexpressed in tumor tissue compared with adjacent non-malignant tissue. Furthermore, HIF-1α mRNA expression was positively correlated with serum IL-6 concentration. The results of the present study suggest that IL-6 promotes CAC progression, in the early stage of the disease, through HIF-1α regulation.

Introduction

Colitis-associated colorectal cancer (CAC) is a subtype of colorectal cancer that is associated with inflammatory bowel diseases (IBD), including ulcerative colitis (UC) and Crohn’s disease (1). Over 20% of patients with IBD develop CAC within 30 years of the disease onset and >50% of these will succumb to CAC (2). The primary risk factors for CAC progression are the duration and severity of intestinal inflammation (3). Increasing evidence supports the importance of pro-inflammatory cytokines in CAC progression (4-7). Interleukin-6 (IL-6), a pro-inflammatory cytokine, is a key regulator in the development of CAC (8). A previous study demonstrated that IL-6 is overexpressed in patients with active UC and CAC (9). However, despite evidence implicating the importance of IL-6 in the progression of CAC, the underlying molecular mechanisms of this association remain unclear.

Hypoxia is a common feature of solid tumors, including CAC, to which tumor cells frequently adapt through changes in their gene expression patterns (10). Hypoxia-inducible factor 1α (HIF-1α), an inducible transcription factor, is typically considered to be the primary regulator of the hypoxic adaptive response (10) and is regulated by the partial pressure of oxygen (11). In addition, a number of growth factors, including human epidermal growth factor receptor-2 and insulin-like growth factor-2, and cytokines, including tumor necrosis factor-α, have been demonstrated to regulate the expression of HIF-1α (12-14). Furthermore, previous studies identified that the activation of oncogenes and loss of function of tumor suppressor genes affects HIF-1α expression (15,16). HIF-1α induces the expression of hundreds of genes and serves important roles in various aspects of cancer biology, including angiogenesis, invasion and metastasis (17-19). In clinical studies, HIF-1α was shown to be overexpressed in multiple types of cancer, including colorectal (20,21). In addition, HIF-1α was identified to be overexpressed in patients with IBD and in the early stages of colorectal cancer (22). Furthermore, patients carrying one or both of the common HIF-1α polymorphisms, C1772T and G1790A, displayed a significantly higher risk for the development of CAC (23). These results indicate that HIF-1α serves a significant role in the progression of CAC. However, the association between IL-6 and HIF-1α during the progression of CAC remains unclear.

In the present study, the association between IL-6 and HIF-1α in the development of CAC was investigated. The results of the current study demonstrated that, in a CAC mouse model, the anti-IL-6 receptor antibody used inhibited the progression of CAC and decreased the expression of HIF-1α. Further analysis in human samples revealed that the serum concentration of IL-6 was positively correlated with HIF-1α.

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mRNA expression in CAC tumor tissues. In conclusion, the results of the present study indicate that IL-6 promotes CAC progression through the regulation of HIF-1α.

Materials and methods

Mouse CAC model. Male BALB/c mice (6-8 weeks old; 18.3±1.4 g), purchased from the Shanghai Laboratory Animal Center (Shanghai Institutes for Biological Sciences, Chinese Academy of Sciences, Shanghai, China), were used to produce the CAC model. Mice were housed at 22±1°C, with a 12 h light/dark cycle, and ad libitum access to food and water. Mice were divided into two groups; the immunoglobulin G (IgG) group (control group, n=8) and the anti-IL-6 receptor antibody (both eBioscience, Inc., San Diego, CA, USA) group (treatment group, n=8). The protocol to induce CAC in a mouse model was performed as previously described by Neufert et al (24). Briefly, mice were peritoneally injected with azoxymethane (AOM, 10 mg/kg; Sigma-Aldrich; Merck Millipore, Darmstadt, Germany) at the beginning of the first week. Then, mice were given drinking water supplemented with 2% (w/v) dextran sulfate sodium (DSS; Sigma-Aldrich; Merck Millipore) for 1 week, followed by water alone for 2 weeks, this was repeated 3 times (Fig. 1A). Simultaneously, mice in the treatment and control groups were injected peritoneally every 2 days with 10 µg anti-IL-6 receptor antibody or 10 µg IgG, respectively, diluted in 200 µl normal saline.

The mice were sacrificed via cervical dislocation at the end of week 9. Subsequently, large intestines (from the ileocecal junction to the anal verge) were collected, longitudinally cut open along the main axis and washed with normal saline. Visible neoplasms were counted and collected, the length and width of each neoplasm was measured using a digital micro ruler, and the neoplasm area was calculated by multiplying the length and width. Subsequently, neoplasms were cut into halves, where 1 half was prepared for histopathological analysis and immunohistochemistry, and the other half was prepared for reverse transcription-quantitative polymerase chain reaction (RT-qPCR) analysis and western blotting. All procedures were carried out in accordance with the guidelines and regulations for the Use of Experimental Animals by the Chinese Academy of Sciences (Beijing, China). The current study was approved by the Animal Ethics Committee of Zhongshan Hospital of Fudan University (Shanghai, China).

Collection of patient samples. A total of 13 patients (males n=8, females n=5), aged between 41 and 76 years old (58.6±10.9 years), diagnosed with CAC and receiving surgical treatment at Zhongshan Hospital of Fudan University between June 2012 and December 2014 were included in the present study. Tumor tissue and adjacent non-malignant tissue was collected during surgery and stored at -80°C until required for further analysis. A total of 5 ml blood was collected from each patient prior to surgery and immediately centrifuged at 2,400 x g for 10 min at room temperature. The serum obtained was preserved at -80°C until required. The patients included in the present study had not received chemotherapy or radiotherapy prior to surgery. Informed consent was obtained from all participants included in the current study and ethical permission for the study was obtained from the Zhongshan Hospital of Fudan University.

Histopathological analysis. Biopsy specimens were fixed with 10% formalin, embedded in paraffin and cut into 4-µm-thick sections. Hematoxylin and eosin staining was performed on the sections to distinguish between colorectal adenomas and adenocarcinomas. All results were confirmed by 2 pathologists.

Measurement of serum IL-6 concentration. The serum concentration of IL-6 was assessed using Canine IL-6 DuoSet ELISA (R&D Systems, Inc., Minneapolis, MN, USA; cat. no. DY1609), according to the manufacturer's protocol. The detection range of this kit was between 0.156 and 10.0 pg/ml, with a sensitivity of 0.11 pg/ml.

Immunohistochemistry. Immunohistochemical staining of colorectal tissue sections was performed as previously described by Yuan et al (25). Briefly, tissue sections were treated with xylene and a series of graded alcohols. Antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) at 95°C for 20 min. Following antigen retrieval, the sections were incubated with primary mouse monoclonal anti-HIF-1α antibody (1:100; Abcam, Cambridge, UK; cat. no. ab113642) at 4°C for 12 h. Following three washes with 0.01 mmol/l PBS, sections were incubated with goat anti-mouse IgG H&L (FITC) biotinylated secondary antibody (1:50; Abcam; cat. no. ab6785) at 37°C for 1 h and horseradish peroxidase-conjugated streptavidin (1:10,000; Abcam; cat. no. ab7403) for at room temperature 15 min. Then, sections were treated with diaminobenzidine and hydrogen peroxide chromogen substrate plus diaminobenzidine enhancer for 10 min, followed by counter staining with Mayer's hematoxylin. The percentage and intensity of stained immunoassayed cells was evaluated over 5 visual fields, selected at random, at a magnification of x400 using a light microscope. The degree of staining was evaluated independently by 2 pathologists. Expression of HIF-1α was indicated by the presence of cytoplasmic and/or membranous staining.

Total RNA extraction and RT-qPCR analysis. Total RNA was isolated from human and mouse colorectal tissue samples using the RNAprep Pure Tissue Kit (Tiangen Biotech Co., Ltd., Beijing, China; cat. no. DP431) and cDNA was synthesized from 1 µg total RNA using the cDNA Synthesis Kit (Takara Bio, Inc., Otsu, Japan; cat. no. DRR820A) on the StepOnePlus Real-Time PCR System (Applied Biosystems; Thermo Fisher Scientific, Inc., Waltham, MA, USA). Thermocycling conditions were as follows: 95°C for 2 min; 40 cycles of 30 sec at 95°C; 10 sec at 60°C; 30 sec at 72°C; and 10 min at 72°C. The expression level of the HIF-1α gene relative to β-actin was determined using the SYBR Green-based comparative CT method (2-ACT) (26). The primer sequences used were as follows: Human HIF-1α sense, 5’-ACT TCTGGATGCTGGTATGATT-T-3’ and antisense, 5’-GTCGTC CGAATAATACACTC-3’; mouse HIF-1α sense, 5’-AGCCCT AGATGGAAGGTTGGTA-3’ and antisense, 5’-TATCGAGGC TTGTCGACTG-3’; and human and mouse β-actin sense, 5’-CAGATGGAAGGCGCCGACCTCATC-3’ and antisense, 5’-TAAGACCTCTATGCCAACACAGT-3’.
Western blotting. Preparation of total protein lysates and western blot analysis was performed as previously described by Yuan et al (25). To detect HIF-1α, monoclonal anti-HIF-1α antibody (1:1,000) was used. Tubulin expression, detected using an anti-tubulin antibody (1:2,000; Sigma-Aldrich; Merck Millipore) was used as an endogenous control.

Statistical analysis. Results are presented as the mean ± standard error of the mean. Data from the present study was analyzed with GraphPad Prism (version 5.0; GraphPad Software, Inc., La Jolla, CA, USA) using a Student’s t-test. The correlation between serum IL-6 concentration and HIF-1α mRNA expression was analyzed using the Pearson product-moment correlation coefficient. P<0.05 was considered to indicate a statistically significant difference.

Results

Anti-IL-6 receptor antibody inhibits the development of CAC. CAC was successfully induced in the mice using AOM and DSS (Fig. 1A). Macroscopically, the majority of colorectal neoplasms were located in the middle and distal colon (Fig. 1B). Mice in the treatment group developed significantly fewer and smaller neoplasms compared with mice in the control group (P<0.05; Fig. 1C and D). Following histopathological examination, colorectal neoplasms were divided into adenomas and adenocarcinomas (Fig. 1E). The incidence of colorectal adenocarcinomas was 87.5% (7/8 mice) in the treatment group and 100% (8/8 mice) in the control group, while the incidence of colorectal adenomas was 100% (8/8 mice) in the treatment and control groups. These results suggest that the anti-IL-6 receptor antibody used inhibits the development of CAC.

Anti-IL-6 receptor antibody downregulates HIF-1α in colorectal adenocarcinomas. To identify the potential mechanisms by which the anti-IL-6 receptor antibody inhibits the development of CAC and to investigate whether this effect is through the regulation of HIF-1α, the expression of HIF-1α in colorectal adenocarcinomas was examined. RT-qPCR analysis demonstrated that HIF-1α mRNA expression was significantly downregulated in colorectal adenocarcinomas following treatment with the anti-IL-6 receptor antibody (P<0.05 vs. control; Fig. 2A). In addition, western blotting identified a corresponding downregulation of HIF-1α protein levels (Fig. 2B). Furthermore, immunohistochemistry revealed a notable decrease in HIF-1α protein levels in the treatment group compared with the control group (Fig. 2C).

Anti-IL-6 receptor antibody downregulates HIF-1α in colorectal adenomas. To investigate whether the anti-IL-6 receptor antibody used downregulates HIF-1α in early stage CAC, the expression of HIF-1α in colorectal adenomas was examined. Similarly to the results observed in colorectal adenocarcinomas, HIF-1α was found to be downregulated in colorectal adenomas following treatment with the anti-IL-6 receptor antibody (Fig. 3A-C). This indicates that IL-6 regulates HIF-1α expression in early stage CAC.

HIF-1α mRNA expression positively correlates with serum IL-6 concentration in patients with CAC. As HIF-1α expression was significantly downregulated by the anti-IL-6 receptor antibody used in the mouse CAC model, HIF-1α expression was analyzed in patients with CAC to determine whether it positively correlated with serum IL-6 concentration. RT-qPCR analysis revealed that HIF-1α mRNA levels were significantly increased in CAC tumor tissues compared with adjacent non-malignant tissues (P<0.05; Fig. 4A). The fold change in HIF-1α mRNA expression in tumor tissues compared with adjacent non-malignant tissues was between 0.8 and 5.2 (median, 2.1). In addition, western blotting confirmed
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that there was an overexpression of HIF-1α protein in the malignant tissues (Fig. 4B). Then, serum IL-6 concentration was examined in the participants, which identified that serum IL-6 concentrations were between 1.2 and 9.7 pg/ml (median, 6.4 pg/ml). Furthermore, the Pearson product-momentum correlation coefficient showed that serum IL-6 concentration positively correlated with HIF-1α mRNA expression in the tumor tissue of these patients with CAC (r=0.58; P=0.04; Fig. 4C).

Discussion

IL-6 overexpression is a well-known risk factor for CAC and the role of HIF-1α during CAC progression is well documented (22,27). However, whether IL-6 contributes to CAC progression through the regulation of HIF-1α remains unclear. The results of the present study indicate that inhibition of IL-6 using an anti-IL-6 receptor antibody hindered the development of CAC and decreased HIF-1α expression, even in early stage...
CAC. In addition, HIF-1α was demonstrated to be overexpressed in human CAC tumor tissue and positively correlated with serum IL-6 concentration. These results indicate that IL-6 promotes CAC progression by regulating HIF-1α expression during the early stages of CAC development.

Following the combination of exposure to AOM and 3 cycles of DSS treatment, CAC was successfully induced in BALB/c mice. Similarly to a previous study (28), the tumor incidence rate was 100% in the mice. In addition, following the inhibition of IL-6 using the anti-IL-6 receptor antibody, tumor size and number was significantly reduced, which was consistent with the results of a previous study (29).

In recent years IL-6 has emerged as a potential therapeutic target in patients diagnosed with malignant disease (30). For example, anti-IL-6 antibodies have been used to treat patients with multiple myeloma or acquired immune deficiency syndrome-associated Kaposi's sarcoma since the 1990s (31,32). In 2014 siltuximab, a chimeric anti-IL6 antibody, was approved by the U.S. Food and Drug Administration (FDA; Silver Spring, MD, USA) for the treatment of patients diagnosed with Castleman disease (33). In addition, siltuximab was used in clinical trials for a number of malignant tumors, including metastatic renal cell cancer, prostate cancer and colorectal cancer (34-36). Although tocilizumab, an anti-IL-6 receptor antibody, has not yet been approved for the treatment of malignant tumors, it has been approved by the FDA for the treatment of chronic inflammatory diseases, including rheumatoid arthritis and juvenile idiopathic arthritis (37).

In the present study, the anti-IL-6 receptor antibody used was observed to have a protective effect in a mouse CAC model. However, further investigations are required to validate the effectiveness of anti-IL-6 receptor antibody treatment of CAC.

Signal transducer and activator of transcription-3, suppressor of cytokine signaling-3 and vascular endothelial growth factor receptor-2 were reported to be the downstream proteins through which IL-6 exerts its tumor promoting effect in CAC (38-41). However, there is little literature on the association between IL-6 and HIF-1α in the development of CAC. In the present study, HIF-1α was downregulated in colorectal adenocarcinomas and adenomas in the mouse CAC model by the anti-IL-6 receptor antibody, which indicates that IL-6 stimulates the development of CAC through the regulation of HIF-1α. However, whether IL-6 exerts its tumor promoting effect directly through the regulation of HIF-1α remains unclear.

In patients with CAC, HIF-1α was overexpressed in tumor tissues compared with adjacent non-malignant tissues, which is consistent with the results of a previous study (18). In addition, a positive correlation between HIF-1α mRNA expression and serum IL-6 concentration in patients with CAC was observed. However, these findings should be interpreted with caution due to the relatively small number of patients included in the present study. Further research with a larger sample size is required to confirm these results.

In conclusion, the present study provides evidence that IL-6 promotes the development of CAC through the regulation of HIF-1α during the early stages of CAC progression. In addition, the results of the current study demonstrate that anti-IL-6 receptor antibodies are a potential therapeutic agent for CAC that warrant further investigation.

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References


