Expression of *Reg IV* in inflammatory bowel disease and the predictive effect on lesion activity.

Junwei Zhang*, Jing Bai, Limin Zhang, Aibin Cheng

North China University of Science Technology Affiliated Hospital, PR China

Abstract

Purpose: This article aims to observe the expression of the Regenerating gene (*Reg*) IV in intestinal tissues of patients of Inflammatory Bowel Disease (IBD) and explores the correlation between Reg IV expression level and the activity of IBD.

Method: 65 cases of IBD patients confirmed during January 2013-October 2013 as well as 30 cases of healthy people tested by colonoscopy inspection during corresponding period are taken for comparisons, including 30 cases of Ulcerative Colitis (UC) and 35 cases of Crohn's Disease (CD). The immunohistochemistry method is adopted to detect the protein expression level of Reg IV. The RT-PCR method is adopted to inspect the mRNA expression level of Reg IV. In addition, the correlation with IBD is further analysed.

Results: According to the quantitative results of immunohistochemistry staining, Reg IV has weak protein expression in health people, and Reg IV protein level of UC during active stage is significantly higher than that of remission stage (P<0.01); there is no significant difference on Reg IV protein expression level of CD during active stage and remission stage (P>0.05). The RT-PCR result is consistent with that of the immunohistochemistry.

Conclusion: Reg IV has a significantly higher expression in IBD, illustrating that Reg IV may be involved in the occurrence and development of IBD.

Keywords: Regenerating gene, Inflammatory bowel disease, Ulcerative colitis, Crohn's disease.

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Introduction

With common symptoms of diarrhea and stomach-ache, some patients of Inflammatory Bowel Disease (IBD) even have bloody stools. IBD includes Ulcerative Colitis (UC) and Crohn's Disease (CD) [1]. There has been no clear definition on the etiology and pathogenesis on IBD at present, which may be involved in environmental factors, genetic factors and microorganism infection, with changes on body immunity state [2-4]. The Regenerating gene (Reg) IV belongs to the calciumdependent phytohemagglutinin superfamily, and the protein coded belongs to the secreted protein, with expressions mainly in gastrointestinal tract and pancreas. According to existing researches, there is a close correlation between Reg IV and the occurrence and development of digestive system neoplasms [5]. However, little research on roles of Reg IV in the occurrence and development of IBD has been reported in literatures. Therefore, the immunohistochemistry method and the RT-PCR method are adopted in this research to detect the protein and mRNA expressions of Reg IV protein in intestinal tissues. In addition, analysis on the correlation between the expression level of Reg IV protein and the activity of diseases is conducted, so as to find out new indexes for assessment of IBD activity.

Materials and Methods

Clinical data

65 cases of IBD patients confirmed during January 2013-October 2013 were collected, including 30 cases of Ulcerative Colitis (UC) and 35 cases of Crohn's Disease (CD). The diagnosis of all the patients was in accordance with the diagnostic criteria for IBD of 2007 [6]. At the same time, 30 healthy people with colonoscopy inspection were collected for contrast.

Immunohistochemistry staining

The samples were baked at the temperature of 65-70°C for 45 min, followed by gradient dewaxing and dehydration as well as washing for 3-5 times with distilled water. Then citrate buffer solution was added into the samples which are then put into microwave oven at 95-98°C for 10-15 min for antigen retrieval. The samples were then taken out and cooled to indoor temperature, followed by washing. Then 0.6% H₂O₂ (H₂O₂ diluted with absolute methanol) was added. Ten minutes later, the samples were washed with distilled water for 2 times and then with PBS for 2 times.

The samples were then sealed (with 5% Bovine Serum Albumin) at indoor temperature (18-20°C) for 10 min. They were then placed for a night at 4°C after addition of primary antibodies. They were cleaned with PBS thoroughly for 3 times after taking out, and 5 min for each time. They were then added with second antibodies at 37°C for 25 min. They were cleaned with PBS for 3 times after taking out, and 5 min for each time. They were cleaned with tertiary antibodies (streptomycin) at 37°C for 30 min. They were cleaned for 3 times after taking out, and 5 min for each time. Then substrate was added for coloration.

RT-PCR detection method

The tissues were grinded to powders in liquid nitrogen. Then the Trizol method was adopted to extract the total RNA, and the RT-PCR method was utilized to measure the tissue expression quantity of the chemotactic factor. Please refer to the literature for the calculating method for the expression quantity of the chemotactic factor [7].

Quantitative method of the immunohistochemistry result

Please refer to the literature for specific methods [1].

Statistical method

The SPSS 16.0 statistical software was adopted for data analysis. T-test was adopted for comparisons between the data of 2 groups, and One-Way ANOVA test was adopted for comparisons among the data of multiple groups. P<0.05 meant the statistical significance of the difference.

Results

Reg IV protein expression in the detection with the immunohistochemistry method

As shown in Table 1, Reg IV protein expression levels in comparison: no expression (0 cases), low expression (23 cases), moderate expression (5 cases), and high expression (2 cases). Reg IV protein expression levels in UC patients: no expression (0 cases), low expression (5 cases), moderate expression (9 cases), and high expression (16 cases). Reg IV protein expression levels in CD patients: no expression (0 cases), low expression (7 cases), moderate expression (12 cases), and high expression (16 cases). According to the representative immunohistochemistry staining results in Figure 1, healthy people had only low Reg IV protein expression; Reg IV protein level of UC during active stage was significantly higher than that of remission stage (P<0.01), and there was no statistical significance on Reg IV protein level of UC during remission stage than that of healthy people (P>0.05). There was no significant difference between Reg IV protein expression level of CD during active stage and that of remission stage (P>0.05); however, Reg IV protein expression levels of both of them were significantly higher than that of healthy people (P<0.01). There was certain correlation between

 Table 1. Reg IV protein expression levels in comparison.

Group	Reg IV expression			
	-	+	++	+++
Healthy control group	0	23	5	2
UC group	0	5	9	16
CD group	0	7	12	16

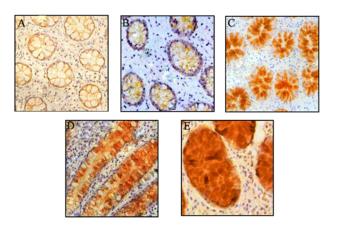


Figure 1. Detection on Reg IV protein expression level in intestinal tissues with the immunohistochemistry method (A. Healthy control; B. Remission stage of UC; C. Active stage of UC; D. Remission stage of CD; E. Active stage of CD).

Detection on Reg IV mRNA expression level with the PT-PCR method

As shown in Figure 2, according to the RT-PCR results, healthy people had relatively low Reg IV mRNA expression level and Reg IV mRNA level of UC during active stage was significantly higher than that of the emission stage. There was no significant difference on Reg IV mRNA level of CD during active stage and emission stage, but Reg IV mRNA level of both of them was significantly higher than that of the healthy people.

Correlation between Reg IV expression and the serum CRP concentration of UC patients

As shown in Table 2, the CRP concentration of the low expression group of Reg IV was $(5.83 \pm 4.67 \text{ mg/L})$; the CRP concentration of the moderate expression group of Reg IV was $(14.83 \pm 10.94 \text{ mg/L})$; the CRP concentration of the high expression group of Reg IV was $(81.74 \pm 39.82 \text{ mg/L})$. The concentration of CRP gradually increased with the strengthening of Reg IV protein expression level (P<0.01).

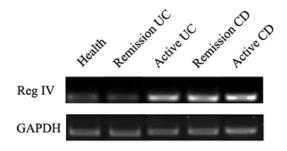


Figure 2. Detection on Reg IV mRNA expression level in intestinal tissues with the RT-PCR method.

 Table 2. Correlation between Reg IV protein expression level and serum CRP concentration of UC patients.

Number of cases	CRP concentration
5	5.83 ± 4.67
9	14.83 ± 10.94
16	81.74 ± 39.82
27.826	
0.001	
	5 9 16 27.826

Correlation between Reg IV expression level and Mayo index of UC patients

As shown in Figure 3, the Mayo index of the low expression group of Reg IV was 4.40 ± 1.14 ; the value of the moderate expression group was 6.89 ± 1.05 ; the value of the high expression group was 8.44 ± 1.59 . Mayo index gradually increased with the strengthening of Reg IV protein expression level (P<0.01).

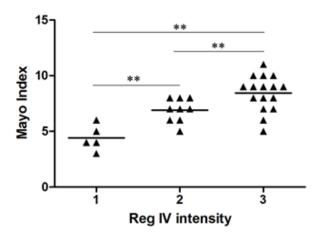


Figure 3. Correlation between Reg IV protein expression intensity and Mayo index of UC patients.

Discussion

With common symptoms of diarrhea and bellyache, some patients of Inflammatory Bowel Disease (IBD) even have

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bloody stools. Generally speaking, the diseases involved in ileum, rectum and colon are known as Inflammatory Bowel Disease (IBD). IBD includes Ulcerative Colitis (UC) and Crohn's Disease (CD) [1]. There has been no clear definition on the etiology and pathogenesis on IBD at present, which may be related to environmental factors, genetic factors and microorganism infection, with changes on body immunity state [2-4]. Clinically, the diagnosis on IBD is mainly made according to the clinical presentations of patients, the endoscopy results and the laboratory inspection results. In addition, in order to make a definite diagnosis on IBD, it is needed to exclude other diseases of immune system and intestinal tract. Therefore, the IBD auxiliary diagnosis and indexes related to the state of IBD are helpful to clinical diagnosis and judgment on disease state of IBD.

The Regenerating gene (*Reg*) IV belongs to the calciumdependent phytohemagglutinin superfamily, and the coded protein belongs to the secreted protein, mainly expressed in gastrointestinal tract and pancreas [8-10]. Little research has been published on the roles of *Reg IV* in the occurrence and development of IBD. According to the findings of this research, *Reg IV* has low expression in intestinal tissues of healthy people, which significantly increases in that of IBD patients; in addition, Reg IV protein level of UC patients during active stage is significantly higher than that of remission stage (P<0.01). The subsequent RT-PCR experiment further verifies that Reg IV protein has low expressions in healthy people, but has high ones in patients of IBD.

According to the above experiment, there is difference on Reg IV expression levels only between the UC patients during active stage and that during remission stage, and there is no significant difference on Reg IV protein expression levels between the CD patients during active stage and that during remission stage (P>0.05). It illustrates that Reg IV expression level may be related to the disease activity of UC patients. In order to work out the correlation between Reg IV expression level and UC disease activity, the correlation between Reg IV expression level and the serum CRP and Mayo index of UC patients is observed in this article. According to research findings, there is certain correlation between CRP level and UC disease state [11-15]. In addition, with the rise on Reg IV expression level, the serum CRP level of the patients gradually increases; besides, Mayo index, the disease activity index, also increases with the rise on Reg IV expression level. It illustrates that Reg IV expression can illustrate the activity of the disease of UC patients to certain extent.

In conclusion, according to the findings of this research, *Reg IV* has significantly high expressions in IBD patients. In addition, Reg IV protein level of UC patients during active stage is significantly higher than that of remission stage; there is no significant difference on Reg IV protein expression level between CD patients of active stage and that of remission stage. According to further analysis on the correlation between *Reg IV* expression level and the disease activity of UC patients, the serum CRP level and the Mayo index of the patients gradually increase with the rise on *Reg IV* expression level. It

illustrates that *Reg IV* may be involved in the occurrence and development of IBD, especially that of UC.

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*Correspondence to

Junwei Zhang

North China University of Science Technology Affiliated Hospital

PR China