Serum IP-10 Levels Correlate with the Severity of Liver Histopathology in Patients Infected with Genotype-1 HCV

Chan Ran You*, Su-Hyung Park†, Sung Won Jeong‡, Hyun Young Woo§, Si Hyun Bae*, Jong Young Choi*, Young Chul Sung†, and Seung Kew Yoon*

*Department of Internal Medicine, WHO Collaborating Center of Viral Hepatitis, The Catholic University of Korea College of Medicine, Seoul, †Postech-Catholic Biomedical Institute, Seoul, ‡Department of Internal Medicine, Institute for Digestive Research, Soonchunhyang University College of Medicine, Seoul, and §Department of Internal Medicine, Medical Research Institute, Pusan National University College of Medicine, Busan, Korea

Background/Aims: Interferon-γ-inducible protein 10 (IP-10) plays important roles in the pathogenesis of hepatitis C virus (HCV) infection. We investigated the association between serum IP-10 levels and liver pathology in patients with chronic HCV infection. Methods: The serum IP-10 concentration was assessed in 85 patients with chronic HCV infection using a solid phase sandwich enzyme-linked immunosorbent assay, and a liver biopsy specimen was obtained. The pathology was scored using the Knodell histologic activity index (HAI). Results: Of the 85 patients, 58 had genotype 1 HCV infection, 21 had genotype non-1, and 6 were undetermined. The serum IP-10 levels did not differ between patients infected with genotype 1 and genotype non-1 (p=0.472). In patients with genotype 1 infection, the total HAI score and the stage of fibrosis were highly correlated with the serum IP-10 level (r=0.555, r=0.578, p<0.001). Furthermore, the serum IP-10 concentrations of patients with severe fibrosis (stages 3, 4) were higher than those of patients with mild fibrosis (stages 0 to 2; 214.4 vs. 72.3 pg/mL, p=0.002) among patients with genotype 1 infection. However, in patients without genotype 1 infection, the histopathology was not associated with the serum IP-10 level. A multivariate analysis showed that serum IP-10 was an independent predictor of fibrosis (stages 3, 4) in patients with genotype 1 infection (odds ratio, 1.034; 95% confidence interval, 1.006 to 1.064; p=0.018). Conclusions: Serum IP-10 concentration was significantly correlated with the severity of liver histology in genotype 1 HCV infection. (Gut Liver 2011;5:506-512)

Key Words: Chronic hepatitis C; Fibrosis; Interferon-γ-inducible protein 10

INTRODUCTION

The immunological and pathological mechanisms of liver injury caused by the hepatitis C virus (HCV) remain unclear. Several reports have suggested that HCV-induced hepatic injuries result from the lysis of virus-infected cells by the host immune system, T cells and some cytokines, rather than direct destruction by HCV.1-3 The host immune response to HCV infection can lead to 2 opposite results: spontaneous eradication or chronic HCV infection. A powerful and specific cellular response may clear HCV, but most patients fail to eradicate HCV and become chronically infected with the virus. Persistent HCV infection results in chronic inflammation and fibrosis of the liver, cirrhosis, and hepatocellular carcinoma. Of the patients with chronic HCV infection, 20% progress to cirrhosis and approximately 4% progress to hepatocellular carcinoma.

Chemokines are relatively small cytokines that play various roles under physiological and pathological conditions. They are involved in the immune response, tumor development and progression, and tissue regeneration.4 Chemokines are divided into 4 families according to 2 characteristic N-terminal cysteine residues: CC, CXC, CX3C, and C.5 Chemokine receptors are expressed mainly in leukocytes, and the interaction of chemokines with their receptors induces leukocyte migration and activation.

Many chemokines and chemokine receptors are involved in immune reactions in response to HCV infection. Of these, the
non-EXR (Glu–Leu–Arg)-CXC chemokines and CC chemokines may be particularly important in the pathogenesis of HCV infection. 5,9 In addition, the mRNA expression of CCL2, CCL3, and CCL5, which are CC chemokines, and their receptors, CCR2 and CCR5, are elevated in HCV infection. 5,10 CXCR3 is expressed on activated T and natural killer (NK) cells, and CCR5 is expressed on activated memory T cells. 6 Interferon-γ-inducible protein-10 kDa (IP-10/CXCL10) is a non-EXR-CXC chemokine 12,13 and is a chemo-attractant for T lymphocytes, monocytes, and NK cells. 14,15 Recently, it was found that the expression of IP-10 in liver specimens from patients with chronic hepatitis C was higher than in patients with chronic hepatitis B and normal healthy controls. 16 Furthermore, the expression of IP-10 mRNA by hepatocytes strongly correlated with the serum IP-10 levels. 17 These studies suggest that IP-10 may be involved in the immune response to chronic hepatitis C. In this study, we evaluated the possible association between serum IP-10 levels and liver histology in terms of inflammation and fibrosis to examine the role of IP-10 in patients with chronic hepatitis C.

MATERIALS AND METHODS

1. Patients

Eighty-five patients with chronic hepatitis C were enrolled in this study. All of the patients were adults (median age, 52 years; range, 26 to 77 years), and consisted of 43 men and 42 women. The diagnosis of chronic hepatitis C was based on sero-positivity for anti-HCV antibody using third-generation enzyme immunoassay, the confirmation of HCV-RNA using reverse transcription-polymerase chain reaction (RT-PCR), and the histology of liver biopsy specimens. Patients with a history of alcohol abuse (more than 60 g/day) or taking herbal medicine and those diagnosed with hepatic failure (hepatic encephalopathy within 3 months after onset of jaundice), decompensated cirrhosis, hepatocellular carcinoma, other malignant tumors, or severe systemic diseases were excluded from the study. In addition, patients were excluded if they had taken immunosuppressants within the previous 6 months. The baseline characteristics of the patients are listed in Table 1.

### Table 1. Baseline Characteristics of the Patients

| Characteristic                        | Value                  |
|---------------------------------------|------------------------|
| No. of patients                       | 85                     |
| Mean age, yr*                         | 51.5±11.2 (26–77)      |
| Gender, M/F                           | 43/42                  |
| HCV genotype                          |                        |
| Genotype 1                            | 58                     |
| Genotype non-1                        | 21                     |
| Undetermined                          | 6                      |
| HCV-RNA titer, log copies/mL          | 6.3±0.7 (3.3–7.8)      |
| AST, IU/L                             | 79.2±54.0 (19–314)     |
| ALT, IU/L                             | 124.7±101.3 (12–492)   |
| Albumin, g/dL                         | 4.0±0.4                |
| Pathologic findings                   |                        |
| Knodell histologic activity index     | 5.9±3.2 (0–14)         |
| Mean grade                            | 4.1±2.3 (0–11)         |
| Mean stage                            | 1.8±1.2 (0–4)          |
| Median serum IP-10 level, pg/mL       | 103.8 (3.1–939.5)      |

Data are presented as mean±SD (range). HCV, hepatitis C virus; AST, aspartate transaminase; ALT, alanine transaminase; IP-10, interferon-γ-inducible protein 10.

Fig. 1. Serum interferon-γ-inducible protein 10 (IP-10) concentrations did not differ significantly between patients with genotype 1 (n=58) and those without genotype 1 (n=21).
4. IP-10 quantification

Serum samples for IP-10 quantification were obtained on the day of the liver biopsy and then stored at –70°C until analysis. The level of secreted IP-10 was measured using a DuoSet ELISA Development kit (R&D Systems, Minneapolis, MN, USA) according to the manufacturer’s protocol.

5. Statistical analyses

The Spearman rank test was used to analyze the correlations between serum IP-10 levels and histopathology, HCV-RNA titer, and laboratory results. Logistic regression was used in univariate and multivariate analyses to determine factors related to severe fibrosis. p-values of less than 0.05 were deemed to be significant. All statistical analyses were conducted using SPSS version 14.0 for Windows (SPSS Inc., Chicago, IL, USA).

RESULTS

The median serum IP-10 level for the 85 patients was 103.8 pg/mL (range, 3.1 to 939.5). The serum IP-10 level was not related to age (≤50 years vs. >50 years, 132.1±182.9 pg/mL). The serum interferon-γ-inducible protein 10 (IP-10) concentration positively correlated with histopathological severity in patients with genotype 1 hepatitis C virus infection.

Fig. 2. The serum interferon-γ-inducible protein 10 (IP-10) concentration positively correlated with histopathological severity in patients with genotype 1 hepatitis C virus infection.

HAI, histologic activity index.

Fig. 3. The correlations between serum interferon-γ-inducible protein 10 (IP-10) concentration and histopathological findings in patients with genotype 1 hepatitis C virus infection: piecemeal necrosis (A), intralobular degeneration (B), portal inflammation (C), and fibrosis (D). In patients with genotype 1 HCV infection, the serum IP-10 concentration was significantly correlated with the scores for piecemeal necrosis, portal inflammation, and fibrosis (A, C, and D).
vs. 116.4±76.8 pg/mL, p=0.587) or gender (male vs. female, 134.8±163.3 vs. 110.3±81.3 pg/mL, p=0.386).

1. Genotype of HCV and HCV RNA

Of the 85 patients, 58 had genotype 1 HCV infection, 21 had non-1 genotypes, and 6 were undetermined. Serum IP-10 levels were similar in HCV patients with genotype 1 and non-1 infections (130.2±148.5 vs. 105.6±76.7 pg/mL, p=0.472, Fig. 1). The mean HCV-RNA titer was 6.3±0.7 log copies/mL (range, 3.3 to 7.8 log copies/mL). The HCV-RNA titer did not correlate with serum IP-10 levels (r=0.063; p=0.579).

2. Association between histological severity and serum IP-10 level in genotype 1

For the 58 patients with HCV genotype 1, a positive correlation between serum IP-10 concentration and the Knodell HAI score was identified (r=0.555, p<0.001, Fig. 2). In terms of the relationship between the grade of necroinflammation and the serum IP-10 concentration in genotype 1 patients, the serum IP-10 concentration correlated with the degree of piecemeal necrosis (r=0.416, p=0.001) and portal inflammation (r=0.381, p=0.004), but not intralobular degeneration (r=0.060, p=0.660, Fig. 3). There was also a significant correlation between the serum IP-10 concentration and the stage of fibrosis (r=0.577, p<0.001, Fig. 3).

Of the patients infected with HCV genotype 1, the serum IP-10 concentration in the patients with severe fibrosis (stages 3, 4) was significantly higher than in the patients with mild fibrosis (stages 0 to 2, 214.4 vs 72.3 pg/mL, p=0.002, Fig. 4). The receiver operating characteristic (ROC) curve for severe fibrosis in genotype 1-infected patients had an area under the curve (AUC) of 0.846 (95% confidence interval [CI], 0.740 to 0.952, Fig. 5).

Fig. 4. The serum interferon-γ-inducible protein 10 (IP-10) concentrations were higher in patients with severe fibrosis (stages 3, 4) than in patients with mild fibrosis (stages 0 to 2, 214.4 vs 72.3 pg/mL, p=0.002).

Fig. 5. The receiver operating characteristic (ROC) curve of the serum interferon-γ-inducible protein 10 (IP-10) concentrations in patients with genotype 1 infection and severe fibrosis (stages 3, 4). The area under the curve (AUC) was 0.846 (95% confidence interval, 0.740 to 0.952), and the cutoff value of the serum IP-10 concentration was 100.3 pg/mL (sensitivity, 82.6%; specificity, 72.7%).

Table 2. Factors Related to Severe Fibrosis in Genotype 1 HCV Infection

| Parameter              | Univariate analysis | Multivariate analysis |
|------------------------|---------------------|-----------------------|
|                        | OR (95% CI)         | p-value               | OR (95% CI)         | p-value               |
| Age                    | 1.03 (0.99–1.08)    | 0.185                 | 1.16 (0.94–1.44)    | 0.170                 |
| Gender                 | 1.48 (0.51–4.32)    | 0.473                 |                       |                       |
| Viral load             | 0.33 (0.11–1.05)    | 0.060                 | 0.88 (0.10–8.03)     | 0.913                 |
| Serum IP-10 level      | 1.02 (1.01–1.03)    | <0.001                | 1.03 (1.00–1.06)     | 0.018                 |
| AST                    | 1.04 (1.02–1.06)    | <0.001                | 1.03 (0.96–1.11)     | 0.356                 |
| ALT                    | 1.01 (1.00–1.02)    | 0.009                 | 1.02 (0.98–1.07)     | 0.244                 |
| Serum albumin          | 0.03 (0.00–0.31)    | 0.004                 | 0.09 (0.00–8.46)     | 0.301                 |
| Total bilirubin        | 0.71 (0.30–1.70)    | 0.440                 |                       |                       |
| PT INR                 | 5.5×10^7 (248–1.2×10^9) | 0.002 | 2.8×10^10 (0.15–5.2×10^10) | 0.065 |
| Platelet count         | 0.99 (0.98–1.00)    | 0.053                 | 0.98 (0.94–1.02)     | 0.297                 |

HCV, hepatitis C virus; OR, odds ratio; CI, confidence interval; IP-10, interferon-γ-inducible protein 10; AST, aspartate transaminase; ALT, alanine transaminase; PT INR, prothrombin time international normalized ratio.
The cutoff serum IP-10 level for severe fibrosis in genotype 1-infected patients was 100.3 pg/mL (sensitivity 82.6%, specificity 72.7%). In a multivariate analysis of factors related to severe fibrosis, the serum IP-10 level was an independent predictor of fibrosis in genotype 1-infected patients (odds ratio, 1.034; 95% CI, 1.006 to 1.064; \( p = 0.018 \), Table 2).

In contrast to patients with genotype-1 HCV infection, no significant correlation was observed between serum IP-10 concentrations and Knodell HAI scores in the 21 patients with genotype non-1 HCV infection \((r = 0.034, p = 0.883, \text{Fig. 6})\). The serum IP-10 concentration did not correlate with the degree of piecemeal necrosis, intralobular degeneration, portal inflammation, or stage of fibrosis (Fig. 7).

### 3. Association between liver enzyme and peripheral blood mononuclear cell count and serum IP-10 level

In the patients with genotype-1 HCV infection, as the serum aspartate aminotransferase (AST) and alanine aminotransferase (ALT) levels became elevated, the serum IP-10 concentration also increased, and these correlations were statistically significant \((r = 0.463, p < 0.001; r = 0.484, p < 0.001, \text{respectively})\). The

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**Fig. 6.** The serum interferon-\( \gamma \)-inducible protein 10 (IP-10) concentration did not correlate with the severity of pathology in patients with genotype non-1 hepatitis C virus infection. HAI, histologic activity index.

**Fig. 7.** The correlations between the serum interferon-\( \gamma \)-inducible protein 10 (IP-10) concentration and pathology findings in patients with genotype non-1 hepatitis C virus infection: piecemeal necrosis (A), intralobular degeneration (B), portal inflammation (C), and fibrosis (D). In these patients, the serum IP-10 concentration did not correlate with the scores for piecemeal necrosis, intralobular degeneration, portal inflammation, or fibrosis.
serum IP-10 concentration of the patients with AST >100 IU/L was higher than that of the patients with AST ≤100 IU/L (median, 155.4 vs. 83.1 pg/mL, p=0.048). A similar result was obtained for the serum ALT (median, 140.8 vs. 80.5 pg/mL, p=0.014). A slightly positive (statistically nonsignificant) correlation was observed between the peripheral lymphocyte count and serum IP-10 concentration (r=0.250, p=0.058).

In patients with genotype non-1 HCV infection, serum IP-10 concentrations did not correlate with the peripheral lymphocyte count (r=0.117, p=0.615) or with serum AST (r=0.402, p=0.071) or ALT (r=0.233, p=0.308) levels.

**DISCUSSION**

IP-10 is a CXC chemokine. Unlike other CXC chemokines, IP-10 lacks chemotactic activity for neutrophils, but instead targets T lymphocytes, NK cells, and monocytes. IP-10 is produced by many cell types, including T-helper 1 (Th1) lymphocytes. In HCV infection, the inflammatory cells that infiltrate the liver are mainly antigen-nonspecific T-helper 1 (Th1) lymphocytes, which secrete cytokines such as interferon-gamma (IFN-γ) and IL-2, which activate monocytes and macrophages. In other words, Th1 lymphocytes initiate a delayed hypersensitivity reaction by producing IFN-γ and IL-2. This immune reaction results in ongoing tissue damage and progressive liver disease. IP-10 is produced by hepatocytes and recruits stimulated CD4+ T cells, NK cells, and monocytes to the target site, rather than CD8+ T cells and neutrophils. IP-10 may be involved in the Th1-dominant immune response of chronic hepatitis C.

In our study, the serum IP-10 level correlated with the histological severity of necroinflammation and fibrosis in patients with chronic genotype 1 HCV infection. Interestingly, the increasing of serum IP-10 level was independently related to severe fibrosis in genotype 1-infected patients. Thus, the IP-10 level is closely related to the progression of fibrosis in chronic HCV-mediated disease. A higher concentration of serum IP-10 may recruit more CD4+ T cells to the liver, inducing more vigorous immune reactions. This is supported by the correlation between the serum IP-10 level and the liver enzyme levels. Additionally, our results may indicate a potential association between activation of hepatic stellate cells and chemokines. The activation of hepatic stellate cells is known to be a key pathogenic feature of liver cirrhosis. Many cytokines, platelet-derived growth factors, and chemokines are involved in the activation of hepatic stellate cells. However, whether hepatic stellate cells express a specific receptor for IP-10 is unclear. It is a new challenge to discover the exact mechanism as to how IP-10 is involved in hepatic fibrosis.

Several other studies have demonstrated the relationships between the pathological findings and serum IP-10 levels. Diago et al. suggested that the serum IP-10 levels are higher in patients with advanced fibrosis than in those with mild fibrosis with chronic HCV infection, and that this relationship is more significant for genotype 1. Romero et al. demonstrated a correlation between IP-10 levels and necroinflammatory activity and fibrosis stage in patients with chronic hepatitis C. Recently, not only serum IP-10, but also intrahepatic IP-10, was reported to correlate with hepatic inflammation and fibrosis in chronic hepatitis C. Moreover, elevated intrahepatic mRNA expression of IP-10 in chronic HCV-infected patients was found to be associated with increased necroinflammation and fibrosis. This result suggests that serum IP-10 levels correlate with intrahepatic IP-10 level, as suggested by Mihm et al. The findings of the three studies mentioned above are consistent with our results. Furthermore, in our study, we evaluated the association between serum IP-10 and the severity of histology in patients with genotype non-1 HCV infection. The result was that serum IP-10 in patients with genotype non-1 HCV infection was not correlated with the severity of necroinflammation or fibrosis, in contrast to patients with genotype-1 HCV infection. The reason for this difference remains unclear, but may be attributable to the different mechanisms of pathogenesis involved under the 2 conditions. Generally, the prognosis of genotype-1 HCV infection is poorer than that of genotype non-1 HCV infection. In genotype-1 HCV infection, more powerful immune reactions may be induced, and IP-10 may be closely related to disease progression. Another issue is that we could not completely evaluate and make comparisons in terms of the association between serum IP-10 levels and the histological severity because of the limited number of patients with genotype non-1 HCV.

Patients with HCV infection show very diverse progressions in terms of hepatic injury. Some patients will only have mild inflammation, without fibrosis, despite the long duration of HCV infection, whereas others may progress to severe fibrosis or decompensated cirrhosis. According to our study, in patients with genotype-1 HCV infection, the serum IP-10 level could be a potential prognostic marker for fibrosis.

In conclusion, we showed that the serum IP-10 concentration was correlated with the severity of fibrosis and necroinflammation and that a high serum IP-10 level was an independent predictor of severe fibrosis in genotype 1 HCV infection. Further studies must determine whether IP-10 has the potential to be an important factor in the clinical management of patients with chronic hepatitis C such as in the development of new therapeutic strategies or the prediction of the prognosis.
CONFLICTS OF INTEREST

No potential conflict of interest relevant to this article was reported.

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