The pathological alternation of hepatic tissues in HBV infected patients with negative peripheral antigen and the expression profile of tissues HBV DNA.

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Abstract

Objective: During viral hepatitis B, serum HBsAg and HBeAg were important indexes reflecting viral replication and liver inflammation. However, patients with negative serum HBeAg still developed viremia or inflammatory necrosis. Negative HBsAg patients also had serum HBV DNA. This study thus analysed patients who were negative for HABsAg or HBeAg, to provide evidences for diagnosing chronic hepatitis with unknown reasons.

Patients and methods: A total of 22 patients were collected to detect HBV marker using HBV specific test kit. Biopsy was performed by ultrasonic guidance, followed by hematoxylin-eosin staining. Fluorescent quantitative PCR measured the copy number of HBV DNA from hepatic tissues and peripheral blood samples.

Results: 3 out of 22 cases (13.6%) showed positive HBV DNA in peripheral blood, while the positive rate of hepatic HBV DNA was 86.4%. In serum model positive for anti-HBs or anti-HBs+anti-HBc, pathological examination revealed no significant abnormality. In serum models of anti-HBs+anti-HBc +anti-HBe, anti-HBc or anti-HBc+anti-HBe, partial infiltration of inflammatory cells could be observed. The activity of hepatic tissues was the highest in hepatic tissues positive for anti-HBc, accompanied with higher fibrosis. A further correlation analysis revealed positive relationship between replication number of hepatic HBV DNA copy number and activity/fibrosis of hepatic tissues (r=0.56 or 0.34, p<0.05 in both cases).

Conclusion: In serum models with anti-HBc, anti-HBc plus anti-HBe, the activity and fibrosis degree of hepatic tissues were relatively higher, and were correlated with HBV DNA copy numbers.

Keywords: Antigen negative, HBV infection, HBV DNA.
**Inclusive criteria:** (1) Not received HBV vaccine; (2) No history of chronic hepatitis B; (3) Not received anti-viral treatment; (4) Negative for HBsAg and HBeAg but positive for serum core antibody for HBV (HBcAb) and/or HBeAb and/or HBsAb; (5) Serum enzymatic index for liver less or equalled to 2 ULN (80 U/L), and Total Bilirubin (TBIL) less or equal to 85.5 μmol/L.

**Exclusive criteria:** (1) Accompanied with other viral infection; (2) Alcohol abuse patients; (3) With auto-immune or metabolic liver disorders; (4) Abnormal liver development; (5) Complicated with major disorder (tumor, chronic kidney disease or diabetes); (6) Long term usage of liver-toxic drugs; (7) Using immune suppressant drugs; (8) Pregnant women.

The experimental protocol has been pre-approved by the ethical committee of Sichuan Mianyang 404 Hospital and written consents have been obtained from all patients and healthy volunteers.

**Reagents**

HBV marker test kit (Xinbo Bio, China); ZTLYB nucleic acid extraction kit (Tianlong Bio, China); Hematoxylin and eosin dyes (Beisuo Bio, China); Automatic biochemical analyzer (Beckman, US).

**Serum HBV marker assay**

The detection of HBV markers in peripheral blood was performed using HBV marker test kit following the manual instruction.

**Liver tissue biopsy**

Liver biopsy was performed under the guidance of colored ultrasound (indicator: serum TBIL<85.5 mmol/L, PTA>60%, excluding congestive liver lesion, hepatic echinococcosis, hepatic vascular disease, severe jaundice/ascites, infection on right pleura, worse general condition or severe anemia patients). Tissues collected were fixed in 4% formaldehyde, embedded in paraffin, and stained by hematoxylin-eosin. The morphology of tissues was examined by standards. The judgement of liver inflammation and fibrosis was deduced based on the standard stipulated by Chinese Medical Association (2000).

**HBV DNA extraction and assay**

Paraffin tissues sections were de-waxed, rinsed in ethanol to remove xylene, weighted and lysed. DNA was precipitated, filtered in column for binding. The column was rinsed to wash out, centrifuged and eluted to keep DNA at -20°C. HBV reaction mixture from the test kit was mixed for centrifugation. The reaction system consisted for 19.4 μl HBV reaction mixture and 0.6 μl Taq polymerase. Fluorescent quantitative PCR was performed under the following conditions: 95°C for 3 min, followed by 45 cycles each containing 94°C for 15 s and 60°C for 30 s.

**Statistical method**

SPSS16.0 software was used to analyse all collected data. Spearman was used for correlation analysis. The significant level was defined as 0.05.

**Results**

**Serum levels of HBV markers**

HBV surface markers were divided into 5 groups based on serum test results. As shown in Figure 1, in all HBV patients negative for HBsAg and HBeAg, there were 3 cases (13.6%) with positive peripheral HBV DNA, suggesting the existence of HBV replication even in some patients who were negative for HBeAg or HBsAg. In all 6 patients negative for HBsAg, no one showed HBV DNA existence (Table 1).

From HBV DNA test results on hepatic tissues from biopsy, there were a total of 19 cases which were positive for HBV DNA replication (86.4%). Therefore there were still lots of HBV infected patients even with negative expression for HBsAg and HBeAg, which should draw attention in clinical diagnosis and treatment (Table 2).

**Hepatic tissue pathology and HBVM serum modes**

In hepatic tissues samples collected from patients with different HBV serum markers, hematoxylin-eosin staining was performed (Figure 1). In anti-HBs (+) or anti-HBs (+) plus anti-HBc (+) patients, no significant abnormality was observed. In tissues with anti-HBs (+) plus anti-HBc (+) plus anti-HBe (+), or anti-HBc (+), or anti-HBc (+) plus anti-HBe (+), infiltration of inflammatory cells were observed.

**HBV serum modes and hepatic tissue fibrosis**

In different HBV serum modes, we analysed the activity of hepatic tissues and fibrosis. Results showed the highest activity in anti-HBc (+) patients, who also had highest fibrosis grade (Table 3). A further correlation analysis revealed the positive relationship between HBV DNA replication and activity/fibrosis of hepatic tissues (r=0.56 and 0.34, p<0.05 in both).

**Table 1. Peripheral HBV DNA expression in various serum modes of HBV patients.**

<table>
<thead>
<tr>
<th>HBVM serum mode</th>
<th>N</th>
<th>HBV DNA (+)</th>
<th>HBV DNA (+) rate</th>
</tr>
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<tbody>
<tr>
<td>Anti-HBs</td>
<td>6</td>
<td>0</td>
<td>0%</td>
</tr>
<tr>
<td>Anti-HBs+Anti-HBc</td>
<td>7</td>
<td>2</td>
<td>28.60%</td>
</tr>
<tr>
<td>Anti-HBs+Anti-HBc + Anti-HBe</td>
<td>2</td>
<td>0</td>
<td>0%</td>
</tr>
<tr>
<td>Anti-HBc</td>
<td>5</td>
<td>1</td>
<td>20.00%</td>
</tr>
<tr>
<td>Anti-HBc+Anti-HBe</td>
<td>2</td>
<td>0</td>
<td>0%</td>
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**Table 2. Analysis of hepatic tissue expression of HBV DNA across HBV infection modes.**

<table>
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<td>2</td>
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</table>
Most studies so far have confirmed the chronic hepatitis B with response, facilitating continuous replication of HBV in peripheral blood, suggesting the existence of HBV for HBsAg or HBeAg. Due to the limited sample size (N=22), caused by the mutation of anterior C domain and basic core start codon. Lots of studies showed that the mutant strain with negative serum HBsAg or HBeAg expression. In a total of 22 patients, there were 3 of them (13.6%) with positive HBV infection, as shown by re-activation of hepatitis and elevated HBV DNA replication quantification even in those patients who were negative for HBsAg but negative for HBsAg contents, which had higher lower limit in current assays. Most studies have found relatively lower HBV DNA in serum and lower viral copy number in hepatocytes in those patients who were positive for HBV DNA but negative for HBsAg [12-14]. Moreover, some studies found the negative correlation between HBsAg and age, indicating the possible suppression of HBsAg antibody level with aging [15].

Tissue pathological examination for patients with different HBV-surface markers found no significant abnormality in anti-HBs (+) or anti-HBs (+) plus anti-HBe (+) patients by hematoxylin-eosin staining, and partial inflammatory cell infiltration in tissues with anti-HBs (+) plus anti-HBe (+) or anti-HBe (+), or anti-HBe (+) plus anti-HBe (+). Severer inflammatory response existed in the existence of anti-HBe. Some scholars calculated the sensitivity of anti-HBe and found its sensitivity as 22% in all 2505 research objects, as four-fold of the sensitivity of HBsAg [16]. Therefore it is speculated that during HBV infection diagnosis, HBsAg may not be the sole sensitive marker due to individualized difference.

We further analysed hepatic tissue activity and fibrosis across different HBV serum modes. Results showed the highest activity in hepatic tissues with anti-HBc, which also had higher fibrosis level. Further correlation analysis revealed the positive relationship between HBV DNA replication number or hepatic tissue activity with fibrosis level, suggesting that anti-HBe might work as one index for HBV replication. Other study revealed no existence of such correlation between HBV DNA and hepatic tissue activity in HBsAg negative or positive patients [17]. Such inconsistency might be attributed to: (1) This study selected HBV patients with HBsAg and HBeAg; (2) This study selected Chinese population while the abovementioned study recruited Bangladesh people. Hepatic HBV DNA level is one important parameter measuring the treatment efficacy for HBsAg positive patients [5]. For HBsAg negative patients, however, no reliable evidence has been found so far. Other study also revealed the potency of HBV DNA in predicting hepatic fibrosis in HBV patients who were negative for HBeAg.

Another study revealed the existence of the correlation between HBV DNA and hepatic injury only in those patients with anterior C domain mutation [18]. Normally e-antigen may develop with the progression of chronic viral hepatitis. Therefore it is unreliable for differentiating between active hepatitis and non-active carriers solely based on the single

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<table>
<thead>
<tr>
<th>HBV serum modes</th>
<th>Hepatic tissue activity</th>
<th>Fibrosis</th>
</tr>
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<tbody>
<tr>
<td>Anti-HBs</td>
<td>1.2 ± 0.3</td>
<td>0.2 ± 0.1</td>
</tr>
<tr>
<td>Anti-HBs+Anti-HBc</td>
<td>1.3 ± 0.2</td>
<td>0.4 ± 0.2</td>
</tr>
<tr>
<td>Anti-HBs+Anti-HBc+Anti-HBe</td>
<td>1.8 ± 0.4</td>
<td>2.4 ± 0.6</td>
</tr>
<tr>
<td>Anti-HBc</td>
<td>2.6 ± 0.2</td>
<td>2.6 ± 0.2</td>
</tr>
<tr>
<td>Anti-HBc+Anti-HBe</td>
<td>1.5 ± 0.2</td>
<td>1.5 ± 0.3</td>
</tr>
</tbody>
</table>

Figure 1. Hepatic tissue pathology of various HBVM serum modes. A: anti-HBs (+); B: anti-HBs (+) plus anti-HBe (+); C: anti-HBs (+) plus anti-HBe (+); D: anti-HBc (+); E: anti-HBc (+) plus anti-HBe (+).

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value of HBV DNA [19,20]. The combination between clinical symptom, serum study and HBV DNA replication number should benefit effective diagnosis of viral hepatitis B.

Acknowledgments
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