T Cell Receptor (TCR) Gene Rearrangement During the Course of Chronic Hepatitis B. A Preliminary Report.

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Abstract

T cells recognize specific antigens through the T cell receptor (TCR) and self-MHC molecules. Structural variation in the TCR to recognise the enormous diversity of antigens is effected by maturation rearrangement of TCR genes in a manner analogous to immunoglobulin genes. Although it is believed that T cells play a major role in the seroconversion of eAg positive hepatitis B carriers, the mechanisms are still unclear. To examine the role of TCR rearrangements in chronic hepatitis B we have carried out Southern transfers of DNA aliquots from peripheral blood mononuclear cells (PBMC) using TCR chain specific probes. Rearrangements affecting as few as 5% of PBMC could be detected. Rearrangements were more common in eAg positive than negative. Of several rearrangement patterns a 5.5 kb EcoRI and 5.5 kb Hind III were most common. Longitudinal studies of 6 patients suggested the appearance of the rearrangement band preceded seroconversion and could be detected in PBMC for 6 months after seroconversion. It is postulated that during seroconversion the appearance of a clone of T cells with a particular rearrangement is associated with seroconversion and characterisation of these clones is currently being undertaken.

Keywords: T cell receptor, Gene rearrangement, Chronic hepatitis B

INTRODUCTION

The disease course of chronic hepatitis B is variable, with patients fluctuating between remission and active disease, as monitored by markers of liver destruction and virus replication. It is important to understand T cell activity during the course of chronic hepatitis B, as it may provide a means of monitoring the disease, and possibly will be of benefit for immunological intervention in care or treatment strategies.

From several studies, it appeared that different phases of disease progression can be recognized over the years during the long course of chronic hepatitis B. These phases reflect variations in host
immune response to HBV during different occasions of chronic infection.

Application of TCR gene rearrangement analysis during the course of chronic HBV infection would be useful in defining the disease progress, as it may reflect the immunological state in different phases of chronic disease. The observation of TCR gene rearrangement bands may thus be linked to the features most predictive of outcome in chronic hepatitis B. As discussed in several publications, these are serum aminotransferase level indicating liver destruction, and serologic markers of virus replication indicated by HBeAg present in the serum.

The possibility that there are changes of T cell clonality during the course of chronic HBV infection and the use of serum aminotransferase level and a virus replication marker as the indicators of disease chronicity, prompted the application of TCR gene rearrangement analysis in association with serum transaminase level and the presence of HBeAg in the serum of patients from whom blood had been taken on several occasions. This examination may provide a means for the evaluation of host immunity in virus-induced chronic hepatitis B, that may be advantageous for the strategy of chronic hepatitis B treatment.

SUBJECTS AND METHODS

Subjects

Subjects were 6 patients with chronic hepatitis B who had been tested serologically for HBeAg and anti-HBc as virus replication markers; the test was performed using commercial ELISA kit (Abbott laboratories, IL). Subjects were also tested for serum alanine transcrasase (ALT) level as a marker for liver destruction, using a standard method. The tests were carried out by the Pathology Department, Royal Brisbane Hospital, over a period ranging from 15 to 37 months. The blood was taken at least once in 2 or 3 months.

Methods

Genomic DNA was isolated from 10 ml of heparinized blood. 10 ug of genomic DNA was then digested completely with EcoR1 or Hind III in an appropriate buffer concentration overnight. DNA fragments were size fractionated by electrophoresis in 0.8% agarose and the separated DNA fragments were transferred to Hybond Nylon membrane (Amersham). The filter containing DNA fragments was then hybridized with the 32P-labelled TCR β cDNA probe overnight. At the end of the hybridization the filter was washed several times with 2 x down to 0.1 x SSC containing 0.1% SDS, and finally the filter was autoradiographed.

RESULTS

To determine whether the detection of TCR β gene rearrangements might be correlated with the pathogenesis of chronic hepatitis B, the appearance of rearrangement bands during the disease course of these six patients was associated with the disease progress which was assessed by regular testing of ALT level and serum HBeAg. The pattern of TCR β gene rearrangement, ALT level and serum HBeAg were plotted in one chart, and the results are shown in Figure 1.

As shown in case 1 (MY), the presence of rearrangement bands of 5.0 kb EcoR1 and / or 5.5 kb Hind III DNA fragments coincided with high ALT level and the presence of HBeAg in the serum, detected in the period of 5/88 to 2/89. As the patient underwent seroconversion and the ALT dropped to moderate level, the rearrangement bands become undetectable as shown in 9/89. Nine months later (6/90) the serum was again HBeAg positive, and the ALT level increased to a higher level, although not as high as in the period before seroconversion; rearrangement bands appeared again, but the size was different from those before seroconversion.

In case 2 (MF), the 5.0 kb EcoR1 and 5.5 kb Hind III rearrangement bands were also detectable and coincided with high ALT level and HBeAg positive serum as indicated in the period of 8/88 to 1/89. At the time of seroconversion between 6/89 and 11/89, the ALT dropped to normal level. At this time the presence of rearrangement bands was not assessed. Four months later (3/90) the ALT increased to a high level and reached a maximum in 5/90; the serum become HBeAg positive again, the 5.0 kb EcoR1 and 5.5 kb Hind III rearrangement bands also reappeared. The 5.0 kb EcoR1 band was undetectable in the beginning of ALT elevation in 3/90, probably because the amount of the T cell clone of interest at that time was less than the threshold of detection sensitivity. Four months later (9/90) the patient underwent seroconversion and the ALT dropped to normal level; this stage persisted until 9/91, during which rearrangement bands of TCR β gene were undetectable.

In case 3 (BW), there were no detectable EcoR1 rearrangement bands during 23 months of chronic disease course. the appearance of the 5.5 kb Hind III rearrangement band coincided with serum being HBeAg positive and with a high ALT level as in 3/89 and 9/89. As patient underwent seroconversion and ALT dropped to normal level in 1/90 the Hind III
rearrangement band became undetectable, and remained so through 2/91.

In case 4 (DD), the patient had not cleared the virus completely following 20 months of disease course. The 5.0 kb EcoRI and 5.5 kb Hind III rearrangement bands were detectable, coinciding with serum being HBeAg positive and with high ALT level as seen in the period between 10/88 and 3/88. These rearrangement bands disappeared as the ALT dropped to normal level in 8/89, when HBeAg was still present in trace amount in the serum (borderline). This stage persisted until 6/90.

In case 5 (DHD), the HBeAg was always negative during the course of chronic disease. ALT level increased slightly in the first 5 month period and reached a maximum in 6/89, during which 5.0 kb EcoRI and 5.5 kb Hind III rearrangement bands were detectable. When ALT dropped to normal level in 8/89 the rearrangement bands were undetectable. This situation proceeded until 3/90, when ALT level increased to moderate level and the 5.0 kb EcoR1 and 5.5 kb Hind III rearrangement bands reappeared. Six months later, the rearrangement bands disappeared again, coincident with the decreased ALT level.

Case 6 (KC) is unique in that the rearrangement of TCR β gene in peripheral blood T cells was detectable simultaneously with the rising ALT level and become undetectable as ALT dropped to normal level. However, the HBeAg was always negative during the course of the disease.

The results of monitoring TCR β gene rearrangement during the disease course on these 6 patients is summarized in table 1 and the general feature of the relationship of TCR β gene rearrangement, ALT level and serum HBeAg can be illustrated in Figure 2.

DISCUSSION

The appearance of TCR gene rearrangement bands in peripheral blood DNA, is associated with clonal expansion of a T cell clone to a proportion of at least 5% of total mononuclear cells, as has been confirmed previously. Clonal expansion of T cells would occur if there were a specific stimulation that induced specific T cells to become sensitized and undergo proliferation. Some previous experiments have reported that peripheral blood lymphocytes of chronic HBeAg positive patients are sensitized to HBeAg. Furthermore, some experiments suggest that the peripheral blood mononuclear cell compartments in patients with chronic active hepatitis B (CAH.B) contain cytotoxic T cells specific for HBeAg. HBeAg induces important immune reactions to cytotoxic class I restricted T lymphocytes responsible for the elimination of HBV-infected hepatocytes. In addition to cellular cytotoxicity, lymphokine secretion is another important consequence of T cell activation, which contribute to the outcome of viral disease. One of the important lymphokines is gamma-interferon, which has been well known for its antiviral effect, but may also contribute to cytotoxic events by upregulating the expression of HLA molecules on the susceptible target cells. This finding is likely to be relevant to the result of this study. The 5.0 kb EcoRI and 5.5 kb Hind III rearrangement bands may correlate with clonal expansion of cytotoxic T cells specific to HBeAg or regulatory T cells that mediate immune response by releasing lymphokines.

This study examined the hypothesis that TCR β gene rearrangement in peripheral blood T cells is a manifestation of immune progression in chronic hepatitis B. The experimental approach was based on the consideration that clonal proliferation of T cells in peripheral blood correlate with cell mediated immunity to virus infection, and linked to the activity of T cells in the site of inflammation in the liver.

It was found in this study that only few rearrangement bands of TCR β genes were observed in peripheral blood of chronic hepatitis B, indicating few clones of T cells to proliferate and detectable by direct examination of TCR β gene rearrangement of white blood cells. Dominant clones carrying TCR gene detectable as 5.0 kb EcoRI and 5.5 Hind III DNA fragments appear in most patients examined. This rearrangement bands change dynamically during the disease course concurrent with immune attack to infected liver in the process of viral clearance. A previous study by Moebius et al found that liver infiltrating lymphocytes distribute polyclonally as detected by TCR β gene rearrangement analysis on T cell clones from liver of chronic active hepatitis B (CAH) patients. It is likely that sensitized T cell clones expand in peripheral blood are part of those present in the site of inflammation in the liver. There are only small proportion of sensitized T cells in peripheral blood as most of them are sequestered into the liver for clearance of virus infected cells, and only cells with considerable expansion would be detectable by the present method.

The activity of T cells, as indicated by TCR β gene rearrangement detectable in peripheral blood was examined for association with liver destruction and virus clearance during disease progression. As shown in figure 1, presenting 6 cases, the TCR β gene rearrangement bands of 5.0 kb EcoRI and/or 5.5 kb Hind III DNA fragments were usually detectable several months before seroconversion, coinci-
Figure 1. Time course of the appearance of markers virus replication (HBeAg) and seroconversion (anti-HBe) in the serum, liver damage (indicated by serum ALT level) and TCR β gene rearrangements detectable when peripheral blood mononuclear cell DNA was digested with EcoRI or Hind III, in 6 patients with chronic hepatitis B. The presence of HBeAg or anti-HBe in the serum is indicated by positive (+) or negative (-) signs. Changes of ALT level are indicated by graph lines. Dates on the abscissa indicate date of blood collections. Detectable rearrangement bands are indicated by a positive (+) sign along with the size of rearrangement band (in brackets). A negative (-) sign indicates that rearrangement bands were not detectable (case 1, 2, 3, 4, 5 and 6).

CASE 1. Name : MY, Sex : Male, Age : 37 years

<table>
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<tr>
<th>HBeAg</th>
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<th>+</th>
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<tr>
<td>Anti HBe</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>+</td>
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ALT (U/l) vs. Date (1988-1991)

EcoRI

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<tr>
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Hind III

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<td>+</td>
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<td>(5.5)</td>
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(+18) (+18)
CASE 2. Name: MF, Sex: Male, Age: 45 years

CASE 3. Name: BW, Sex: Male, Age: 50 years
CASE 4. Name: DD, Sex: Male, Age: 51 years

CASE 5. Name: DHD, Sex: Male, Age: 19 years
CASE 6. Name: KC, Sex: Male, Age: 50 years

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<thead>
<tr>
<th></th>
<th>Case 1</th>
<th>Case 2</th>
<th>Case 3</th>
<th>Case 4</th>
<th>Case 5</th>
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<tbody>
<tr>
<td>HBeAg</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>+</td>
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<tr>
<td>Anti HBe</td>
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<td>+</td>
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Table 1. TCR β gene rearrangement analysis during the disease course of six patients, monitored in association with serum ALT level and the presence of HBeAg in the serum.

<table>
<thead>
<tr>
<th>ALT level reached</th>
<th>Average Period</th>
<th>e or e'</th>
<th>Case 1</th>
<th>Case 2</th>
<th>Case 3</th>
<th>Case 4</th>
<th>Case 5</th>
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</thead>
<tbody>
<tr>
<td>Prior to peak</td>
<td>5 mth</td>
<td>+</td>
<td>5/5.5</td>
<td>4.5/-</td>
<td>-5.5</td>
<td>5/5.5</td>
<td>-/</td>
<td>5/5.5</td>
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<tr>
<td>At peak</td>
<td>&lt;1 mth</td>
<td>+</td>
<td>5.5/5.5</td>
<td>5/5.5</td>
<td>-5.5</td>
<td>5/5.5</td>
<td>5/5.5</td>
<td>5/5.5</td>
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<tr>
<td>After peak</td>
<td>5 mth</td>
<td>+</td>
<td>-/-</td>
<td>5/-</td>
<td>-/-</td>
<td>-/-</td>
<td>-/-</td>
<td>-/-</td>
</tr>
<tr>
<td>Trough</td>
<td>≥7 mth</td>
<td>-</td>
<td>-/-</td>
<td>-/-</td>
<td>-/-</td>
<td>-/-</td>
<td>-/-</td>
<td>-/-</td>
</tr>
<tr>
<td>Prior to peak</td>
<td>5 mth</td>
<td>+</td>
<td>-/18</td>
<td>-5.5</td>
<td>-/</td>
<td>-/</td>
<td>-</td>
<td>5/5.5</td>
</tr>
<tr>
<td>At peak</td>
<td>&lt;1 mth</td>
<td>+</td>
<td>21/18</td>
<td>5/5.5</td>
<td>-/</td>
<td>-/</td>
<td>-</td>
<td>-/</td>
</tr>
<tr>
<td>After peak</td>
<td>5 mth</td>
<td>+</td>
<td>4.5/-</td>
<td>-/-</td>
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<td>Trough</td>
<td>&gt;7 mth</td>
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<td>-/-</td>
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Note: E = EcoR1 rearrangement band  
H = Hind III rearrangement band  
e = e antigen (HBeAg)
dent with the increase of ALT level. In some cases rearrangement bands of different size were detectable six months after seroconversion (as in case 1). The results indicate that in some chronic HBV carriers, the expansion of peripheral blood T cell clones with 5.0 kb EcoRI and 5.5 kb Hind III TCR β gene fragments is associated with cytotoxic clearance of virus infected hepatocytes and the loss of HBeAg from circulation. The clonal expansion of T cells detectable by TCR β gene rearrangement apparently represents the progressive development of immunity to virus infection in chronic HBV carriers. The expanded T cell clones are most likely linked to the activity of cytotoxic T cells with the potential to eliminate substantial numbers of virus infected liver cells. During this elimination phase, the destruction of liver tissue is marked by the elevation of ALT level in the serum. The elimination of virus infected hepatocytes by T cells is the effector mechanism of virus eradication, that consequently prevents viral replication and therefore reduced the release of virions to peripheral blood. This reduction of virus replication is characterized by the disappearance of HBeAg and in many patients by seroconversion from being HBeAg positive to being anti-HBe antibody positive. When there is no sign of viral replication and the patient becomes anti-HBe positive, liver cell lysis is decreased, indicated by normal or slightly elevated ALT level. The specific T cells responsible for liver cell lysis would be reduced in number, as antigenic stimulation leading to clonal proliferation is over. Consequently the TCR β gene rearrangement become undetectable, since the number of such T cells would be reduced to less than the threshold of detection sensitivity for TCR gene rearrangement.

Case 5 (DHD) and 6 (KC) are unique in that the rearrangement of the TCR β gene in peripheral blood was detectable simultaneously with rising ALT level, however the HBeAg was always negative during the course of the disease. As there was evidence of cytotoxic liver cell lysis (high ALT level), virus replication must have occurred previously, since production of viral antigen on the liver cells drives the cytotoxic reaction. The absence of HBeAg during the replication phase of the disease could be associated with the failure of the virus to produce and release e antigen to the circulation. This situation may occur due to mutation in the virus genes responsible for the

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**Figure 2. General pattern of the relationship of TCR β gene rearrangement in peripheral blood, ALT level and HBeAg in the serum during the course of chronic hepatitis B infection.**
secretion of HBeAg. HBeAg is part of the nucleocapsid precore-core gene product; when the full precore-core gene is transcribed, the transcript will be processed and the resulting mRNA is translated into a secretory protein. Transcription of the nucleocapsid genes starts at the second initiation codon, yielding HBc protein which is not secreted from cells due to lack of precore encoded peptide which functions as leader directing this protein to the cell membrane, ultimately to be secreted from liver cells.14 Mutation in the precore region would lead to failure of production of leader peptide necessary for secretion of HBeAg. Patients infected with such mutant virus would therefore never have detectable HBeAg in their serum, despite the presence of HBV-DNA and DNA polymerase simultaneously. This is presumably the case in patients 5 and 6.

This study describes the relationship of TCR B gene rearrangement with the mechanism of immune mediated liver injury and virus clearance in chronic hepatitis B. It might be suggested that specific TCR B gene rearrangement of 5.0 kb EcoR1 and / or 5.5 kb Hind III DNA fragments in peripheral blood are detected in chronic hepatitis B patients just prior to seroconversion. The 21 kb EcoR1 DNA fragment was found at the peak of ALT level, if the 5.0 kb EcoR1 or 5.5 kb Hind III DNA fragments were not detectable. This band may represent clonal expansion of T cells with the activity similar to those expressing the 5.0 kb EcoR1 or 5.5 kb Hind III DNA fragment. However, as the presence of this band is not common, its significance is unclear. As seroconversion is completed, the 5.0 kb EcoR1 and / or 5.5 kb Hind III DNA fragments usually become undetectable and this is accompanied by dropping of ALT to normal level. This general pattern of the relationship of TCR B gene rearrangement in peripheral blood with ALT level and serum HBeAg during the course of chronic hepatitis B infection was summarized as in table 1 and is shown diagrammatically in Figure 2. In general, the TCR B gene rearrangement band usually of 5.0 kb EcoR1 and / or 5.5 kb Hind III DNA fragments or sometimes 4.5 kb, 21 kb EcoR1 and 18 kb Hind III DNA fragments are detectable during the viral clearance phase of chronic disease course. This is indicated by the flare up of ALT level and continue to resolution phase where the ALT level decreasing to normal level. The rearrangement band disappears when ALT dropped to normal level and patient undergoes seroconversion, when the marker of virus replication is undetectable in the peripheral blood.

In conclusion, the appearance of specific TCR B gene rearrangement in peripheral blood T cells of chronic hepatitis B patients apparently represents the progression of host immunity to clear the virus. The TCR B gene rearrangement may reflect clonal expansion of T cells responsible for elimination of virus infected hepatocytes in the process of viral clearance, bringing the patient to seroconversion.

REFERENCES