Which is More Important and Insidious in Dialysis Patients? Occult Hepatitis B or Occult Hepatitis C?

Diyaliz Hastalarında Hangisi Daha Önemli ve Sinsidir? Okült Hepatit B veya Okült Hepatit C mi?

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ABSTRACT

Objectives: The aim of this study was to investigate the presence of occult hepatitis B infection (OBI) and occult hepatitis C infection (OCI) in hemodialysis patients and to determine whether there is an activation in the follow-up or not.

Materials and Methods: Demographic data, causes of renal failure, access to hemodialysis, duration of hemodialysis, alanine aminotransferase (ALT) levels, hepatitis indicators of 100 HD patients with normal ALT levels were recorded in this study. Serum anti-hepatitis B core antibody (anti-HBc) immunoglobulin G (IgG) was tested with ELISA (Architecht, Abbott). Serum hepatitis B virus (HBV)-DNA, HCV-RNA [in peripheral blood mononuclear cells (PBMCs)] were studied with “real-time” polymerase chain reaction method.

Results: Anti-HBc IgG positivity was detected in 27% of patients, but with no isolated anti-HBc IgG positivity. In 4% of the patients, HBV-DNA positivity and OBI infection were detected. None of the patients showed HCV-RNA positivity in serum and in PBMCs, therefore OCI was not detected. None of the patients developed OBI or OCI activation in five-years follow-up. Renal transplantation was performed in one of the OBI patients and lifelong prophylaxis was planned with oral antiviral medication.

Conclusion: Presence of OCI is lower than OBI in hemodialysis patients.

Keywords: Occult Hepatitis B, occult hepatitis C, hemodialysis
**Introduction**

In patients with chronic renal failure (CRF), infections are important causes of morbidity and mortality. These patients are particularly at risk of parenterally transmitted viral hepatitis (1). Hepatitis B and C viruses (HBV and HCV) are primarily transmitted parenterally in dialysis patients. Chronic hepatitis B (CHB) Chronic hepatitis C and (CHC) are more common infectious agents in patients with CRF compared to the normal population. These infections are also causes morbidity and mortality in patients with CRF and in patients undergoing renal transplantation (2,3). According to the Turkish Nephrology Association; hepatitis B surface antigen (HBsAg) positivity was 2.65% and anti-HCV positivity was 3.94% in hemodialysis patients in Turkey at 2017 (3).

HBV infection recovery defined as HbsAg dissapearance with HBV-DNA negativity in case of anti-Hbs positivity. The evaluation of serological markers for determining the infection is important but may be insufficient. Sensitive polymerase chain reaction (PCR) techniques have shown a low level of HBV-DNA in some patients who have spontaneously and serologically lost their HBsAg in serum and/or liver. Therefore, this condition, which defines chronic HBV infection (by PCR) with undetectable HBsAg levels, is called occult HBV infection (OBI) (4). OBI is divided into two groups according to anti-Hbc and anti-HBs positivity.

The actual cause of approximately 10% of liver enzyme abnormalities is unknown. In the last decade, OCI has been defined with studies have been conducted to identify patients with chronic liver disease whose etiology has not been clarified. Firstly HCV-RNA was detected in liver cells when anti HCV and HCV-RNA were negative in serum. Thereafter HCV-RNA was found in liver and in peripheral blood mononuclear cells (PBMNC) with undiagnosed high liver function tests. Viral RNA can be detected in PBMNC over 70% of patients with OCI (5,6).

OCI, firstly defined by Castillo et al. (6) HCV-RNA is detected in liver cells, while serum anti-HCV and HCV-RNA was negative. In the following years, Fabrizi and Martin (7,8) defined OCI, in patients with elevated liver enzyme; serum anti HCV and HCV-RNA were negative in serum. Thereafter HCV-RNA was found in liver and in peripheral blood mononuclear cells (PBMNC) with undiagnosed high liver function tests. Viral RNA can be detected in PBMNC over 70% of patients with OCI (5,6).

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The aim of this study was to investigate the presence of OBI and OCI in hemodialysis patients in Çanakkale, and follow up the reactvation of OBI and OCI.

**Materials and Methods**

This study was approved by the Çanakkale Onsekiz Mart University Ethics Commitee (approval number: 2014/03, date: 05.02.2014). We included 100 patients over 18 years of age and written informed consent was obtained from the patients. Patients were selected who had normal alanine aminotransferase (ALT) levels and shows seronegativity for HbsAg and anti-HCV antibody tests. The demographic data, ALT levels, hemodialysis periods, hepatitis B vaccination history, HBsAg, anti-Hbs and anti-HCV indicators were recorded.

Peripheral venous blood samples were collected from 5 mL each of 3 separated biochemistry tubes for anti-HBC Immunoglobulin G (IgG), HBV-DNA, HCV-RNA, and an amount of 9 mL blood in EDTA tube for PBMNC separation.

Anti-HBC IgG test was carried out with the Architect anti-HBC II Reagent kit. Blood samples for HBV-DNA and HCV-RNA isolation were centrifuged at 1500 rpm for 15 minutes. The obtained sera were stored at -20 °C until isolation of DNA and RNA.

Whole blood (9 mL) was taken into the EDTA tube for further differentiation of PBMNC. Histopaque® (R)-1077 (9 mL) was added to 50 mL falcon tube. Gently drop whole blood with sterile pasteur pipette from the edge of the falcon tube onto the Histopaque®-1077. According to the manufacturer’s recommendations, it was centrifuged at 400 G cycle for 30 minutes. After centrifugation, the cells in the cloud appearing in the middle of the tube were identified as PBMNC, and these cells were taken to the microviva lid cryo tubes by taking 3 mL with the help of micropipette. RNA was stored at -20 °C until isolation.

Prepared serum and PBMNCs after DNA/RNA isolation using HBV-DNA and HCV-RNA isolation kit (Magnesia®-2448 nucleic acid isolation and PCR setup robot) in Anatolia Diagnostic and Biotechnology R&D laboratory Montan®4896 real time (RT)-PCR Bosphore® HBV/HCV quantification (analytical sensitivity is 25 IU/mL and its linear range is 1x10 duy-1x 10 v IU/mL) was performed using Kit V1. The Bosphore® HBV Quantification Kit v1 (analytical sensitivity of 10 IU/mL and a linear range of 1x10 analytic-1x10 Kit IU/mL).

OBI was defined as HBV-DNA positivity in patients with HBsAg negative and with normal ALT levels.

OCI was defined as HCV-RNA positivity in patients with anti-HCV negative in patients with normal ALT levels.

**Statistical Analysis**

SPSS 20.0 package program used for data collection, recording and analysis.

**Results**

The study included 100 patients with normal ALT levels and HBsAg negative, anti-HCV negative in one dialysis center in Çanakkale province. Demographic data of the patients included in the study are given in Table 1. Fiftyeight (58%) of the patients were male and 42 (42%) were female. The mean age was 63.5±12.5 years. Eighty-five (85%) patients underwent dialysis through arteriovenous fistula. Other patients underwent dialysis with a...
permanent hemodialysis catheter. The mean duration of dialysis was 67.6 months. 95% of the patients had dialysis 3 days a week and 5% had 2 days dialysis. When the serological tests of the patients are examined; 27 patients (27%) were showed anti-HBc IgG and anti-HBs positivity together. None of the patients had anti-HBc IgG positivity alone. HBV-DNA was positive in 4/100, 4% of all patients. OBI diagnosed patients’ characteristics were shown at Table 2. When the hepatitis B vaccination history were examined, it was seen that 55% had at least three doses.

There was not any clinical and laboratory activation of hepatitis B in five-years follow-up of these four patients. Serum and PBCMN were investigated by PCR for OCI and HCV-RNA was not detected in any of the patients. The general characteristics of 27 patients with anti-HBc IgG positive are in Table 2. The mean age of the patients was 57±16.2 years. The mean duration of hemodialysis was 72.7±37.5 months. ALT levels were within normal limits and the mean was 9.7±2.3 IU/mL.

### Discussion

Although HBsAg positivity was decreased in hemodialysis patients, HBV viremia OBI was shown by PCR tests. The prevalence of OBI varies from 1% to 87% in different regions of the world (11).

The incidence of OBI is different in every country, for example it has been reported between 0%-36.4% in blood donors in our country (12). According to other studies, OBI was reported actullay between 3.4% and 19% in hemodialysis patients (13). In this study, we investigated the presence of OBI and OCI with RT-PCR in hemodialysis patients. In our study, the incidence of OBI was found 4%. But OCI was not obtained in our study group. In Egypt, Helaly et al. (14) found anti-HBc IgG positivity in all patients who had detected OBI. Therefore, in the presence of anti-HBc IgG positivity, patients should be investigated for possible OBI by molecular methods (14). But in our study none of the OBI patients did not showed this antibody positivity for core antigen. In our opinion, the main cause of this situtaion is; core antigen production was irregular in hepatocytes. If the replication continues regularly, you can detect antibody response, but if not antibody shows negativity. In hemodialysis patients this irregular sythesis might be in maximum level because of the CRF.

In the study conducted by Fontenele et al. (15) in Brazil, 79% of 301 patients with CRF who had hemodialysis showed anti-HBs positivity and isolated anti-HBs positivity was detected in only 35% of the patients. They found OBI in three patients with anti-HBs positivity alone. Anti-HBs positivity was found in 95% of our patients and 68% of patients have been showed anti-HBs positivity alone. All of OBI patients showed anti-HBs positivity alone.

Although the exact cause is unknown, the presence of anti-

### Study Limitations

Our study population was not enough to make a general recommendation for management of OBI in all dialysis patients.

### Conclusion

In our study, we found the incidence of OBI 4% in seronegative hemodialysis patients, but OCI was not obtained. When the infectious properties of these patients are also taken into consideration, it is inevitable that HBV negative patients will be infected by dialysis. It is an important risk factor that may adversely affect morbidity and mortality in these patients whose quality of life has decreased significantly due to CRF. Since our key diagnostic method for detection of OBI is HBV-DNA, it is essential to standardize the technique and used method. During the follow-up in dialysis units, once a year, viral DNA analysis with PCR-based method can be helpful in preventing the problems that may occur in the expected for organ transplantation.

### Acknowledgements

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### Ethics

Ethics Committee Approval: This study was approved by the Çanakkale Onsekiz Mart University Ethics Commitee (approval number: 2014/03, date: 05.02.2014).

Informed Consent: Written informed consent was obtained from the patients.

Peer-review: Externally and internally peer-reviewed.

### Table 2. Occult Hepatitis B diagnosed patients’ characteristics

<table>
<thead>
<tr>
<th>Patient Number</th>
<th>Age</th>
<th>Gender</th>
<th>HD way</th>
<th>Total HD time (month)</th>
<th>HBV DNA level (IU/mL)</th>
<th>Anti Hbs titer (mIU/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>(42)</td>
<td>33</td>
<td>M</td>
<td>AVF</td>
<td>59</td>
<td>61.4</td>
<td>1000</td>
</tr>
<tr>
<td>(58)</td>
<td>61</td>
<td>M</td>
<td>AVF</td>
<td>112</td>
<td>56.9</td>
<td>270</td>
</tr>
<tr>
<td>(98)</td>
<td>67</td>
<td>F</td>
<td>AVF</td>
<td>93</td>
<td>60</td>
<td>220</td>
</tr>
<tr>
<td>(100)</td>
<td>67</td>
<td>M</td>
<td>AVF</td>
<td>27</td>
<td>48.6</td>
<td>21</td>
</tr>
</tbody>
</table>

HD: Hemodialysis, AVF: Arteriovenous fistula, M: Male, F: Female
Authorship Contributions

Conflict of Interest: The authors declare no conflict of interest.

Financial Disclosure: The authors declare that this study has not received any financial support.

References