Roles of Toll-like Receptor 7 and 8 in Prevention of Intrauterine Transmission of Hepatitis B Virus

Ting Tian   Dandan Sun   Peng Wang   Hanzhi Wang   Xiaoxia Bai   Xiaofu Yang   Zhengping Wang   Minyue Dong

Women’s Hospital, School of Medicine, Zhejiang University, Key Laboratory of Women’s Reproductive Health of Zhejiang Province, Key Laboratory of Reproductive Genetics of the Ministry of Education, Hangzhou, China

Key Words
Hepatitis B virus • Intrauterine transmission • Placenta • Pregnancy • Toll-like receptor

Abstract
Background: Approximately 5% of newborns were infected by hepatitis B virus (HBV) via intrauterine transmission, but most of the infants born to HBV-positive mothers are protected from infection. However, the mechanisms by which intrauterine transmission is avoided remain elusive, and the roles of toll-like receptors (TLRs) have been proposed. The aims of this study were to clarify if TLR 7 and 8 are involved in the prevention of intrauterine transmission of HBV.

Methods: Real time polymerase-chain reaction (PCR) was used to determine the expression of TLRs and cytokines in placenta and trophoblasts. The expression of MyD88 was interfered with small interfering RNA (siRNA) in trophoblasts. An in intro model mimicking trophoblast barrier was established to evaluate the effect of MyD88 siRNA on HBV transmission across trophoblast barrier.

Results: There were significant differences in placental expression of TLR7 (F=3.263, P=0.048) and TLR8 (F=3.257, P=0.048) among control (HBV-negative women), non-infected group (HBV-positive women whose infants were not infected) and infected group (HBV-positive women whose infants were infected). The expression of TLR7 was significantly higher in non-infected group than infected group (P=0.039) and control (P=0.043). There was a significant difference in TLR8 expression between non-infected group and control (P=0.014), and the difference was close to but not significant (P=0.074) between non-infected and infected groups. Exposure of trophoblast to HBV significantly induced the expression of TLR7 (P<0.001), TLR8 (P=0.005), MyD88 (P=0.004), interferon (IFN)-α (P=0.004), IFN-β (P<0.001) and interleukin (IL)-8 (P=0.001). When MyD88 was interfered by siRNA, the expression of IFN-α (P<0.001), IFN-β (P=0.01) and IL-8 (P<0.001) was significantly decreased while the amount of HBV transcytosed across trophoblastic barrier significantly increased (P=0.03).

Conclusions: TLR7 and TLR8 on trophoblastic cells play an important role in the prevention of intrauterine HBV transmission by inhibiting HBV translocation across trophoblast.
Introduction

Hepatitis B virus (HBV) infection is a serious public health problem and chronic HBV infection affects over 350 million people in the world, which leads to a series of liver diseases including chronic hepatitis, fulminant hepatic failure, liver cirrhosis and hepatocellular carcinoma and over one million individuals die annually from these diseases [1]. Intrauterine infection is one of the most important reasons for the high prevalence of HBV infection in endemic regions. It is estimated that intrauterine infection of HBV occurs in 5-10% of infants of HBV-positive women [2]. Majority of infants of HBV-positive women are not infected via intrauterine route, indicating unexplained mechanisms protect the fetus from intrauterine HBV infection. The mechanisms by which HBV translocates across placental barrier and infects the fetus remain unknown, although it has been evidenced HBV infects trophoblast in vivo and in vitro and this infection is the first and most important step of HBV intrauterine infection [3, 4].

Placenta is considered the pregnancy-specific component of innate immune system and comprises the mechanical and immunological barrier restricting microbes from accessing the fetus [5]. Recently, it has been revealed placental trophoblast functions as a macrophage in recognizing and responding to pathogens through the expression of toll-like receptors (TLR) [6-8] and primary placental trophoblasts are more resistant to virus infection than non-trophoblastic cells [9]. The mechanisms by which placental trophoblast combats virus including HBV need elucidating; however, TLRs on trophoblast may play an important role.

TLRs are a family of innate immune receptors that recognize pathogen-associated molecular patterns (PAMPs) expressed by microbes and subsequently mediate an immune response [10, 11]. To date, 10 functional human TLRs have been characterized [10, 11], and their expression is not restricted to immune cells but is also found in a wide range of non-immune cells including placental trophoblast [12]. Following ligand recognition, TLRs activate a common intracellular signaling pathway by recruiting the intracellular signaling adapter protein MyD88. A subsequent kinase cascade triggers the activation of the nuclear factor κB (NF-κB) pathway, which results in the generation of an inflammatory response [13, 14].

In the current investigation, it was hypothesized that TLRs on trophoblastic cells is involved in the prevention of intrauterine transmission of HBV. To verify this hypothesis, mRNA expression of TLR7 and TLR8 was first determined in the placenta of HBV-positive and -negative pregnant women. Then, the response of trophoblast to HBV regarding mRNA expression of TLR7 and TLR8 and their down-stream anti-viral cytokines was observed and the effects of TLR-mediated inflammatory response in preventing HBV translocation across placental barrier were evaluated in an in vitro model.

Materials and Methods

Subjects

Seven HBV-positive women whose infants were infected via intrauterine transmission composed infected group (infected) and 30 randomly selected HBV-positive women whose infants were not infected comprised non-infected group (non-infected). In addition, 30 HBV-negative pregnant women served as control (control).

There were no significant differences in maternal age (P=0.675), gestational age at delivery (P=0.251), maternal body weight (P=0.933), neonatal birth weight (P=0.130) and neonatal gender (P=0.303) among three groups (Table 1). Maternal HBV-DNA load (P=0.306) and the prevalence of HBeAg (P=0.606) were not significantly different between infected and non-infected groups. HBV-DNA load ranged from 2,400 to 25,700 copies/ml infected neonates. No subjects received anti-HBV drugs.

Chronic maternal HBV infection was diagnosed on the basis of the seropositive hepatitis B surface antigen (HBsAg), the presence of HBV-DNA in maternal serum, normal liver tests and absence of symptoms of acute hepatitis during pregnancy. Intrauterine infection of HBV was defined as the seropositive HBsAg and the presence of HBV-DNA in serum of neonates taken on the third days after delivery. To exclude the possible
effects of labor on the intrauterine transmission of HBV, only mothers with selective Cesarean section were included. The indication for selective cesarean section was maternal HBV infection, breech presentation cephalopelvic disproportion or maternal request. All subjects were negative for serum markers of hepatitis A, C and E. Exclusion criteria included acute HBV infection, presence of pregnancy complications (such as preterm labor, preterm rupture of membrane and preeclampsia), fetal distress, auto-immune diseases, abnormal liver and kidney tests, human immunodeficiency virus (HIV), syphilis, infectious diseases and other diseases.

Standard passive and active immunoprophylaxis strategy was provided for all infants born to HBV-positive mothers. Vaccination strategy included the administration of 100 IU HBlg (human hepatitis B immunoglobulin) and HBV vaccine (5 μg) within 12 hours of birth. Vaccination series were later completed with two additional doses of HBV vaccine (one at 4 weeks of age and one at 6 month of age).

The protocol of the current investigation was approved by the Ethics Committee of Women's Hospital, School of Medicine, Zhejiang University and the informed consents were obtained from all the participants.

Sample collection

For the assay of HBV markers, maternal blood samples were taken at the admission to hospital, and neonatal blood samples on the third day after delivery. Blood samples were centrifuged after standing for at least 30 minutes in room temperature and serum was separated. For trophoblastic response test, blood samples were taken from high level HBV carriers (HBV-DNA > 1.0 × 10^8 copies/mL) with normal liver function test and healthy volunteers with a negative serum HBV marker and serum were separated aseptically. A 0.22 μm filtration device (Costar Co., Ltd.) was used for further sterilization. The complement was inactivated at 56°C for 30 min and serum was stored at -80°C until assay.

Placental samples were collected immediately after the placentas were delivered. Placental cotyledons were dissected at middle zone, washed thoroughly with ice-cold normal saline after amniotic membranes, decidua and connective tissues were removed. They were snap-frozen with liquid-nitrogen, and then stored at -80°C until assay.

Cell culture and HBV exposure test

Placental trophoblast cell lineage Swan 71 was cultured in DMEM/F12 medium supplemented with 10% fetal bovine serum and antibiotics (penicillin 100 units/ml and streptomycin 100 μg/ml) at 37°C/5% CO_2. Cells were seeded at a concentration of 2×10^5 cells per ml in tissue culture dishes and cultured overnight and then the medium was changed.

When the cells reached a confluence of 50%-60%, serum containing high quantity of HBV was added (corresponding to approximately 100 viral genomes equivalent/cell). After 48 hours of incubation, cells were then washed three times with PBS, trypsinized, and collected after extensively washed.

Interference of MyD88

A recombinant lentiviral vector expressing LV-MYD88-RNAi was constructed by Shanghai Genechem (Shanghai, China). LV-MYD88-RNAi was introduced into GV248 RNAi vector that carried the green

| N | Maternal age (y) | Gestational age (w) | Maternal body weight (kg) | Primiparity | Neonatal birth weight (g) | Neonatal gender (F/M) | Maternal HBsAg (+) | Maternal HBeAg (+) | Maternal DNA level (×10^8 copies) |
|---|-----------------|---------------------|--------------------------|--------------|--------------------------|----------------------|-------------------|-------------------|---------------------------------|
| N | Control 30      | Non-infected 30     | Intrauterine infected 7   |              |                          |                      |                   |                   |                                 |
|   | 29.1±3.08       | 28.4±2.88           | 28.8±4.85                | 0.675        | 39.0±1.13                | 38.7±0.79            | 38.4±1.27         | 0.251             |                                 |
|   | 70.0±6.39       | 70.6±8.65           | 69.5±8.56                | 0.933        | 29                       | 27                   | 5                 | 0.105             |                                 |
|   | 3510±397        | 3397±348            | 3164±430                 | 0.138        | 15/15                    | 19/11                | 3/4               | 0.303             |                                 |
|   | 0               | 30                   | 7                        | --           | 0                        | 18                   | 4                 | 0.606             |                                 |
|   | 62.5 (22.6, 170.3) | 67.8 (27.4, 159.0) |                            |              |                          |                      |                   |                                 |
fluorescent protein (GFP) reporter gene driven by the U6 promoter. Three vectors were designed: LV-MYD88-RNAi (8920-1), LV-MYD88-RNAi(8922-1), and LV-MYD88-RNAi (8923-1). The suppression of mRNA expression was analyzed by real-time reverse transcription-polymerase chain reaction (RT-PCR), and MyD88 protein levels were detected by Western blotting. The most efficient recombinant vector was used in later experiments. hU6-RNAi vector, which produces a non-targeting sequence TTC TCC GAA CGT GTC ACGT, was used as the negative control (GV248-NC). The recombinant vectors and the shRNA-LVs were co-transfected into 293 T-cell, and the titer of recombinant lentivirus was 5E+7 TU.

**HBV transcytosis study**

To observe the effect of MyD88 siRNA on HBV translocation across trophoblastic cell barrier, an in vitro transcytosis study system was established and verified as described by Bhat et al [3]. Briefly, Swan 71 cells were seeded to a confluence of 60-70% on six-well PET cell culture inserts (Corning Inc., USA) in a two-chamber system. HBV-positive serum (10^7 copies of HBV) was added to the apical domain of the polarized monolayers when the corrected transepithelial resistance measurements exceeded 100 Ωcm^2, and all of the apical and basolateral supernatant were collected respectively 48 hours later. The amount of HBV-DNA copies was determined with quantitative PCR.

**Assays**

Serum markers of hepatitis A, B, C and E including HAV IgM, HBsAg, HBeAg, anti-HBc, anti-HBe, HCV IgG and HEV IgM were routinely detected with enyme-linked immunoabsorbent assay (ELISA) (Sino-American Biotechnology Inc., Beijing, China) and HBV-DNA with real time PCR (Zhongshan University DaAn Gene Company, Zhongshan, China).

Total RNA was isolated by using the Trizol RNA reagent (Invitrogen Life Technologies, CA, USA). Reverse transcription was performed with random primers using PrimeScript RT reagent kit (Takara Bio Inc, Tokyo, Japan). The RNA and cDNA samples were stored at -80°C. The sequences of primers and PCR conditions for the detection of TLR7, TLR8 and the internal control genes were listed in Table 2. All primers were synthesized by Shanghai Sangong Biological Engineering Technology & Services Co. Ltd (Shanghai, China). Real-time PCR was performed on the ABI7900 HT Fast real-time PCR System (Applied Biosystems, USA) using the SYBR(R) Premix EX Taq (Takara Bio Inc, Tokyo, Japan) according to the manufacturer’s instructions. cDNA template of 2μl was used for each reaction in a system of 25μl. All PCR reactions were performed in triplicate. Thermal cycling conditions included pre-incubation at 95°C for 10 min followed by 40 cycles of 95°C for 15 s, 60°C for 30 sec and 72°C for 10 s. LightCycler collected data automatically and analyzed the value of Threshold Cycle (Ct). The fold changes of mRNA expression were detected by using 2^−ΔΔCt method.

**Statistical analysis**

Data distribution was tested with Kolmogorov-Smirnov test and data were presented in mean and SD or median and quartiles according to their distribution. Significance was tested with one-way ANOVA.
Results

As shown in Fig. 1, there were significant differences in placental mRNA expression of TLR7 ($F=3.263$, $P=0.048$) and TLR8 ($F=3.257$, $P=0.048$) among the three groups. The expression of TLR7 mRNA was significantly higher in non-infected group than infected group ($P=0.039$) and control ($P=0.043$). There was a significant difference in TLR8 expression between non-infected group and control ($P=0.014$), and the difference was close to but not significant ($P=0.074$) between non-infected and infected groups.

To determine if HBV up-regulates the expression of TLR7, TLR8 and downstream molecules, Swan 71 cells were cultured in the absence (control) or presence (HBV exposure) of HBV and detected the expression of TLR7, TLR8, MyD88, IL-8, IFN-α and IFN-β mRNA. We found significant increases in mRNA levels of TLR7 ($P<0.001$), TLR8 ($P=0.001$), MyD88 ($P=0.004$), IFN-α ($P=0.001$) and IFN-β ($P=0.001$) when trophoblast cells exposed to HBV.
Fig. 3. The effect of MyD88 siRNA. The expression of MyD88 was significantly decreased (P=0.04) when MyD88 was interfered by siRNA.

Fig. 4. The effect of MyD88 siRNA on the expression of IFN-α, IFN-β and IL-8 and HBV transcytosis. The silence of MyD88 significantly reduced the expression of IFN-α (P<0.001), IFN-β (P=0.01) and IL-8 (P<0.001) but significantly increased (P=0.03) the amount of HBV transcytosed across trophoblastic monolayer.

TLR8 (P=0.005), MyD88 (P=0.004), IFN-α (P=0.004), IFN-β (P<0.001) and IL-8 (P=0.001) when trophoblast cells exposed to HBV compared to control (Fig. 2). When MyD88 was silenced by siRNA, the expression of MyD88 protein was significantly reduced (P=0.04) (Fig. 3), the expression of IFN-α (P<0.001), IFN-β (P=0.01) and IL-8 (P<0.001) decreased significantly while the amount of HBV transcytosed across trophoblastic monolayer increased significantly (P=0.03) (Fig. 4).

**Discussion**

As a pattern recognition receptor, TLR recognizes and responds to HBV. It has been reported that TLR signaling inhibits HBV replication in vivo and in vitro. Isogawa et al [15] showed HBV replication was almost completely abolished in HBV transgenic mice by administration of a single intravenous injection of ligands specific for TLR3, TLR4, TLR5, TLR7, or TLR9 within 24 hours in an IFN-dependent manner. In vitro, supernatants of TLR3
or TLR4-stimulated murine Kupffer cells (KCs) and TLR4-stimulated sinusoidal endothelial cells (LSECs) suppressed HBV replication in HBV-Met cells [16]. In hepatoma cell HepG2, Xia et al [17] observed that TLR2, 3, 4, 7, 9 and MyD88 were abundantly expressed, and activation of these TLRs induced apparent production of antiviral cytokines including IFN-α and β and inhibited HBV replication. Further, the over-expression of MyD88 which is the universal adaptor for all TLRs with the except for TLR3 induced antiviral response and inhibited HBV replication in HepG2 and Huh7 cells [18]. In addition, the expressions of TLR7 [19], TLR9 [19, 20] and signaling molecules (IRAK1, IRAK4, TRAF3 and IRF7) [21] were significantly decreased in PBMC of chronic HBV infected subjects compared with control. These findings indicate that TLRs and its induced responses play important roles in the progress of HBV infection.

Placental trophoblast expresses functional TLRs. It has been reported that normal placental tissue expresses TLR1-10 at the mRNA level [8]. At the protein level, TLR2 and TLR4 are expressed in term placenta, however, there are contradictory findings in cellular localization of TLR4 [8]. Following ligation of TLR2 or TLR4 by zymosan or LPS, trophoblast cells from term placental explants can produce IL-6, IL-8, nitric oxide which has potent anti-microbial properties [22, 23]. Studies on first trimester trophoblast cells have shown that treatment with LPS induces the production of granulocyte colony stimulating factor (G-CSF) and regulated upon activation normal T cell expressed and secreted factor (RANTES) [24]. Messenger RNA of TLR1-10 is expressed on human chorioncarcinoma cell lines. Lipopolysaccharide (LPS) and DNA oligonucleotides containing unmethylated CpG motifs (CpG) induced the enhanced expression of TLR2 mRNA and TLR2 surface protein, activated signaling components (NF-κB, the phosphorylation of ERK1/2 and p38 MAP kinases) and increased the secretion of interleukin-8 [25]. Taken together, these findings suggest that trophoblast cells function similarly as immune cells and are able to recognize and respond to the pathogens at the maternal-fetal interface.

However, the study on the roles of placental TLRs in the prevention of intrauterine transmission of HBV is still little. In the current investigation, we described for the first time that placental expression of TLR7 and 8 was markedly increased in HBV-positive women (in vivo) and the in vitro exposure to HBV enhanced the expression of TLR7 and 8 as well as signaling molecule and antiviral cytokines of trophoblastic cell, indicating that TLR7 and 8 on trophoblasts function in a pattern similar to immune cells in recognizing and responding to HBV. We also revealed that the up-regulation in the expression of TLR7 and 8 was absent in the placenta of women whose infants were infected via intrauterine transmission, and silence of MyDD88 decreased the expression of antiviral cytokines and increased the amount of HBV translocating trophoblastic barrier, implying TLR and downstream molecules are involved in the prevention of intrauterine transmission of HBV. Given that placenta is the pregnancy-specific component of innate immune system [5] and TLRs and their downstream events are important part of innate immune system[3, 4, 7-9, 11, 13, 15, 22], our findings suggest that immune response to HBV by placental trophoblast is central in preventing intrauterine infection of HBV and up-regulation of TLR7 and TLR8 expression is part of the immune response to HBV infection.

Conclusions

In summary, toll-like receptor and mediated inflammatory response are necessary for the prevention of intrauterine HBV transmission and absence of TLR-mediated response to HBV is one of the mechanisms leading to HBV infection via intrauterine route.

Abbreviations

HBV (hepatitis B virus); TLR (toll-like receptor); PCR (poly-chain reaction); siRNA(small interfering RNA); IFN (interferon); IL (interleukin); PAMPs (pathogen-associated molecular patterns); NF-κB (nuclear factor κB); HBsAg (hepatitis B surface antigen); HIV (human immunodeficiency virus); HB Ig (human hepatitis B immunoglobling).
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Disclosure Statement

The authors declare that they have no competing interests

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