Beta2-glycoprotein I cooperate with hepatitis B surface antigen promotes hepatocellular carcinogenesis via the nuclear factor kappa B signal pathway were enhanced by the lipopolysaccharide

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Abstract

Aim: We aimed to elucidate whether beta2-glycoprotein I (β2GPI) cooperation with hepatitis B surface antigen (HBsAg) promoted hepatocellular carcinogenesis enhanced by the lipopolysaccharide (LPS) via activation of nuclear factor kappa B (NF-κB) and expression of tumor necrosis factor-α (TNF-α), interleukin-1β (IL-1β), and alpha fetal protein (AFP) in liver cancer cells.

Methods: Liver cancer cells (SMMC-7721) were transiently transfected with β2GPI and/or HBsAg and were subjected to LPS treatment. TNF-α, IL-1β, and AFP expression were measured in all groups by ELISA. NF-κB activation was assessed by non-radioactive electrophoretic mobility shift assay (EMSA) and was quantified in all groups.

Results: Cells transfected with β2GPI and/or HBsAg induced activation of NF-κB, with the highest activation seen in the doubly β2GPI- and HBsAg-transfected cells treated with LPS. Non-transfected cells treated with LPS exhibited lower activation compared to either β2GPI- or HBsAg-transfected cells with LPS treatment. In addition, cells transfected with β2GPI and/or HBsAg induced significantly increased expression of TNF-α, IL-1β and AFP, with the highest levels again seen in the doubly β2GPI- and HBsAg-transfected cells treated with LPS.

Conclusion: These observations suggest that the activity of NF-κB induced by β2GPI and HBsAg was enhanced by LPS. Expression of TNF-α, IL-1β and AFP increased in β2GPI and HBsAg cotransfected liver cancer cells.
INTRODUCTION

Beta2-glycoprotein I (β2GPI) also known as apolipoprotein H (apoH), is an abundant glycoprotein in the plasma\(^1\). To date, most studies of β2GPI have focused on its role in anti-phospholipid antibody-thrombosis syndrome\(^2,3\), lipid metabolism, coagulation, and/or regulation of the fibrinolysis system\(^4\). Mehdi et al.\(^5\) found hepatitis B surface antigen (HBsAg) bound β2GPI, an interaction that has been of interest to our research group. We previously found that there was a substantially increased level of β2GPI in hepatitis B-related hepatocellular carcinoma (HCC) tissue\(^6\). The combination of β2GPI and HBsAg substantially activated nuclear factor kappa B (NF-κB)\(^6\), suggesting that β2GPI played a role in the pathogenesis of hepatitis B-related HCC.

A recent study\(^7\) showed that lipopolysaccharide (LPS) specifically interacted with β2GPI, activating NF-κB via toll-like receptor 4 (TLR4) signaling pathway in macrophages. NF-κB is a pleiotropic transcription factor involved in inflammation-associated tumor promotion and progression in HCC\(^8\). Most hepatitis B-related liver cancer patients experience dysbacteriosis, resulting in increased levels of and sensitivity to LPS. In the present study, we further examined whether LPS enhanced the effect of β2GPI and HBsAg on activation of NF-κB, as well as the expression of cytokine factors in the liver cancer cells.

METHODS

Experimental groups

The human hepatoma cell line SMMC-7721 maintained in our laboratory were gifts from the central laboratory of the First Affiliated Hospital of Jilin University. The cells were incubated with Iscove’s modified Dulbecco’s medium (IMDM) culture media purchased from Gibco, containing 10% fetal bovine serum (FBS), and maintained at 37 °C in a 5% CO\(_2\) incubator. All cells were grown to adherence and were passaged every 2-3 days. Cells in the logarithmic growth phase were selected for experimental use. SMMC-7721 cells were divided into six experimental groups. Group A was the control group, neither transfected nor treated; group B was co-transfected with β2GPI- and HBsAg plasmids without LPS treatment; group C was treated with 500 μL (100 ng/mL) LPS and incubated for 6 h\(^9\); group D was transiently cotransfected with β2GPI- and HBsAg plasmids after treatment with 500 μL (100 ng/mL) LPS and incubated for 6 h; group E was transiently β2GPI-transfected after treatment with 500 μL (100 ng/mL) LPS and incubated for 6 h; group F was transiently HBsAg-transfected after treatment with 500 μL (100 ng/mL) LPS and incubated for 6 h.

Cell transfection

Groups B, D, E, and F were respectively transfected. The vector pcDNA3.1(-) was obtained from Invitrogen. The pcDNA3.1(-)-beta2-GPI and pcDNA3.1(-)-HBsAg eukaryotic expression plasmids were constructed previously in our laboratory. The recombinant plasmids, pcDNA3.1(-)-β2GPI, or pcDNA3.1(-)-HBsAg at 1 μg/well, and both at 3 μg/well (1:3) were dissolved in 50 μL IMDM basal media that was mixed to become Solution A. 2 μL FuGENE HD transfection reagent was dissolved in 50 μL IMDM basal media, mixed gently, incubated at room temperature for 5 min, labeled as Solution B. Solution A and Solution B were mixed gently to become Solution C, incubated at room temperature for 20 min. The cells were washed 3 times in serum-free IMDM culture media, and Solution C was slowly added to the cells that were incubated at 37 °C in a 5% CO\(_2\) incubator. Transfection media was removed after 6-8 h and was replaced with 500 μL 10% FBS IMDM media. Cell supernatants were collected at 24 h after transfection. A previous study\(^6\) from our lab found β2GPI protein expression was the highest 24 h after transfection.
Enzyme-linked immunosorbent assay analysis

Enzyme-linked immunosorbent assay (ELISA) detection of targets of interest was performed according to the manufacturers’ instructions. β2GPI was measured in groups A, B, D, and E; HBsAg in groups B, D and F; and tumor necrosis factor-α (TNF-α), interleukin-1β (IL-1β), and alpha fetal protein (AFP) in all groups. Once β2GPI reached the highest expression level, as determined from previous studies[^6^], cell supernatants from each group were collected for ELISA analysis. Triplicates of standards, samples and blank groups were prepared. The optical density (OD) value of each well was measured at 450 nm. Data were presented as means ± SD.

Non-radioactive NF-κB EMSA and NF-κB relative quantification

Assays were performed only with nuclear extracts according to the manufacturer’s instructions. Nuclear extracts (5 μg) were used for each reaction with 400 fmol bio-labeled (hot) oligonucleotide NF-κB probe (5’-AGT TGA GGG GAC TTT CCC AGGC-3’) and unlabeled (cold)-NF-κB probe (5’-AGT TGA GGG GAC TTT CCC AGGC-3’). Poly(dI-dC): poly(dI-dC) was used as a nonspecific competitor. A 25-fold molar excess of unlabeled homologous oligonucleotide was used as a specific competitor. Non-homologous oligonucleotide sequences were also used to validate the specificity of the binding of each transcription factor in the competition assays. Binding reaction resolved by 6.5% acrylamide/bis (30:1 ratio) electrophoresis in 0.25× TBE on ice. The gel was transferred to nitrocellulose membranes in 0.5× TBE. The membrane was then UV crosslinked for 10 min, blocked with 1× blocking buffer for 30 min, and then incubated with streptavidin-HRP in blocking buffer (1:750) at room temperature for 30 min. The membrane was washed four times with 1× washing solution and was equilibrated with 1× equilibration solution for 5 min with shaking. Finally, the membranes were incubated with chemiluminescence substrate buffer, and the bands were visualized using Viagene CoolImager (Viagene Biotech Co., China). NF-κB relative quantification was based on relative activity of the combination of NF-κB and DNA. The last result was represented by ΔΦ (gray value). The gray values of the image were measured after film exposure by the imaging system CoolImger. Data were presented as means ± SD.

Statistical analysis

SPSS 22.0 software was used for data processing and statistical analysis. Cell assay data were presented as means ± SDs and the variance was analyzed. Comparison between groups was measured using Fisher’s least significant difference (LSD) test. Differences were significant at P < 0.05.

RESULTS

Expression of β2GPI and HBsAg in transfected cells

We used ELISA to measure expression of β2GPI and HBsAg 24 h after transfection of recombinant plasmids in cell supernatants. β2GPI protein expression was found in group B, D, and E, significantly different from non-transfected, non-treated group A (P < 0.001). There were no differences in expression levels of β2GPI in groups B, D, and E (P > 0.05) suggesting similar transfection efficiency. HBsAg protein expression was found in groups B, D, and F. Expression was determined using a cutoff value (COV) that equal to the average absorbance value of the negative control (0.532). The absorbance of specimen ≥ COV indicated positive expression of HBsAg.

Activation of NF-κB in β2GPI- and/rHBsAg-transfected cells following LPS stimulation

A representative image of non-radioactive NF-κB EMSA in the six groups is shown in Figure 1, and NF-κB relative quantification was represented by gray value is shown in Figure 2. Groups B, C, D, E, and F induced differential levels of activation of NF-κB, with the highest relative activity of NF-κB observed in group D (1404.5 ± 11.28); this was significantly different compared with the other five groups (P < 0.05). The relative activity of NF-κB in group B was 914.57 ± 12.51, significantly higher than levels in groups A, C, E, and F (P < 0.05). The levels in group E (867.76 ± 6.27) and F (882.52 ± 7.92) were much higher than those of group...
Figure 1. Detection of non-radioactive NF-κB by EMSA in six groups. Group A: non-transfected, non-treated cells; group B: transient β2GPI- and HBsAg-transfection without LPS treatment; group C: non-transfected cells treated with 100 ng/mL LPS; group D: transient β2GPI- and HBsAg-transfection and treated with 100 ng/mL LPS; group E: transient β2GPI-transfection and treated with 100 ng/mL LPS; group F: transient HBsAg-transfection and treated with 100 ng/mL LPS. β2GPI: beta2-glycoprotein I; HBsAg: hepatitis B surface antigen; LPS: lipopolysaccharide

Figure 2. NF-κB relative quantification in six groups. Group A: non-transfected, non-treated cells; group B: transient β2GPI- and HBsAg-transfection without LPS treatment; group C: non-transfected cells treated with 100 ng/mL LPS; group D: transient β2GPI- and HBsAg-transfection and treated with 100 ng/mL LPS; group E: transient β2GPI-transfection and treated with 100 ng/mL LPS; group F: transient HBsAg-transfection and treated with 100 ng/mL LPS. Data presented as means ± SD: a: groups B, C, D, E, and F compared with group A, P < 0.05; b: group B compared with groups A, C, E, and F, P < 0.05; c: group D compared with other five groups, P < 0.05. β2GPI: beta2-glycoprotein I; HBsAg: hepatitis B surface antigen; LPS: lipopolysaccharide
C (590.4 ± 9.49) (P < 0.05). The level of NF-κB activation in group E and F were similar (P > 0.05). Taken together, these data suggest that LPS alone induced activation of NF-κB, which enhanced by either β2GPI- or HBsAg-transfection. However, the highest effect was seen in doubly-transfected cells, suggesting synergism between LPS, β2GPI and HBsAg with respect to activation of NF-κB in HCC.

DISCUSSION

HCC, one of the most common tumors, is currently the fifth most common malignant tumor worldwide, with morbidity increasing every year. Hepatitis B virus (HBV) and hepatitis C virus (HCV) are major causes of HCC\textsuperscript{[10]}. Therapeutic options include etiological treatment, resection, percutaneous ablation, trans-arterial chemoembolization (TACE), and targeted therapy. The overall efficacy of these therapies is poor, and five-year survival rates for early treatment of HCC are not favorable\textsuperscript{[11]}. Therefore, understanding the pathogenesis of HCC (abnormal neovascularization, genomics, proteomics and signal transduction pathways) is necessary to understand how HCC occurs and to develop new therapeutic approaches.
β2GPI is synthesized by liver cells and plays roles in anticoagulation, cell clearance, and lipid metabolism under normal physiological conditions. β2GPI is also involved in the pathogenesis of chronic viral hepatitis, alcoholic liver disease, autoimmune liver disease, liver cirrhosis and liver cancer. A previous study showed that a fraction with maximal apoH (β2GPI)-binding predominantly contained full Dane particles in HBV patients. Gao et al. found there was a specific binding event between HBV and β2GPI. Gao et al. provided the first evidence that a protein existed on SMMC-7721 cell membrane that could specifically bind β2GPI. The binding protein was later identified as annexin II. A previous study from our lab, demonstrated strong β2GPI expression in hepatitis B-related HCC tissue. In addition, the combination of β2GPI and HBsAg was shown to significantly activate NF-κB and expression of AFP, suggesting that β2GPI may be involved in the pathogenesis of hepatitis B-related HCC. However, it is unknown whether β2GPI directly interacts with HBsAg or if other proteins are involved in NF-κB activation.

β2GPI is physically closed in a circular conformation, with low activity. β2GPI opens and adopts a J-like conformation and becomes active when combined with antibodies or anionic phospholipids. In a study, it was found that LPS opened β2GPI, exposed its binding sites in domain V, and interacted with β2GPI to participate in physiology and pathology. The β2GPI and LPS complex relied on the TLR4 signaling pathway to activate NF-κB in macrophages. A previous study from our lab found that LPS enhanced signal transduction in β2GPI in liver cancer cells leading to activation of NF-κB, triggering downstream signal transduction and increasing the expression of downstream factors. This activation was related with LPS concentration. This suggests that LPS enhancement of β2GPI signal transduction may participate in the development of liver cancer.

LPS, a component of the cell wall of gram-negative bacteria, is an important mediator of the host inflammatory response to infection. A study of 169 patients with chronic hepatic disease found elevated levels of LPS in 27%, 85%, and 41% of patients with chronic hepatitis, chronic hepatitis with acute exacerbation and cirrhosis, respectively. In patients with chronic liver diseases, elevated levels of LPS in the portal and/or systemic circulation are common because of increases in intestinal permeability and bacterial translocation. LPS from gut microbiota contributed to HCC promotion by activating TLR4 signaling. Classically, TLR4 recognizes microbial lipids in homodimer configuration, thus activating various intracellular signaling pathways, such as the NF-kB and MAPK pathways. TLR4 has been identified in HCC and may play a role in progression of HCC. LPS-induced activation of TLR4 signaling promoted HCC cell survival and proliferation associated with regulation of the activation of the NF-kB and MAPK pathways.

In the present study, we demonstrated substantial activity of NF-kB in cells transfected with both β2GPI and HBsAg and treated with LPS. Our data suggested that the combined action of β2GPI and HBsAg were enhanced by LPS in the progression of carcinogenesis. Constitutive expression of NF-kB is emerging as a hallmark of cancer. In fact, constitutive NF-kB activation is generally associated with cancer proliferation, survival, chemoresistance, and progression of HCC.

NF-kB is another pro-inflammatory transcription factor that triggers downstream signal transduction and increases expression of downstream factors. In the present study, inflammatory cytokines (TNF-α, IL-1β, and AFP) were substantially elevated in cells transfected with both β2GPI and HBsAg and treated with LPS, more so than by single transfections with either factor. The action of various inflammatory mediators is known to occur in carcinogenesis. TNF-α has been postulated to have a crucial role in the pathogenesis of various cancers. It is one of the most important pro-inflammatory cytokines involved in the growth, differentiation, cellular function and survival of many cells. It is produced by several types of cells, including macrophages, neutrophils, fibroblasts, keratinocytes, NK cells, T and B cells, and tumor cells. IL-1β is also known to mediate several immune responses in HCV/HBV infection. There is a network of TNF-α and IL-1β secretion and interactive bio-functions in immune responses.
We found that LPS enhanced the effect of β2GPI- and HBsAg in development of liver cancer by increasing the activity NF-κB and elevating levels of TNF-α, IL-1β, and AFP. We predict that LPS may be an initiating agent in the pathogenesis of HCC, combining with β2GPI to activate and expose β2GPI binding sites to HBsAg, in turn interacting with HBsAg to further modulate NF-κB. Further studies are needed to uncover the specific mechanisms of interaction of β2GPI, HBsAg and LPS, and the role of β2GPI in liver cancer and other hepatic diseases.

DECLARATIONS

Authors’ contributions
Designed the study protocol, performed the studies and wrote the manuscript: Jing X, Ding XL
Gathered the data and performed the statistical analyses: Han NJ, Yang L, Yu YN
Improved the final version of the manuscript: Tian ZB, Gao PJ

Data source and availability
The data presented is original and obtained in our laboratory. It is available with the authors and can be made available if required.

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Conflicts of interest
There are no conflicts of interest.

Patient consent
Not applicable.

Ethics approval
Not applicable.

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