Silencing of Hypoxia-Inducible Factor-1β Induces Anti-Tumor Effects in Hepatoma Cell Lines under Tumor Hypoxia

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Abstract

Dimerization of hypoxia-inducible factor-1 beta (HIF-1β) [aryl hydrocarbon receptor nuclear translocator (ARNT)] with HIF-1α is involved in various aspects of cancer biology, including proliferation and survival under hypoxic conditions. We investigated the in vitro mechanism by which silencing of HIF-1β leads to the suppression of tumor cell growth and cellular functions. Various hepatocellular carcinoma (HCC) cell lines (Huh-7, Hep3B, and HepG2) were transfected with small interfering RNA (siRNA) against HIF-1β (siHIF-1β) and cultured under hypoxic conditions (1% O2 for 24 h). The expression levels of HIF-1α, HIF-1β, and growth factors were examined by immunoblotting. Tumor growth was measured using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide assay, and tumor activity was measured by terminal deoxynucleotidyl transferase dUTP nick end labeling, tumor cell invasion, and migration assays. Under hypoxic conditions, silencing of HIF-1β expression suppressed tumor cell growth and regulated the expression of tumor growth-related factors, such as vascular endothelial growth factor, epidermal growth factor, and hepatocyte growth factor. Suppression of tumor cell invasion and migration was also demonstrated in HIF-1β-silenced HCC cell lines. Silencing of HIF-1β expression may induce anti-tumor effects under hypoxic conditions in HCC cell lines.

Introduction

Tumor hypoxia is one of the distinguished features of tumor microenvironment found in hepatocellular carcinoma (HCC) [1,2]. Hypoxic microenvironment usually occurs in a rapidly proliferating tumor, mainly due to its increased metabolic rate and oxygen consumption [3,4]. Hypoxia plays an important role in tumor progression through angiogenesis and resistance to apoptosis [2]. Cellular response to hypoxia is mediated, at least in part, by a family of transcription factors known as hypoxia-inducible factors (HIFs) [2,3].

HIF-1 is a heterodimeric transcription factor, and is composed of two subunits, the oxygen-sensitive HIF-1α and constitutively expressed HIF-1β (also called the aryl hydrocarbon receptor nuclear translocator [ARNT]) [5,6]. Under low oxygen tension, the activated transcription factor, HIF-1α, upregulates diverse hypoxia-inducible genes through dimerization with HIF-1β, a co-activator of HIF-1α, and binds to the hypoxia-responsive element in the promoter of target genes [5,6].

HIF-1β plays an important role in the differentiation and development of many cells including T-cells, neurons, and hepatocytes under normoxia [5]. HIF-1β together with HIF-1α forms a complex with many other proteins and participates in various functions in the cell [7,8]. It is also a co-activator of several activators, and plays the role of receptor and transcription factor by connecting activators [6,9]. Similar to HIF-1α up-regulation, the expression of HIF-1β is increased by approximately two folds under hypoxic conditions [10]. The interaction of HIF-1α and HIF-1β is critical in the process of tumor survival. Although the influences of HIF-1α on tumor cells have been widely studied [3], the role of HIF-1β expression in tumor cell survival have been reported in few literature and therefore remains to be investigated. In this study, we demonstrated the effects of silencing HIF-1β expression on HCC cells, and found that HIF-1β-silencing regulates dimerization with HIF-1α under hypoxic conditions, leading to the suppression of tumor cell growth, invasion, and migration.
**Materials and Methods**

**Cell culture**

Huh-7 (KCLB60104, Korean Cell line Bank), Hep3B [11], and HepG2 (KCLB88065, Korean Cell line Bank) cells were cultured at 37°C with 5% CO₂ in Dulbecco’s Modified Eagle Medium (DMEM; Gibco, Grand Island, NY) supplemented with 10% fetal bovine serum (FBS; Welgene, Daegu, Korea), 4.5 g/L glucose, L-glutamine, and 1% penicillin/streptomycin.

**Small interfering RNA (siRNA) and transfection**

siRNA was synthesized using the following sequences: siHIF-1α: (Forward) 5'-GUG GUU GGA UCU AAC ACU A-3', (Reverse) 5'-UAG UGU UAG AUC CAA CCA C-3'; siHIF-1β: (Forward) 5'-CAG ACA AGC UAA CCA UCU U-3', (Reverse) 5'-AAG AUG AGC UUG UGU U-3'. Cells were transfected with respective siRNAs using Fugene HD transfection agent (Promega, Madison, WI, USA) according to the manufacturer’s instructions. After transfection, the cells were maintained at normoxic conditions (21% O₂) for 24 h, then replaced with fresh culture medium, followed by 24 h of culture under hypoxic conditions (1% O₂, 5% CO₂, 94% N₂).

**Cell growth assay**

Tumor cell growth rate was measured by the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT; Amresco, Solon, OH, USA) assay according to the manufacturer’s instructions. Briefly, the cells were seeded in a 96-well plate at 5×10⁴ cells/well, incubated at 37°C for 24 h, transferred to serum-free medium, and transfected with siRNA. After incubation at 21% O₂ for 24 h, the culture medium was replace with fresh medium supplemented with 10% FBS. Hypoxia was induced by incubation under 1% O₂ at 37°C for 24 h. After addition of MTT to each well, the plates were incubated at 37°C for 3~4 h for sufficient staining of cells, followed by medium removal and dimethylsulfoxide (Sigma-Aldrich, St. Louis, MO, USA) treatment. Cell viability was measured with absorbance at 595 nm using a spectrophotometer (Molecular Devices, Toronto, Canada).

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**Figure 1. Suppression of tumor cell growth by silencing of hypoxia-inducible factors-1α and -1β**. Huh-7 cells were transfected with small interfering RNAs against HIF-1α, HIF-1β, or green fluorescent protein as control (siHIF-1α, siHIF-1β, and siGFP, respectively), followed by exposure to hypoxic conditions (1% O₂). (A) Tumor cell growth after silencing of HIF-1α or -1β was measured by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay. Tumor cells were susceptible to growth retardation under hypoxic conditions when more than 100 nM of siHIF-1β was transfected. However, normoxic conditions (100 nM) did not show significant difference. (B) Transfection of siHIF-1α (100 nM) or siHIF-1β (100 nM) suppressed cell growth when maintained at hypoxic conditions. As compared to control, the growth inhibition was more prominent with increase of siHIF-1β concentration. NT, non-target; siGFP, siRNA against green fluorescent protein; siHIF1-α, siRNA against HIF-1α; siHIF-1β, siRNA against HIF-1β. *, P<0.05; **, P<0.01.

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Real-time quantitative polymerase chain reaction (PCR)

Total RNA was extracted from HCC cells using Trizol reagent (GIBCO BRL, Grand Island, NY, USA) according to the manufacturer’s instructions. cDNA was synthesized from 1 μg of RNA using reverse transcriptase (Clontech, Mountain View, CA, USA), according to the manufacturer’s instructions. Gene expression was measured by real-time quantitative PCR.

Immunoblot assay

The effects of silencing HIF-1α and HIF-1β expression on the expression of proteins related to cell proliferation and angiogenesis were assessed by immunoblot assays. Total proteins were extracted from HIF-1α- or HIF-1β-silenced HCC cells after 24 h of hypoxia induction. The proteins were separated according to their molecular weight via sodium dodecyl sulfate–polyacrylamide gel electrophoresis (SDS-PAGE), transferred to a polyvinylidene fluoride membrane (GE Healthcare/Amersham, Buckinghamshire, UK), and blotted with mouse monoclonal antibodies specific for the proteins of interest. The blots were developed using the enhanced chemiluminescence technique (PerkinElmer, Boston, MA, USA) according to the manufacturer’s instructions, and the level of expression of each protein was quantified and compared.

Immunoprecipitation

Total cell lysate was extracted from HCC cells using RIPA cell lysis buffer (Cell signaling, Danver, MA, USA) according to the manufacturer’s instructions. HIF-1α was bounded with mouse anti-HIF-1α (Cell signaling, Danver, MA, USA) using IgG magnetic beads (Novex, Oslo, Norway), according to the manufacturer’s instructions. The dimerized proteins were detected according to western blot, and blotted with the other target-mouse monoclonal antibodies-HIF-1α (Bethyl, Montgomery, TX, USA) specific for the proteins of interest. The blots were developed using the enhanced chemiluminescence technique (PerkinElmer, Boston, MA, USA) according to the manufacturer’s instructions, and the level of expression of each protein was quantified and compared.

Tumor cell invasion assay

The effects of HIF-1α and HIF-1β knockdown on tumor cell invasiveness were investigated in Huh-7 cells grown in serum-free DMEM. The invasiveness of tumor cells was assessed in vitro using a transwell chamber (Corning Costar, Cambridge, MA, USA). Each transwell chamber was plated with 1 × 10^5 cells, and the invading cells were stained with hematoxylin and eosin. The total number of invaded cells on the lower side of the filter was counted under the microscope (Olympus America, Melville, NY, USA).

Migration assay

The mobility of cells was assessed by scratch and wound healing assay. Each experimental result was observed by optical microscope.

Cell death assay

Cells were stained with FITC-labeled annexin V and propidium iodide (PI), and tumor cell death was assessed by terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL, Promega) assay and flow cytometry.

Statistical analysis

Results were expressed as means ± standard error of the mean (SEM) or frequency (%). Independent t-test was performed to compare the difference of the mean between control and experimental groups. All statistical analysis was done using SPSS version 12.0 (SPSS, Inc., Chicago, IL). A p value of less than 0.05 was considered statistically significant.
Results
Silencing of HIF-1α and HIF-1β suppresses tumor cell growth

After transfection of HCC cells with various concentrations of siHIF-1β, tumor cell growth was assessed by MTT assay. Forty-eight hours after transfection, tumor cell growth was significantly suppressed as compared to control, in a dose-dependent manner. The negative effect of HIF-1β silencing on tumor cell growth was more prominent under hypoxic conditions, especially when more than 100 nM of siHIF-1β or siHIF-1α was transfected to the tumor cells (Fig. 1A, and File S1). Although the tumor cell growth was maintained at higher doses of siHIF-1β transfection under normoxic conditions, exposure to hypoxic environment resulted in significant suppression of tumor cell growth. Suppression of tumor cell growth under hypoxic conditions by HIF-1β-silencing was more pronounced by prolonged exposure to hypoxic environment. (Fig. 1B). Of note, consistent with the previous findings [12], silencing of HIF-1α also displayed suppression of tumor cell growth under hypoxic conditions, which is probably mediated by inhibition of several other targets related to cell proliferation.

These findings demonstrate that tumor cell growth is suppressed by silencing of HIF-1β under hypoxic conditions.

Silencing of HIF-1β affects expression of tumor growth-related genes

Since tumor cell growth was inhibited by HIF-1β-silencing, we measured the mRNA levels of growth factors involved in tumor cell growth in HIF-1β-silenced tumor cells. Expression of HIF-1β was reduced by >60% in the group treated with siHIF-1β compared with that in the Control group, in a cell-density-dependent manner. Therefore, siHIF-1β inhibits expression of HIF-1β. HIF-1α expression was induced by hypoxia regardless of HIF-1β knockdown. To determine the influence of HIF-1β knockdown on genes related to tumor growth, the mRNA expression levels of epidermal growth factor (EGF), fibroblast growth factor (FGF), and hepatocyte growth factor (HGF) were quantified by real-time quantitative PCR (Fig. 2). Under hypoxic conditions, silencing of HIF-1β produced diminished expression of EGF and HGF by 52% and 36% (p-value < 0.05), respectively, compared to control. However, mRNA levels of FGF2 expression...
was not affected by HIF-1β-silencing. Collectively, these data suggest that under hypoxic conditions, HIF-1β expression regulates the expression of tumor growth-related factors, namely EGF and HGF, but not FGF2.

Silencing of HIF-1β affects protein expression and secretion of tumor growth-related genes

Based on the aforementioned mRNA data, protein expression of tumor growth-related genes were analyzed using Hep3B cells. Dose-dependent and selective inhibition of HIF-1β expression by siHIF-1β transfection was confirmed by immunoblot assays (Fig. 3A). Interaction of HIF-1β and HIF-1α was confirmed by immunoprecipitation (File S2). IP results demonstrated that decreased protein expression levels of HIF-1β or HIF-1α in HIF-1α tagged group. To determine the influence of HIF-1β inhibition on factors related to tumor cell growth, the levels of protein expression and secretion of EGF, HGF, VEGF, and FGF2 were analyzed (Fig. 3B). ELISA results demonstrated that decreased protein expression levels of EGF, HGF, and VEGF in HIF-1β-silenced cells. Of note, the expression of these molecules was also decreased in HIF-1α-silenced cells. Notably, the expression of FGF2 was not affected by silencing of HIF-1α or HIF-1β. Collectively, these results demonstrate that under hypoxic conditions, HIF-1β expression regulates the expression of various tumor growth-related factors, namely EGF, HGF, and VEGF, but not FGF2.

HIF-1β-silencing suppresses tumor cell invasiveness and motility

Since tumor cells exhibit potential to mobilize and invade into adjacent and distant regions, the effect of HIF-1β-silencing on invasiveness and migration ability of tumor cells was studied using various HCC cell lines. The effect of HIF-1β-silencing on the invasiveness of tumor cells was evaluated by assessing the number of cells that have mobilized and moved across the matrigel-coated trans-well to the gelatin coated-bottom well. Compared to control, siHIF-1β-silenced cells showed remarkably reduced number of cells in the bottom well, denoting suppression of tumor cell invasiveness (Fig. 4A). Likewise, as compared to control, markedly diminished migration of HIF-1β-silenced Huh-7 cells was confirmed by scratch and wound healing assay (Fig. 4B). Of note, silencing of HIF-1α also induced suppression of tumor cell invasiveness and migration. Together, these findings suggest that silencing of HIF-1β affects the invasiveness and migration of tumor cells under hypoxic conditions.

Silencing of HIF-1β sensitizes tumor cells to hypoxic apoptosis

Normally, cells are predisposed to apoptosis upon exposure to prolonged hypoxic environment, a phenomenon known as hypoxic apoptosis. However, tumor cells tend to circumvent apoptosis through various mechanisms, with the aid of the HIF system. To evaluate whether HIF-1β is responsible for the resistance to hypoxic apoptosis, cell death assay was performed in HCC cells silenced for HIF-1α or HIF-1β followed by exposure to hypoxic environment. Cell death assay demonstrated that hypoxic apoptosis was merely induced in the control (Figure 5). On the contrary, hypoxic apoptosis was markedly increased by silencing HIF-1β, a finding similar to the effect of HIF-1α-silencing on tumor cell survival. Collectively, these data suggest that HIF-1β, along with HIF-1α, regulates hypoxic apoptosis of tumor cells under hypoxic conditions.

Discussion

In the tumors, hypoxia is an important mechanism that induces proliferation, metastasis, and neovascularization of tumors.
HIF-1α is a key molecule in hypoxia [12], and is known to be involved in the proliferation of tumors and survival mechanisms such as angiogenesis and anti-apoptosis [15]. HIF-1α is highly regulated by oxygen concentration [12]. Under hypoxia, HIF-1α maintained and concentrated in the cytoplasm [10]. In normoxia, the HIF-1α proteins are rapidly degraded, resulting in essentially no detectable HIF-1α protein [15]. On the other hand, HIF-1β is not affected by oxygen concentration [18]. Although HIF-1β can function by forming a dimer with HIF-1α, few studies of HIF-1β have been performed [15–18]. HIF-1β was initially known as an ARNT factor, and plays an important role in the differentiation and development of many cell types such as T cells, neurons, and hepatocytes, by dimerization with AhR, which is similar to HIF-1α [18–20]. Moreover, the AhR pathway, which involves HIF-1β (ARNT), is known to mediate anti-tumor effects [21]. Among high-risk patients with HCC in one study, HIF-1β expression was high and was associated with cell cycle arrest [22]. We investigated the function of HIF-1β under hypoxic conditions. The hypoxia response, which plays a role in tumor development, presumably prioritizes the function of HIF-1β rather than that of ARNT. Accordingly, further study of HIF-1β is warranted.

This study proceeded under the assumption that even when dimerization of HIF-1α and HIF-1β is inhibited, compared with inhibition of HIF-1α expression, the same result as expression of HIF-1α could be obtained. As a result, siRNA inhibition of HIF-1α expression was performed, and the present study performed the MTT assay to determine cell growth when a hypoxic stimulus was applied. Effect of inhibition for tumor growth similar to case of inhibition for HIF-1α could be identified.

![Figure 5. Silencing of HIF-1β sensitizes tumor cells to hypoxic apoptosis.](#)

Cell death assay demonstrated that hypoxic apoptosis was markedly increased by silencing HIF-1β in tumor cells, a finding similar to the effect of HIF-1α-silencing on tumor cell survival. NT, non-target; siGFP, siRNA against green fluorescent protein; siHIF1-α, siRNA against HIF-1α; siHIF-1β, siRNA against HIF-1β; PI, propidium iodide.

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![Graphs showing cell death assay results.](#)
related to tumor growth through real-time quantitative PCR, Western blotting, and ELISA was decreased [4,14]. EGF is a growth factor that stimulates cell growth, proliferation, and differentiation by binding to its receptor EGFR. HGF regulates cell growth, cell motility, and morphogenesis by activating a tyrosine kinase signaling cascade after binding to the oncofetal c-Met receptor. VEGF is a signaling protein produced by cells that stimulates neo-vascularogenesis and angiogenesis [23]. The above three factors are all related to tumor growth and development, and control of the expression of these three growth factors plays an important role in anti-tumor effects. Based on these findings, the expression of HIF-1α is inhibited and this subsequently inhibits the formation of a dimer with HIF-1β. Therefore, expression control of a sub-gen is another method to inhibit the activation of tumors under hypoxic conditions.

In the present study, the influence of hypoxia on cells and that of HIF-1α knockdown were evaluated using wound healing assay and tumor cell invasion. When HIF-1α expression was inhibited, wound healing was suppressed; similarly, when HIF-1α was knocked down, motility decreased. Moreover, among various liver cancer cell lines, only the Huh7 cell line can be used to assess tumor invasion by measuring the degree of tumor invasiveness that was performed using Huh7 in vitro. Further investigation is needed to determine the cause of such a difference. The current study evaluated the invasion degree of Huh7 cells when expression of HIF-1β is inhibited, and the result was compared with the group for which the expression of HIF-1α was inhibited. When the expression of HIF-1α was inhibited, invasion did not occur as expected. We guessed that inhibition of HIF-1α expression prevented the formation of a dimer with HIF-1γ; in turn, various functions that should be performed under hypoxic conditions, such as inhibited cell survival, cell motility, cell growth, and impeded invasion.

Regarding tumor proliferation, a hypoxic state is important for tumor growth to start. It is thought that HIF-1 expression (HIF-1α and HIF-1β) controls the initiation of tumor growth, and can be important in affecting anti-tumor growth by changing growth to be more malignant in a hypoxic state. Further study is required to determine other possible functions of HIF-1β that are comparatively less known than those of HIF-1α, which has drawn most of the attention until now.

Supporting Information

File S1 Suppression of tumor cell growth by knockdown of hypoxia-inducible factors-1a. Tumor cell growth after knockdown of HIF-1α was measured by MTT assay. Tumor cells were susceptible to growth inhibition under hypoxic conditions when more than 100 nM of siHIF-1α was transfected. However, normoxic conditions (100 nM) did not show significant difference. (TIF)

File S2 Confirmation of dimerization with HIF-1α and HIF-1β by Immunoprecipitation. HIF-1α was bounded with mouse anti-HIF-1α using IgG beads. siHIF-1α or siHIF-1β group weakly detect HIF-1α or HIF-1β band. But, HIF-1α/HIF-1β strongly expressed over 2–3 times in the control group. (TIF)

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Author Contributions

Conceived and designed the experiments: SHC, JYP, KHH. Performed the experiments: SHC, ARC, MSL, SWH. Analyzed the data: SHC, WK, JYP. Contributed reagents/materials/analysis tools: JYP, SHA, KHH. Experiments: SHC, ARC, MSL, SWH. Performed the experiments: SHC, JYP, KHH. Conceived and designed the experiments: SHC, JYP, KHH. The authors are grateful to Dong-Su Jang (Medical Illustrator, Department of Research Affairs, Yonsei University College of Medicine, Seoul, Korea) for his help with the figures.

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