Resistance to cisplatin-induced apoptosis via PI3K-dependent survivin expression in a rat hepatoma cell line

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Abstract. Hepatocellular carcinoma (HCC) is known to be resistant to chemotherapy. Survivin, a member of the inhibitor of apoptosis proteins, is overexpressed in most cancers but is absent in most normal adult tissue. The aim of this study was to investigate whether expression of survivin contributes to resistance to cisplatin-induced apoptosis. We confirmed induction of survivin expression in hepatoma in the N-diethylnitrosamine (DEN) induced rat and in the rat hepatoma cell line (K-251). We examined cell proliferation after treatment with cisplatin (CDDP) in the presence and absence of siRNA or the phosphatidylinositol 3-kinase (PI3K) inhibitor LY294002 to suppress survivin or PI3K/Akt, respectively. Survivin was expressed in DEN-induced rat HCC with RT-PCR and Western blotting. Expression of survivin was observed primarily in the nuclei and in the cytoplasm with immunohistochemistry. However, survivin was not detected in non-tumor tissues. Expression of survivin was also observed primarily in the nuclei and in the cytoplasm of the K-251 rat hepatoma cell line. CDDP induced survivin expression, which was blocked by siRNA. LY294002 also attenuated survivin expression induced by CDDP. Our results indicate that survivin expression via PI3K contributes to resistance to CDDP-induced apoptosis in a rat hepatoma cell line.

Introduction

Hepatocellular carcinoma (HCC) is one of the most common cancers in the world, accounting for approximately 6% of all human cancers and 1 million deaths annually, with an estimated number of new cases of over 500,000 per year (1,2). Although the clinical management for early-stage HCC has improved significantly, the prognosis of HCC is still extremely poor and is the third highest cause of cancer-related mortality. HCC exhibits resistance to chemotherapy, such as cisplatin (CDDP). Therefore, investigating and finding new treatment for HCC is important.

Cell proliferation is determined not only by the rate of cell growth but also by the rate of cell loss. Apoptosis is a major component of this regulatory process, which consists of 2 converging cascades ‘death-receptor pathway’ and ‘mitochondrial pathway’ (3,4). The death-receptor pathway is triggered by the interaction of death ligands of the tumor necrosis factor family with their cognate receptors that assemble the death-inducing signaling complex, resulting in activation of caspase-8. The mitochondrial pathway involves mitochondria that release apoptogenic factors, such as cytochrome c and Smac/DIABLO, in response to noxious stimuli.

Members of the IAP family are known to inhibit death receptors and the mitochondrial pathway by acting as endogenous inhibitors of caspases (4,5). Survivin is a bifunctional protein that controls cell division and inhibits apoptosis (6). The mechanism by which survivin inhibits apoptosis remains in dispute. While it is capable of binding to effector caspases under cell-free conditions, under more physiological conditions it inhibits apoptosis by binding to the second mitochondrial activator of caspases (Smac) (7). Survivin is present in only small amounts in terminally-differentiated normal cells, but is found to be prominently overexpressed in a variety of tumors, including cancers of the lung, stomach, breast, melanoma, pancreas, esophagus and colon (8,15). Overexpression of survivin has been identified as a negative prognostic factor in various cancer types (8,14,16), and is implicated in resistance to the induction of apoptosis by anti-cancer agents (17).

Despite increasing evidence in support of survivin as a promising target for molecular intervention, the mechanism of survivin overexpression in HCC and its implication in drug resistance remain to be investigated. Recent reports have demonstrated that survivin was up-regulated via the PI3K/Akt pathway in myeloid leukemia cells (18), endothelial cells (19), and prostate cancer cells (20). Furthermore, Akt is also activated in response to stress by UV irradiation and chemotherapy (21-23). Whether activation of Akt constitutes a physiological stress response that enables cancer cells to evade apoptosis by up-regulation of survivin remains to be demonstrated.

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The purpose of this study was to clarify whether survivin is overexpressed in rat HCCs and if expression of survivin contributes to resistance to cisplatin-induced apoptosis in HCC cells. We investigated alterations in cell morphology and proliferation in HCC cells in the presence and absence of siRNA against survivin and a PI3K inhibitor.

Materials and methods

Rat DEN-induced HCC and HCC cell lines. Rat HCC was induced with N-diethylnitrosamine (DEN, Sigma, St. Louis, MO, USA). SD rats (4-week-old males, weighing 100 g; SLC, Hamamatsu, Japan) were provided with 100 ppm DEN ad libitum in drinking water for 8 weeks and were maintained for an additional 4 weeks without administration of DEN (24-26).

The DEN-induced rat HCC cell line K-251 was obtained from Cell Resource Center for Biomedical Research Institute of Development, Aging and Cancer Tohoku University. The cell line K-251 was grown in RPMI-1640 with L-glutamine (Invitrogen Corporation, Gibco 21875-034) supplemented with 10% FBS and 1% (w/v) penicillin/streptomycin (PC/SM) at 37°C in a humidified atmosphere containing 5% CO2. Cells were treated with CDDP (Nippon Kayaku, Tokyo, Japan) and LY294002 (Cellbiochem-Novabiochem. Corp.). Cells were treated with the PI3K inhibitor LY294002 dissolved in DMSO.

Quantitative real-time RT-PCR. Total RNA was extracted from the liver tissues or cultured cells with TRizol (Invitrogen, Tokyo, Japan). Total RNA (2 μg) was reverse transcribed to cDNA with an Omniscript RT kit (Qiagen, Tokyo, Japan) with oligo (dT)12 primers (Invitrogen). Real-time PCR was performed with a QuantiTect SYBR Green PCR kit (Qiagen) and an ABI PRISM 7700 Sequence Detection System (Applied Biosystems, Tokyo, Japan). RT-PCR was performed with an initial step at 95°C for 15 min, followed by 40 cycles at 95°C for 20 sec, 59°C for 30 sec, and 72°C for 1 min with a final step at 72°C for 10 min. Two specific primers were used to identify survivin (Forward: 5'-TAAGCCACTTGTC CCAGCTT-3'; Reverse: 5'-CTCATCCACTCCCTTCCTC-3') and G3PDH (Forward: 5'-ACCACGTCATGCCC CAT CAC-3'; Reverse: 5'-TCCACCCGCTTGGCGTA-3'). The expression levels were calculated after conversion to numerical values by ABI PRISM 7700 SDS software and are expressed as ratios relative to the expression of G3PDH.

Western blot analysis. Frozen liver tissues or cultured hepatocytes were homogenized in lysis buffer containing 50 mM Tris-HCl, 2% sodium dodecylsulfate and 10% glycerol, and then boiled for 2 min. After the protein concentration in the sample was determined, 0.1% bromophenol blue and 6% 2-mercaptoethanol were added. Subsequently, 20 μg of each sample was electrophoresed and transferred onto PVDF membranes (Immobilon, Millipore, Billerica, MA). The membranes were blocked with 5% skim milk in PBS and incubated at 4°C overnight with the following primary antibodies and dilutions: anti-survivin antibody (no. sc-10811; Santa Cruz Biotechnology, Santa Cruz, CA; 1:200); anti-cleaved caspase-3 antibody (no. 9661; Cell Signaling Technology; 1:1000); anti-Parp antibody (no. 9542; Cell Signaling Technology; 1:1000); anti-phospho Akt antibody (no. 9271; Cell Signaling Technology; 1:500); anti-XIAP antibody (no. 2042; Cell Signaling Technology; 1:1000); anti-α-tubulin antibody (no. CP06; Calbiochem; 1:1000); and anti-GAPDH antibody (no. sc-20357; Santa Cruz Biotechnology; 1:200). After washing, the membranes were incubated with secondary antibodies conjugated with horseradish peroxidase. Chemiluminescence detection was performed using ECL Western Blotting Detection Regents (Amersham, Buckinghamshire, UK), according to the manufacturer’s instructions. The intensity of the band was quantified with National Institutes of Health Image (NIH-Image) and normalized to GAPDH as an internal control.

Immunohistochemistry. The specimens were fixed in 10% formalin and embedded in paraffin. They were subsequently deparaffinized and endogenous peroxidase was quenched with 0.3% hydrogen peroxide in methanol at room temperature for 15 min. The antigen was retrieved by incubation in citric acid buffer at 90°C for 20 min. After blocking, the sections were incubated with primary antibody recognizing survivin (no. sc-10811; Santa Cruz Biotechnology, Santa Cruz, CA; 1:200 dilution) overnight at 4°C and then with labeled polymer in an Envision+ System HRP (Dako, Tokyo, Japan) at room temperature for 1 h. The sections were examined after incubation with a Liquid DAB Substrate Chromogen System (Dako) for 2 sec.

siRNA for survivin transfection. siRNA with the sequence GGCACUUAAGCACUCAGGAA targeting the survivin mRNA and the negative control sequence ATCCGCGCGAT AGTACGTA were purchased from B-Bridge International, Inc. Cells were transfected with Lipofectamine 2000 (Invitrogen) according to the reverse transfection method (27) using Opti-MEM (Invitrogen Corp., Gibco 31985-070) and Insulin-Transferrin-Selenium (ITS; Invitrogen Corp., Gibco 41400-045). Briefly, cells were plated at 2x10^4 cells/well in 6-well plates. Cells were transfected for 6 h, respectively. After transfection, cells were grown in RPMI-1640 with L-glutamine supplemented with ITS and 1% (w/v) penicillin/ streptomycin. Samples were collected 24 h after transfection. Experiments were repeated at least 3 times.

Cell proliferation assay. Cell proliferation was determined using the Cell Counting kit-8 (CCK-8; Dojindo, Kumamoto, Japan) to count living cells. Cells were cultured and treated in 24-well plates. At 0, 12, 24, 48 and 72 h, the CCK-8 was used to determine cell proliferation according to the manufacturer’s instruction.

Statistical analysis. All data are expressed as the means ± SEM, and the statistical significance of differences between groups was assessed by Mann-Whitney U-test. P<0.05 were regarded as statistically significant.

Results

Expression of survivin in DEN-induced hepatoma and in the rat hepatoma cell line. We first used RT-PCR to investigate the level of survivin messenger RNA (mRNA) in DEN-induced
rat HCC. A 75-bp fragment of survivin was detected in rat HCC (Fig. 1A). However, survivin mRNA was faintly detected in non-tumor tissues. Furthermore, a 16.5 kDa fragment of survivin protein was detected in rat HCC with Western blotting (Fig. 1B), while survivin was not detected in non-tumor tissues. We performed immunohistochemical staining to determine the cellular localization of survivin. Expression of survivin was observed primarily in the nuclei and in the cytoplasm of HCC cells. In contrast, survivin was not expressed in the adjacent non-cancer tissues (Fig. 1C). Furthermore, we examined the survivin expression in rat hepatoma cell line (K-251) derived from DEN-induced HCC using immunocytochemical staining. Expression of survivin was observed primarily in the nuclei and in the cytoplasm (Fig. 1D).

**Down-regulation of survivin by siRNA in the rat hepatoma cell line.** In rat HCC cells, the efficiency of transfection was about 80-90%, using red-labeled double-stranded RNA (dsRNA) oligomer by Lipofectamine 2000 (data not shown). Real-time PCR demonstrated that transfection with siRNA targeting survivin attenuated the expression of survivin mRNA to 68.1±8.5% of the expression level in cells transfected with negative control siRNA (Fig. 2A). Similarly, Western blotting showed that survivin expression was inhibited to 34.5±7.5% of control cells (Fig. 2B). The CCK-8 assay showed that the cell number decreased to 10.6±7.8% and 62.7±6.1% at 24 and 48 h after transfection, respectively, compared with control cells (Fig. 2C). These results indicate that transfection with siRNA significantly inhibited cell proliferation.

**CDDP treatment up-regulates survivin expression and the silencing of RNA for survivin sensitizes CDDP-induced apoptosis in cancer cells.** Treatment with CDDP induced over-expression of survivin protein in K-251 cells (189.6±26.1% expression relative to cells without CDDP). Similarly, the induction of survivin was observed in cells transfected with negative control siRNA and siRNA against survivin (167.7±26.9%, 149.5±23.9%, respectively) (Fig. 3).

We examined fluorescence staining of nuclei using Hoechst 33342 and propidium iodide (PI) at 12 h after transfection with siRNA and treatment with CDDP. Irregular staining of the nuclei, nuclear fragmentation, and blebbing phenomenon were observed in cells treated with siRNA for survivin and CDDP, indicating apoptotic cell death (Fig. 4A). Percentage of apoptotic cells treated with negative control siRNA, negative control siRNA + CDDP, siRNA for survivin, siRNA for survivin + CDDP were 2.7±0.91%, 8.47±0.77%, 25.74±6.57%, 33.8±5.88%, respectively (Fig. 4B). The CCK-8 assay showed that 48 h after transfection with siRNA for survivin and treatment with CDDP cell proliferation was attenuated to 85.69±1.80% compared with cells transfected with negative control siRNA (Fig. 4C). Similarly, 48 h after transfection with siRNA for survivin and treatment with CDDP cell proliferation was attenuated to 78.15±1.12% compared with CDDP alone (data not shown).

**siRNA for survivin induces caspase activation.** Western blot analysis demonstrated that expression of cleaved caspase-3 was up-regulated in cells treated with siRNA for survivin compared to those treated with negative control siRNA. Similarly, siRNA for survivin increased the expression of cleaved caspase-3 in cells treated with CDDP, compared with those treated with negative control siRNA (Fig. 5A). Furthermore, expression of cleaved PARP was also up-regulated in cells treated with siRNA for survivin compared to those treated with negative control siRNA, in cells treated with/without CDDP (Fig. 5B). CDDP up-regulated the expression of phospho-Akt in cells treated with negative...
Figure 2. Down-regulation of survivin by siRNA in the DEN-induced rat hepatoma cell line. (A) Quantitative real-time PCR of survivin mRNA in un-treated HCC cells, in HCC cells treated with negative control siRNA and in HCC cells treated with siRNA for survivin. Data are expressed as the mean ± SEM (n=5). (B) Western blot analysis of survivin protein in un-treated HCC cells, in HCC cells treated with negative control siRNA and in HCC cells treated with siRNA for survivin. Data are expressed as the mean ± SEM (n=4). (C) CCK-8 cell proliferation assay of un-treated HCC cells, HCC cells treated with negative control siRNA, and HCC cells treated with siRNA for survivin. Data are expressed as the mean ± SEM (n=3).

Figure 3. CDDP up-regulates survivin expression. Western blot analysis of survivin protein in untreated HCC cells, HCC cells treated with CDDP (1 μM), HCC cells treated with negative control siRNA (50 μM), HCC cells treated with negative control siRNA (50 μM) + CDDP (1 μM), HCC cells treated with siRNA for survivin (50 μM), and HCC cells treated with siRNA for survivin (50 μM) + CDDP (1 μM). Data are expressed as the mean ± SEM (n=4).
control siRNA or siRNA for survivin. However, the expression level was similar between cells treated with negative control siRNA and siRNA for survivin, indicating that survivin did not affect phosphorylation of Akt in cells in the absence of treatment with CDDP (Fig. 5C). There was no significant difference of XIAP expression between cells treated with negative control siRNA or siRNA for survivin (Fig. 5D).

Regulation of survivin by PI3K/Akt. Rat HCC cells were treated with 25 μM LY294002, phosphoinositide-3 kinase inhibitor, to elucidate the relationship between survivin and PI3K. LY294002 blocked the expression of survivin (Fig. 6A). Similarly, expression of phospho-Akt was down-regulated in cells treated with LY294002 compared to those with treated with DMSO, in cells treated with/without CDDP (Fig. 6B). On the other hand, expression of XIAP was up-regulated in cells treated with LY294002 compared to those with treated with DMSO, and LY294002 increased expression of XIAP in cells treated with CDDP (Fig. 6C).

Discussion

We confirmed overexpression of survivin in DEN-induced rat HCC in vivo and in DEN-induced rat HCC cells in vitro.
There was little survivin expression in non-tumor liver tissues from rat treated with DEN. Overexpression of survivin occurs frequently during cellular promotion of malignant neoplasms and has been identified as a negative prognostic factor in various cancer types (8,14,16). In our study, suppression of survivin expression in rat HCC cells treated with siRNA for survivin attenuated cell proliferation of cancer cells. Survivin was overexpressed in rat HCC cells treated with CDDP, which was down-regulated by treatment with siRNA for survivin. Furthermore, siRNA for survivin sensitized rat HCC cells to CDDP-induced apoptosis. Our data show that CDDP or siRNA for survivin induced apoptosis in rat HCC cells, 24 h after treatment. Data are expressed as the mean ± SEM (n=4).

Figure 5. siRNA for survivin and CDDP induce apoptosis. Western blot analysis of cleaved caspase-3 (A), cleaved PARP (B), phospho-Akt (C) and XIAP (D) in HCC cells, 24 h after treatment. Data are expressed as the mean ± SEM (n=4).
K-251 and that CDDP also activated Akt, resulting in increased survivin and p-Akt levels. Inhibition of the PI3K/Akt pathway decreased overexpression of survivin induced by CDDP and sensitized HCC to CDDP. Therefore, we concluded that survivin is regulated by the PI3K/Akt pathway and that inhibition of survivin expression with siRNA sensitized HCC cells to CDDP-induced apoptosis via the PI3K/Akt-dependent pathway. However, whether PI3K/Akt up-regulates survivin directly, or indirectly remains to be investigated.

We demonstrated that overexpression of survivin and its anti-apoptotic activity in rat HCC cells were regulated by PI3K/Akt signaling. These findings have significant clinical implications, since they provide direct evidence that CDDP itself can trigger resistance in HCC cells which can diminish its therapeutic efficacy. However, inhibition of the PI3K/Akt pathway may have clinical limitations, because of their essential function in all types of cells in a complicated largely unresolved signaling network, which would reduce tumor-specificity. Our results demonstrated that survivin-specific siRNA down-regulates the overexpression of survivin induced by CDDP, and sensitize HCC cells to CDDP-induced apoptosis. The trans-catheter arterial chemo-embolization therapy has already been applied for practical use, and we previously reported that hydro-administration of an siRNA expressing plasmid into the hepatic artery was a valid method for targeting rat HCC (26). Combination therapy of trans-catheter arterial chemo-embolization and siRNA for survivin, or hydro-administration injection and siRNA for survivin may be beneficial in the treatment of HCC. Furthermore, non-cancerous liver tissues may not respond to siRNA for survivin because they do not endogenously express survivin. Therefore, down-regulation of survivin has the potential to enhance the efficacy of chemotherapy in the treatment of HCC and may have significant clinical impact.

References