We already had reported that Bcl-w promotes invasion or migration in gastric cancer cells and glioblastoma multiforme (GBM) by activating matrix metalloproteinase-2 (MMP-2) via specificity protein 1 (Sp1) or β-catenin, respectively. High expression of Bcl-w also has been reported in GBM which is the most common malignant brain tumor and exhibits aggressive and invasive behavior. These reports propose that Bcl-w-induced signaling is strongly associated with aggressive characteristic of GBM. We demonstrated that Sp1 protein or mRNA expression is induced by Bcl-w using Western blotting or RT-PCR, respectively, and markedly elevated in high-grade glioma specimens compared with low-grade glioma tissues using tissue array. However, relationship between Bcl-w-related signaling and aggressive characteristic of GBM is poorly characterized. This study suggested that Bcl-w-induced Sp1 activation promoted expression of glioma stem-like cell markers, such as Musashi, Nanog, Oct4 and sox-2, as well as neurosphere formation and invasiveness, using western blotting, neurosphere formation assay, or invasion assay, culminating in their aggressive behavior. Therefore, Bcl-w-induced Sp1 activation is proposed as a putative marker for aggressiveness of GBM.

INTRODUCTION

Glioblastoma multiforme is the most common and lethal type of malignant brain tumor, exhibiting highly proliferative growth and widespread invasive potential, culminating in vigorous tumor progression. Despite the commonly available treatments of surgical resection, radiation and chemotherapy, GBM is difficult to treat because of its high frequency of recurrence (Fumari et al., 2007).

Current evidence suggests that the B-cell lymphoma 2 homolog (Bcl-w) enhances cell survivability through its anti-apoptotic activity, as well as the migratory and invasive potential of cancer cells. Bcl-w is expressed in a variety of cancer types, including gastric cancer (Lee et al., 2003), colorectal adenocarcinomas (Wilson et al., 2000) and glioblastoma multiforme (Hoellinger et al., 2005; Lee et al., 2013), in a cancer cell-specific manner. We have previously reported that Bcl-w enhances migratory and invasive potential in gastric cancer cells by inducing the production of several types of extracellular matrix (ECM)-degrading proteinases (Bae et al., 2006; 2009). These observations support the claim that expressions of Bcl-w-induced signaling components in GBM are strongly correlated with GBM invasiveness. Growing evidence indicates that, in addition to Bcl-w expression, Sp1 and MMP-2 expressions are upregulated in glioma tissue. Specially, Sp1 plays a critical role in the regulation of multiple genes implicated in tumorigenesis (Safe and Abdelrahim, 2005). In several human cancer types, including breast cancer (Zannetti et al., 2000), thyroid tumor (Chiefari and Abdelrahim, 2004), gastric cancer (Yao et al., 2004), epidermal squamous cell carcinoma (Kumar and Butler, 1999) and pancreatic cancer (Shi et al., 2001), an elevation of Sp1 expression levels has been reported, which, in some instances, has been correlated with poor clinical diagnosis (Guan et al., 2011; Wang et al., 2003). Sp1 transcriptional activity is crucial role for tumor progression such as proliferation, tumor formation, migration, mesenchymal transition, angiogenesis and metastasis in various human cancer cell types (Abdelrahim et al., 2004; Jungert et al., 2007; Lou et al., 2005; Shin et al., 2011; Wang et al., 2005; Wei et al., 2004).

However, no study, to our knowledge, has examined the relationship between glioma cell characteristics and upregulated proteins, such as Bcl-w, Sp1 and MMP-2 in glioma. Based on our findings, we conclude that Bcl-w-induced Sp1 expression is critical for the development of GBM characteristics, because it functions as a positive regulator for the maintenance of stemness. Therefore, our data suggest that Bcl-w and Sp1 represent potential prognostic markers for glioma progression.
MATERIALS AND METHODS

Cell culture, transfection, and treatments
The U251 were obtained from the Korean Cell Line Bank (KCLB) and cultured in Minimum Essential Medium Eagle (MEM) (Mediatech, Inc., USA) containing 10% fetal bovine serum (FBS) (Tissue Culture Biologicals, USA) and penicillin-streptomycin antibiotics (PAA Laboratories GmbH, Austria) under a humidiﬁed 5% CO2 incubator at 37°C. U251 cells were transfected with either empty pcDNA vector or that Bcl-w cDNA. The control and Bcl-w-overexpressing cells were transiently transfected using Lipofectamine 2000 or RNAi MAX (Invitrogen, USA) with the indicated expression constructs or chemically synthesized small interfering RNAs (siRNAs). Small interfering RNAs were purchased from as the following: silencer negative control siRNA, si-Bcl-w (Ambion, USA), si-Sp1 and si-MMP-2 (Santa Cruz Biotechnology, USA).

Western blot analysis
The cells were lysed in a buffer containing 50 mM HEPES, pH 7.4, 100 mM NaCl, 1% Triton X-100, 10% glycerol, 1 mM EDTA, 20 mM l-glutamic acid, 5 mM NaF, 2 μg/ml aprotonin, 2 μg/ml leupeptin and 100 mM phenylmethylsulfonyl ﬂuoride. Prepared cell lysates were subjected to SDS-PAGE and electrophoresed to Immobilon membranes (Millipore, USA), then that incubated with the indicated antibodies and visualized by the ECL detection system (Advanta, USA).

Antibodies and inhibitors
The antibodies were used as follows; anti-Sp1 was purchased from Santa Cruz Biotechnology (USA). Anti-Bcl-w and anti-Musashi were obtained from Cell Signaling technology (USA). Anti-β-actin was purchased from Sigma-Aldrich (USA). Anti-Oct3/4 was obtained from Novous Biologicals (Vittleton, Co). Anti-Nanog was purchased from Chemicon International (USA). Anti-Sox2 was purchased from R & D Systems (USA). Anti-MMP-2 was obtained from Calbiochem (USA). The pharmacological inhibitors were used in this study; mithramycin A (MMA) was purchased from Sigma-Aldrich (USA). Oleyl-N-hydroxylamide (OA-Hy) was obtained from Calbiochem (USA).

Quantitative reverse transcription polymerase chain reaction (RT-PCR) analysis
Total RNA was isolated from cultured U251 cells using Trizol (MRC Gene, USA), of which 100 ng was converted to cDNA with reverse transcriptase. EcoNtAQ PLUS GREEN 2x master mix (Lucigen, USA) was added to the reaction mix, and quantitative RT-PCR analysis was performed using primers for Sp1: forward 5′-TGT AAA AAG GAG TGG TG C-3′, reverse 5′-CTT GGT GCT TAA GTT GAG AG-3′, Musashi: forward 5′-ACA GCC ATT GGA GAG TGA C-3′, reverse 5′-GGT GAA GGC TG TGG GTC CTA C-3′, Nanog forward 5′-GCT TCG TTT TTG AAG CA-3′, reverse 5′-GGTT GTG GCA TCT GCT GGA GG-3′, Oct4: forward 5′-GAG CAA AAG CCG GAG GAC T-3′ reverse 5′-TTC TCT TCT GGC CCT GCA C-3′ and Sox2: forward 5′-ATG CAC CGC TAC GAG GTG A-3′, reverse 5′-CTT TGG CAC CCC TCC CAT T-3′. Quantitative measurement of mRNA was obtained from 3 independent experiments. The expression levels of mRNAs specific for each gene were normalized to the expression of GAPDH (glyceraldehydes 3-phosphate dehydrogenase).

Invasion assay
These assays were conducted as described previously. The 2 × 105 cells were seeded onto the Matrigel-coated polycarbonate ﬁlters which were placed in modiﬁed Boyden chambers (Corning, USA) that contain ECM components (BD Biosciences, USA). After 20-24 h of incubation at 37°C, the cells that had invaded to the lower surface of the membrane were ﬁxed and stained using a Diff-Quick kit (Fisher Scientiﬁc, USA), then counted under a microscope (Mitoti AE31 series, Trinocular inverted MIC). Five independent experiments were analyzed for statistical signiﬁcance using the Student’s t-test. Differences were considered to be statistically signiﬁcant at p < 0.05.

Immunohistochemistry
To determine the levels of endogenous Sp1 in human brain cancer tissues, premade AccuMax array Brain cancer tissues for formalin ﬁxed parafﬁn embedded slides (A221-V) were commercially purchased from ISU Abxis Co. Ltd (Korea). The tissue array contained 35 brain cancer specimens, including astrocytoma (WHO grade I), anaplastic astrocytoma (WHO grade III) and glioblastoma multiforme (WHO grade IV). Tissue sections were incubated with REAL Peroxidase-blocking solution (Dako, Cat # S2023, Denmark A/S produktionsvej 42) to quench the endogenous peroxidase activity. Tissues were immunoreacted with anti-Sp1 antibody (Santa Cruz Biotechnology, USA; 1:200). The levels of endogenous Sp1 were stained with 3,3′-diaminobenzidine (DAB) substrate kit for peroxidase (Vector Laboratories, USA) after conjugation using avidin-biotin-peroxidase complex (VECTASTATIN Universal Elite ABC Kits; Vector Laboratories, USA). Slides were counterstained with hematoxylin, dehydrated, and mounted with Pemount solution (Fisher Scientiﬁc, USA). Images were acquired by Olympus BX53 ﬁrscopery using cellsSens Standard controller software (Olympus). Sp1-positive cells were counted in three different horizontally adjacent ﬁelds. The mean of triplicate experiments was signiﬁcantly different depending on grades of glioma tissues.

Spheres culture
The vector control or the Bcl-w overexpressing U251 cells were suspended in Dulbecco’s modiﬁed Eagles medium ﬂuid-1 (Cellgro, USA) containing 20 ng/ml each of epidermal growth factor (EGF, Biovision, USA), basic ﬁbroblast growth factor (bFGF, Biovision) and B27 (1:50) (GIBCO, USA) as a stem-cell-permissive medium. Spheres were harvested after 2 to 4 days and protein extracts were subject to Western blotting with appropriate antibodies.

Sphere formation assay
For generating sphere formation, sphere cells were dissociated with Accutase (Innovative Cell Technologies, Inc) for expansion. Sphere-forming U251 cells were ﬁrst distributed into 96-well plates at a density of 1-2 cells per well. After 12h, individual wells were visually checked for the presence of a single cell. The clones were grown and tested for the formation sphere. After 20 days until the sphere formation, spheres were attached by horizontally adjacent ﬁelds. The mean of triplicate experiments was signiﬁcantly different depending on grades of glioma tissues.

Gelatin zymography
Conditioned media were prepared by incubating cells in serum-free media for 24 h. The media were supplemented with speciﬁc inhibitors or targeting siRNA. Equal volumes of conditioned media were then subjected to 8% SDS-PAGE containing 0.1%
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RESULTS
Bcl-w-induced Sp1 expression is involved in the invasiveness of GBM

In this study, we investigated the relationship between Bcl-w and Sp1 that resulted in the up-regulation of glioma tissues. To establish the effect of Bcl-w on Sp1 expression and aggressiveness as manifested by invasive potential in glioblastoma, Bcl-w was overexpressed in U251 glioma cells. Bcl-w-overexpressing cells demonstrated enhanced Sp1 protein and mRNA expression and glioblastoma invasiveness, as measured by an invasion assay using Matrigel-coated polycarbonate filters (Fig. 1A). The Bcl-w-induced invasiveness of cells treated with Bcl-w targeted small interference RNAs (siRNAs) was effectively attenuated Sp1 protein and mRNA expression and glioblastoma invasiveness (Fig. 1B). These observations strongly suggest that Bcl-w induces the upregulation of Sp1 expression by the transcriptional mechanism. To confirm the pattern of Sp1 expression in various grades of brain cancer tissues (WHO I, III, and IV), we performed a immunohistochemistry assay using premade AccuMax array slides containing 35 specimen from patients with brain tumors. Sp1 expression was up-regulated in glioblastoma tissues (glioma grade IV) compared with glioma tissues (grade I/III) in every sample examined as low/high magnification (Fig. 1C). Notably, the subcellular localization of Sp1, a transcription factor, was apparent primarily in the nucleus, as expected. These results indicate that Sp1 expression levels are closely correlated with tumor grading in patients with glioma.

Bcl-w-induced Sp1 contributes to stemness of glioblastoma cells

These stemness-related results were used spheres-cultured U251 glioma cells. To examine the effect of Bcl-w-induced Sp1 expression in the enhancement of aggressiveness, Sp1-depleted activation by pharmacological inhibition with mithramycin A (MMA; 10 μmol/L) (Fig. 2A) or Sp1 RNA interference (Fig. 2B) attenuated not only sphere-formation of stem-like cell popu-
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The results suggest that Bcl-w-induced Sp1 expression promotes cancer stem cell markers expression in the transcriptional levels.

Sp1 implicates in Bcl-w-induced invasion by activating MMP-2
To evaluate the signaling mechanism of Sp1 on Bcl-w-induced invasion in glioma, we subsequently performed a Sp1-depleted activation assay by pharmacological inhibition with mithramycin A (10 μmol/L) or RNA interference. Sp1-depleted activation not only inhibited the enzymatic activity of MMP-2, but also repressed Bcl-w-induced invasion (Figs. 3A and 3B). The blockage of MMP-2 by the pharmacological inhibitor OA-Hy (20 μmol/L) or RNA interference attenuated the invasiveness of glioblastoma cells (Figs. 4A and 4B). These sequential events suggest that Bcl-w-induced Sp1 activation contributed to Bcl-w-induced invasion via MMP-2 production in other glioma cell lines including U373 and U87MG (Lee et al., 2013). These results show that Bcl-w-induced Sp1 is implicated in the aggressive characteristics of glioma cells (Fig. 5).

DISCUSSION

Bcl-w is expressed in a variety of cancer types, especially gastric and colorectal adenocarcinomas as well as GBM (Hoelzinger et al., 2005; Lee et al., 2003; Wilson et al., 2000). We previously reported that Bcl-w enhances the migratory and invasive potential, as well as survivability, of gastric cancer cells (Bae et al., 2006). Also, we studied that Bcl-w promotes mesenchymal changes and invasiveness of glioblastoma cells by inducing nuclear accumulation of β-catenin (Lee et al., 2013). In addition, it has been reported that level of Bcl-w or Sp1 is overexpressed in GBM (Guan et al., 2011). However, no study has examined the relationship between Bcl-w and Sp1. We showed that Bcl-w increased Sp1 protein and mRNA expression, resulting in increased migratory and invasive potential in glioblasto-
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Fig. 3. Sp1 is implicated in Bcl-w-induced invasion upstream of MMP-2. (A) The indicated U251 cell transfectants were incubated in a serum-free medium in the presence or absence of the Sp1 inhibitor mithramycin A (MMA; 10 μmol/L) for 1 or 24 h. Expression levels and activities of Sp1 and MMP-2 were compared using Western blotting and zymography assay using β-actin as the loading control or Ponceau S staining, respectively. The invasive potential of treated cells was compared. *p < 0.05 versus untreated control, n = 5. (B) Sp1 siRNAs were introduced into vector or Bcl-w overexpressing cells, and cellular levels of Sp1 and activities of MMP-2 were compared after 24 h of incubation by using Western blotting. The invasive potential of indicated transfectants was compared. *p < 0.05, n = 5.

Fig. 4. Activation of MMP-2 mediates Bcl-w-induced invasion. (A) The indicated U251 cell transfectants were incubated in serum-free medium in the presence or absence of aMMP-2 inhibitor (OA-Hy; 10 μmol/L) for 1 or 24 h. Expression levels of MMP-2 were compared using Western blotting. Invasion assays were performed using MMP-2 inhibitor-treated and untreated cells. **p < 0.005 versus untreated control, n = 5. (B) MMP-2 targeting siRNA was introduced into the indicated cells. The levels of MMP-2 were compared after 24 h of incubation by using Western blotting. The invasive potential of indicated transfectants was compared. *p < 0.05, n = 5.

Sp1 expression level is often greater in various cancer tissues or cells than in normal tissues or cells (Chuang et al., 2009; Davie et al., 2008; Kong et al., 2010). Therefore, we quantitatively examined Sp1 expression levels in grade I, III and IV of glioma tissues in order to determine relationship between expression levels of Sp1 and various grades of glioma tissues. The level of Sp1 expression was upregulated in glioma grade IV compared with grade I/III in every patient examined (Fig. 1C) as well as Bcl-w expression (Lee et al., 2013). These results suggest that Bcl-w and Sp1 expression closely correlated with grading of glioma in the patients. Guan et al’s group reported that Sp1 expression levels, elevated in glioma tissues and cell lines, were correlated with the WHO grading system for cancer cells (Guan et al., 2011).
induced Sp1 is implicated in the aggressive characteristics of glioma cells (Fig. 5). Therefore, Sp1 may be regarded as a putative marker for aggressiveness of GBM. Future studies may investigate whether the Bcl-w-induced signaling components are related to the maintenance of stemness in GBM and how Bcl-w induces Sp1 expression.

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