Abstract. Yes-associated protein (YAP) positivity indicates a poor prognosis in gastric cancer. Transcriptional co-activator with a PDZ-binding domain (TAZ), a YAP paralog, is highly expressed in gastric signet ring cell carcinoma. Verteporfin (VP), a clinical photosensitizer, was recently shown to inhibit YAP/TAZ. In the present study, the therapeutic potential of VP treatment was explored using two gastric cancer cell lines: MKN-45 (TAZ-dominant) and MKN-74 (YAP-dominant). Cell proliferation was evaluated by MTS assay. Vascular mimicry was evaluated by the tube formation assay. Gene and protein expression levels of YAP/TAZ downstream effectors [such as Survivin, Cysteine-rich angiogenic inducer 61 (CYR61), and connective tissue growth factor (CTGF)] were measured. YAP or TAZ localization was evaluated by immunofluorescence. Cell death was assessed by immunofluorescent staining of Annexin V. YAP and TAZ expression were knocked down by small interfering RNA. The current results demonstrate that MKN-45, a poorly differentiated TAZ-dominant gastric cancer cell line, was more sensitive to VP than MKN-74, a moderately differentiated YAP-dominant gastric cancer cell line. VP changed the localization of YAP/TAZ, promoted its degradation and significantly decreased the protein level of Survivin in both cell lines. Cell death was induced by VP treatment in a dose-dependent manner. Vascular mimicry was inhibited in both cell lines. Proliferation in both cell lines decreased in response to YAP/TAZ knockdown. The present study indicated that VP has potential as a therapeutic agent in YAP- and TAZ-dominant gastric cancers due to its ability to suppress the anti-apoptotic protein Survivin via inhibition of YAP and TAZ.

Introduction

Gastric cancer has the fifth highest incidence among all cancers according to GLOBOCAN 2020 data. Globally, it is the fourth most frequent cause of death among malignancies (1). Although the incidence of gastric cancer is decreasing, it remains a significant public health problem (2). Gastric cancer is commonly diagnosed at an advanced stage, except in Japan and Korea where routine screening is performed (3,4). The efficacy of chemotherapy for advanced-stage gastric cancer has increased the survival rate in recent years; however, the degree of improvement is unsatisfactory (5,6).

The Hippo signaling pathway is essential in the control of organ size. It consists of a series of serine/threonine kinases and scaffolding proteins that regulate the subcellular localization and activity of the effector proteins yes-associated protein (YAP) and transcriptional co-activator with a PDZ-binding domain (TAZ) (7). Overexpression of YAP is observed frequently in a variety of cancer types (8). YAP expression is elevated in gastric adenocarcinomas, and its knockdown inhibits gastric cancer cell proliferation (9). High levels of YAP in the nucleus have been linked to chemotherapy resistance in various cancer types (10-13). In gastric cancer cells, YAP is associated with cisplatin and trastuzumab resistance (14,15). Upregulation of YAP correlates with progression, metastasis and poor prognosis in patients with gastric carcinoma (16). Moreover, the expression of TAZ, a YAP paralog, is elevated in gastric signet ring cell carcinoma (17).

Targeting YAP and TAZ, individually or in combination, is likely to have a clinical impact on gastric cancer. Verteporfin (VP), a benzoporphyrin derivative, is used in photodynamic therapy (PDT) for exudative age-related macular degeneration; specifically, against choroidal neovascularization (18).
We have recently reported the efficacy of VP in PDT using gastric cancer cells (19). Other previous studies propose that VP may inhibit cancer cell growth in the absence of photocactivity by inhibiting the YAP/TAZ transcriptional enhanced associate domain (YAP/TAZ-TEAD) complex (20-22). In the present study, the aim was to investigate the effects of VP on two gastric cancer cell lines: MKN-45 (TAZ-dominant) and MKN-74 (YAP-dominant).

Materials and methods

Clinical samples and datasets. The gene expression patterns of YAP and TAZ in cancer and normal tissue were compared on the Gene Expression Profiling Interactive Analysis (GEPIA) server developed by the Zhang Lab at Peking University (http://geopia.cancer-pku.cn/). The survival analysis of YAP and TAZ was performed on the Kaplan-Meier Plotter online tool headquartered at the Semmelweis University in Budapest (http://kmplot.com/analysis/index.php?p=service&cancer=gastriuc). Tissue images are obtained from the Human Protein Atlas (https://www.proteinatlas.org/).

Reagents and antibodies. VP (cat. no. SML0534) was purchased from Sigma-Aldrich; Merck KGaA. The following primary antisera were used for the immunoblotting analysis: Anti-β-actin antibody (cat. no. 13E5; Cell Signaling Technology, Inc.), anti-YAP/TAZ antibody (cat. no. D24E4; Cell Signaling Technology, Inc.), anti-Survivin antibody (cat. no. EP2880Y; Abcam), anti-CTGF antibody (cat. no. L-20; Santa Cruz Biotechnology, Inc.), and anti-CYR61/CCN1 antibody (cat. no. ab24448; Abcam). The following primary monoclonal antibodies were used for immunofluorescence: Anti-YAP antibody (cat. no. sc-101199; Santa Cruz Biotechnology, Inc.), anti-CD31 antibody (cat. no. ab28364; Abcam), Horseradish peroxidase (HRP)-conjugated goat anti-mouse IgG (cat. no. ab7890; Abcam) and goat anti-rabbit IgG H&L (Alexa Fluor® 488; cat. no. ab150113; Abcam) were purchased for western blotting. Goat anti-mouse IgG H&L (Alexa Fluor® 488; cat. no. ab150113; Abcam) was purchased for immunofluorescence.

Cell lines and cultures. Kato III (derived from a signet ring cell carcinoma with no further information), NUGC-4 (derived from an adenocarcinoma, poorly differentiated, signet ring cell carcinoma with lymph node metastasis in a 32-year-old female), MKN-45-Luc (derived from a poorly differentiated adenocarcinoma in a 62-year-old female), MKN-74/CMV-Luc cells (derived from a moderately differentiated tubular adenocarcinoma with liver metastasis in a 37-year-old male), and PANCl-1 (TAZ-dominant pancreatic cancer cell line) were obtained from the JCRB Cell Bank. The Kato III cell line was grown without antibiotics in Roswell Park Memorial Institute (RPMI)-1640 (FUJIFILM Wako Pure Chemical Corporation) medium with Eagle’s minimal essential medium (E-MEM; FUJIFILM Wako Pure Chemical Corporation), supplemented with 10% fetal bovine serum (FBS; Biosera) and 1% L-glutamine solution (FUJIFILM Wako Pure Chemical Corporation). The other cell lines were cultured without antibiotics in RPMI-1640 medium, supplemented with 10% fetal bovine serum and 1% L-glutamine solution. The cells were cultured in a humidified incubator with 5% CO₂ at 37°C. Dimethyl sulfoxide (DMSO) was used as the solvent for VP, and all the negative control groups were treated with a DMSO volume the same as the highest concentration of VP solution. Aluminum foil was used to protect cells treated with VP from light exposure, and all experiments with samples containing VP were performed in the dark.

RNA interference and transfection. YAP and TAZ were transiently knocked down in the MKN-74 and MKN-45 cell lines with validated siRNAs (L-012200-00-0005 ON-TARGETplus Human YAP1 (cat. no. 10413) siRNA-SMARTpool, L-016083-00-0005 ON-TARGETplus Human WW domain containing transcription regulator 1 (WWTR1; also known as TAZ; cat. no. 25937) siRNA-SMARTpool) from GE Healthcare Dharmacon (Horizon Discovery Group). Universal non-targeting control pool of four siRNAs (D-001810-10-05 ON-TARGETplus Non-targeting Pool) from GE Healthcare Dharmacon (Horizon Discovery Group) were used for the negative control group (NT). All the siRNA sequences used in the present study are listed in Table II. The cells were grown on 96-well or 6-well plates and transfected (final concentration, 60 nM) with Lipofectamine RNAiMAX reagent per the manufacturer’s instructions (Thermo Fisher Scientific, Inc.), incubated at 37°C for 96 h, and used for MTS assays, RT-qPCR and caspase-3/7 assays. The efficiency and specificity of the knockdown were assayed by western blotting.

Cell viability assay. VP-induced changes in gastric cancer cell viability were assessed using an MTS assay. MKN-45 and MKN-74 cells were seeded into 96-well plates at a density of 5x10³ cells/ml (200 µl/well) and incubated at 37°C for 24 h. Cells were then treated with 10, 15 and 20 µM of VP and incubated at 37°C for 24, 48 and 72 h, respectively. Cell viability was measured by MTS (3-(4,5-dimethylthiazol-2-yl)-5-[3-carboxymethoxyphenyl]-2-(4-sulfophenyl)-2H-tetrazolium) assay as follows: 100 µl of the culture medium and 20 µl of the proliferation assay solution of a CellTiter 96® AQueous One Solution Cell Proliferation assay (cat. no. G3580; Promega Corporation) were combined and incubated at 37°C for 1 h. Absorbance was then measured at 490 nm using a microplate reader (Viento nano; Sumitomo Dainippon Pharma Co., Ltd.) with an All-in-One Microplate reader Software (Gen 5 ver 2.0.0.18; BioTek), and the viability of the sample relative to control cells was calculated.

Western blot analysis. MKN-45 and MKN-74 cells were treated with 15 µM of VP and incubated at 37°C for 24 h. Whole-cell lysates were collected by adding ice-cold RIPA Lysis and Extraction Buffer (cat. no. 89900; Thermo Fisher Scientific, Inc.) containing Complete™ ULTRA Tablets, EASYpack Protease Inhibitor Cocktail and PhoSTOP (cat. nos. 05892970001 and 4906845001; Roche Diagnostics). Cells were collected by scraping. Samples were vortexed and lysed on ice for ~20 min. The lysed cells were centrifuged at 12,000 x g for 15 min at 4°C to remove cellular debris. Supernatants were transferred to clean tubes, and protein concentrations were determined by a Pierce 660 nm Protein Assay Reagent (cat. no. 1861426; Thermo Fisher Scientific,
Table I. Primer sequences used for reverse transcription-quantitative PCR.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Primer sequences</th>
</tr>
</thead>
<tbody>
<tr>
<td>Survivin</td>
<td>F: 5'-CAAGGACCACCCGCTCCTACTC-3'&lt;br&gt;R: 5'-AGTCTGGCTGTCTCAGTTG-3'</td>
</tr>
<tr>
<td>CTGF</td>
<td>F: 5'-CAAGTGTCTGACTTCGACAACGC-3'&lt;br&gt;R: 5'-GCCATGGCGCTGTCTTGAGTA-3'</td>
</tr>
<tr>
<td>CYR61</td>
<td>F: 5'-GAGTGGGTCTGTGACGAGGAT-3'&lt;br&gt;R: 5'-GTTTGTATAGATGCGAGGCT-3'</td>
</tr>
<tr>
<td>β-actin</td>
<td>F: 5'-GCATCTCACTCCCTGAATGTA-3'&lt;br&gt;R: 5'-TGTGTGAGTTTCTCC-3'</td>
</tr>
</tbody>
</table>

CTGF, connective tissue growth factor; CYR61, Cysteine-rich angiogenic inducer 61; F, forward; R, reverse.

Table II. Sequences of siRNA oligonucleotides.

<table>
<thead>
<tr>
<th>Name</th>
<th>Target sequences</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non-targeting</td>
<td>5'-UGGUGUUACUGUCGACUA-3'</td>
</tr>
<tr>
<td>Pool (NT)</td>
<td>5'-UGGUGUUACUGUUGUGA-3'</td>
</tr>
<tr>
<td>Human YAP1</td>
<td>5'-GCACCAUAUCUCUGAGA-3'</td>
</tr>
<tr>
<td>siRNA</td>
<td>5'-UGGAGAAACAUGACGACCA-3'</td>
</tr>
<tr>
<td>SMART pool</td>
<td>5'-GGUGUACAUGAUCUCAA-3'</td>
</tr>
<tr>
<td>(si-YAP)</td>
<td>5'-CCCAAGACUAGAUAAGA-3'</td>
</tr>
<tr>
<td>Human WWTR1</td>
<td>5'-CAGGAGGCGUCAGAUA-3'</td>
</tr>
<tr>
<td>siRNA</td>
<td>5'-GGGACAAACACCCCAAGA-3'</td>
</tr>
<tr>
<td>SMART pool</td>
<td>5'-AGGAACAAAGUGUCAUU-3'</td>
</tr>
<tr>
<td>(si-TAZ)</td>
<td>5'-CCAAACUCUGAAGAAUA-3'</td>
</tr>
</tbody>
</table>

YAP, Yes-associated protein; WWTR1, WW domain-containing transcription regulator protein 1 (alias of TAZ); TAZ, transcriptional co-activator with a PDZ-binding domain; si, small interfering.

Inc.). Proteins (30-50 μg/lane) were resolved by 4-20% SDS-PAGE and transferred to nitrocellulose membranes for western blotting. Membranes were incubated with primary antibodies at 4°C overnight in 5% BSA-TBS. Primary antibody dilutions were 1:1,000 unless otherwise indicated. Membranes were then washed for 30 min in TBS-0.05% Tween. Membranes were incubated with secondary antibodies (1:5,000) for 1 h at room temperature. Immunoreactive proteins were visualized using Clarity Western ECL substrate (cat. no. 1705061; Bio-Rad Laboratories, Inc.) and an image analyzer (LAS-3000 mini; Fujifilm Co., Ltd.) with an Image Reader (LAS-3000 UV mini ver2.2; Fujifilm Co., Ltd.). When multiple proteins were evaluated, membranes were stripped using Restore PLUS Western Blot Stripping Buffer (Thermo Fisher Scientific, Inc.) before primary antibody incubation.

Crystal violet staining. VP-induced changes in gastric cancer cell viability were also assessed by crystal violet (CV; cat. no. 031-04851; FUJIFILM Wako Pure Chemical Corporation) staining. MKN-45 and MKN-74 cells were seeded onto 6-well plates at a density of 3x10^4 cells/ml and incubated at 37°C for 24 h. Cells were treated with 10 and 15 μM of VP and incubated at 37°C for 24 h. Cells were then fixed for 5 min with 4% paraformaldehyde (PFA) and stained for 30 min at room temperature with 0.05% CV. Samples were then washed twice with tap water and allowed to drain in an inverted position for ~2 min. Staining was recorded by photography, and then a volume of methanol equivalent to one-third to one-half the total well volume was added and solubilize the dye for 30 min at room temperature (20-25°C). The absorbance at 540 nm was measured with aliquots transferred to a fresh plate using a microplate reader (Viento nano; Sumitomo Dainippon Pharma Co., Ltd.) with an All-in-One Microplate reader Software (Gen 5 ver 2.0.18; BioTek).

RNA extraction and reverse transcription-quantitative (RT-q) PCR. Total RNA was extracted from cultured cells using an miRNeasy Mini kit (cat. no. 217004; Qiagen GmbH) and quantified using a Biospec-nano spectrophotometer (Shimadzu Corporation). The extracted RNA samples were stored at -80°C until use. cDNAs were prepared from total RNA using a High-Capacity cDNA Reverse Transcription kit (cat. no. 4374966; Thermo Fisher Scientific, Inc.) according to the manufacturer’s protocols. Reverse transcription reaction mixtures contained 2 μg of total RNA, 1X RT buffer, 4 mM dNTP mix, 1X RT random primer, 50 units MultiScribe reverse transcriptase and 20 units RNase inhibitor. Nuclease-free water was added to adjust the reaction volume to 20 μl. The reaction mixtures were incubated at 25°C for 10 min, followed by 37°C for 120 min and 85°C for 5 min. The real-time-PCR assays were performed in 20 μl aliquots containing 1 μl RT products with 4 μl LightCycler® FastStart DNA Master PLUS SYBR Green I (cat. no. 03515869001; Roche Diagnostics), 0.2 μl (final concentration 0.5 μM) of each primer and 14.6 μl nuclease-free water. Analyses were run on a Real-Time PCR Light Cycler® 1.5 Complete System (Roche Diagnostics). Thermal cycling was initiated with a denaturation step at 95°C for 10 min, followed by 45 cycles of 95°C for 10 sec, 60°C for 10 sec, and 72°C for 10 sec. The cycle threshold (Cq) was recorded for each target mRNA by LightCycler® Software version 3.5.28 (Roche Diagnostics), and β-actin was used as the endogenous control for data normalization. The relative expression was calculated using the formula 2^-ΔΔCq (2^-ΔCq, reagent treatment-ΔCq, control) (23). All the primer sequences used in the present study are listed in Table I.

Verteporfin uptake evaluation. An all-in-one fluorescence microscope (BZ-X800; Keyence Corporation) equipped with an OP-87767 filter (excitation: 405 nm and fluorescence: 630 nm) was used to evaluate VP uptake. MKN-45 and MKN-74 cells were seeded onto 96-well plates at a density of 5x10^4 cells/ml (200 μl/well) and incubated at 37°C for 24 h. Cells were then treated with 15 μM of VP and incubated at 37°C for 30 min, 1 or 2 h. Pixel intensities were measured using ImageJ software ver1.52a (Wayne Rasband; National Institutes of Health) using 20 random microscopic fields (magnification, x20), according to the manufacturer’s instructions (24).
**Immunofluorescence.** MKN-45 and MKN-74 cells were seeded on glass coverslips in 6-well plates at a density of 3x10^5 cells/ml (1 ml/well) and incubated at 37°C for 24 h. Cells were then treated with 15 µM of VP and incubated at 37°C for 24 h. Cells were fixed with 4% PFA and permeabilized with 0.1% Triton X-100 and then incubated in blocking buffer containing 5.0% BSA and 0.1% glycine at room temperature (20-25°C). Cells were incubated with the following primary antibodies in blocking buffer at 4°C overnight; anti-YAP antibody (cat. no. sc-101199; Santa Cruz Biotechnology, Inc.), anti-TAZ antibody (cat. no. ab84927; Abcam), and anti-CD31 antibody (cat. no. ab28364; Abcam). Cells were washed and then incubated with corresponding secondary antibody, Goat anti-mouse IgG H&L (Alexa Fluor® 488; cat. no. ab150113; Abcam) in blocking buffer for 1 h at room temperature in the dark. Cells were washed, and the coverslips were mounted onto slides using ProLong Gold Antifade reagent (Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. Coverslips were mounted with ProLong Gold Antifade reagent with DAPI (Thermo Fisher Scientific, Inc.) and fixed overnight at room temperature in the dark. Slides were analyzed with an all-in-one fluorescence microscope (BZ-X800; Keyence Corporation). Caspase-3/7-positive cells were quantified by calculating the ratio of caspase-3/7-positive to DAPI-positive nuclei (total nuclei) in 10 random microscopic fields (magnification, x20).

**Statistical analysis.** All data represent ≥ three independent experiments using cells from a minimum of three separate isolations. Skewness-Kurtosis was used to check the distribution of the data. The Mann-Whitney U test was used to assess the statistical significance of differences between two groups. Kruskal-Wallis test was used for comparisons between groups and Dunn's test was applied for multiple comparisons. Survival curves were calculated using the Kaplan-Meier method on the Kaplan-Meier Plotter online tool. Statistical analyses were performed using StatFlex software (Windows ver. 6.0; Artech LLC), except for the Log-rank test for survival analysis using the Kaplan-Meier Plotter online tool. P<0.05 was considered to indicate a statistically significant difference and all data are expressed as the mean ± SEM.

**Results**

**YAP and TAZ are prognostic factors of gastric cancer.** YAP and TAZ expression levels in gastric cancer cells are significantly higher than those in normal tissues (Fig. 1A and B). A higher expression of YAP or TAZ in gastric cancer cells is associated with a significantly poorer prognosis than cancer cells with lower expression levels (Fig. IC and D).

**Expression patterns of YAP and TAZ differ among gastric cancer cell lines.** Initially, YAP and TAZ expression levels were examined in four gastric cancer cell lines. Kato III, NUGC-4 and MKN-74 cells exhibited a YAP-dominant expression pattern. By contrast, the MKN-45 cell line showed a TAZ-dominant expression pattern (Fig. 2A). Therefore, MKN-45 was selected as the TAZ-dominant cell line and MKN-74 as the YAP-dominant cell line (Fig. S1A).

**VP suppresses MKN-45 proliferation.** The effect of VP on proliferation in MKN-45 and MKN-74 cells was examined using 10, 15 and 20 µM VP. VP treatment suppressed proliferation in MKN-45 cells (Fig. 2B) more than it does in MKN-74 cells at 24 h (Fig. 2C). In MKN-74 cells, the same level of proliferation suppression was observed at 72 h. VP treatment decreased the amount of CV staining in MKN-45 cells in a dose-dependent manner over 24 h (Fig. 2D). VP uptake was faster in MKN-45 cells than MKN-74 cells (Fig. 2E).

**VP decreases YAP and TAZ proteins after 12 h.** Western blotting demonstrated that VP treatment for 12 h reduces the expression of YAP in MKN-74 cells and TAZ in MKN-45 cells (Fig. 3A). PANC-1 cells were used as a positive control for TAZ expression (Fig. S1A).

**VP decreases Survivin gene and protein expression in MKN-45 and MKN-74 cells.** VP treatment reduces the expression of downstream Survivin gene in MKN-45 and MKN-74 cells at 24 h (Fig. 3B and C). Moreover, immunoblotting
confirmed that only Survivin expression is affected by VP treatment at 24 h in both cell lines (Fig. 3D).

**VP changes the localization of YAP and TAZ from nuclear to cytosolic.** Immunofluorescence analysis demonstrated that exposure to 15 µM VP for 2 h changes YAP and TAZ localization from the nucleus to the cytosol in MKN-45 and MKN-74 cells (Fig. 4A and B).

**VP induces cell death in MKN-45 and MKN-74 cells in a dose-dependent manner.** Annexin V staining demonstrated that treatment with 15 µM of VP for 24 h induces cell death in MKN-45 and MKN-74 cells in a dose-dependent manner (Fig. 5A).

**VP suppresses vascular mimicry.** MKN-45 and MKN-74 are CD31-positive cells that have angiogenic and vasculogenic potential (Fig. S1B). Therefore, the vascular mimicry of both cell lines was examined using tube formation assays. In MKN-45 and MKN-74 cells, exposure to 15 µM of VP for 72 h suppressed the level vascular mimicry, demonstrated by a decrease in the number of tubules formed (Fig. 5B).

**Concurrent YAP/TAZ knockdown decreases cellular proliferation.** RNA interference assays targeting YAP, TAZ and YAP/TAZ in MKN-45 and MKN-74 cells demonstrated that simultaneous knockdown (YAP/TAZ) is required to decrease cell proliferation (Fig. 6A-D). YAP and YAP/TAZ knockdown also decrease Survivin expression. Concurrent YAP/TAZ knockdown increases caspase3/7 activity in both cell lines (Fig. 6E-G).

**Discussion**

In the present study, it was demonstrated that VP affects both YAP-dominant (MKN-74) and TAZ-dominant (MKN-45) cell...
lines. Moreover, in the poorly differentiated, TAZ-dominant gastric cancer cell line MKN-45, VP treatment elicited a more rapid suppression than in MKN-74 cells.

VP is used in PDT as a photosensitizer to eliminate abnormal blood vessels in the eyes of patients with conditions such as macular degeneration (18). It produces highly reactive, short-lived, singlet oxygen species when stimulated by non-thermal red light at 689 nm. VP has also been employed for PDT in oncology. A Phase I/II clinical trial of PDT with VP was conducted in patients with locally advanced pancreatic cancer in 2014 (25). However, light penetration into abdominal tissue is an invasive process. Therefore, the present study focused on gastric cancers because these tumors are more accessible to PDT via endoscopy. Our previous study reported the efficacy of VP in PDT using a gastric cancer cell line (19). More recently, VP has been...
reported to have an anti-cancer effect via YAP inhibition in the absence of light activation (20-22,26,27). Moreover, VP inhibited interactions between YAP/TAZ and the TEA domain transcriptional factor (TEAD), and inhibited YAP function by upregulating the 14-3-3σ sequestering of YAP in the cytoplasm (28).

YAP and TAZ gene expression levels are elevated in a subset of human gastric cancers, most of which are associated with poor clinical outcomes (29-33). Choi et al (34) recently demonstrated that MYC upregulation is a direct consequence of YAP activation in YAP-activated human gastric cancer cells.

YAP and TAZ are usually discussed in the same vein. However, in renal development, YAP-knockout (KO) mice are embryonically lethal (35), whereas TAZ-knockout mice develop severe cystic kidney disease (36). A study using
YAP-KO, TAZ-KO and YAP-/TAZ-KO cell lines generated by the CRISPR/Cas9 technique in 293A cells revealed that YAP inactivation has more significant effects on cellular physiology than TAZ inactivation; namely, cell spreading, volume, granularity, glucose uptake, proliferation and migration (37). TAZ expression is highly elevated in gastric signet ring cell carcinoma (17). Hayashi et al (38) reported an imbalance in TAZ and YAP expression in gastrointestinal cancer cell lines. Another study reported that TAZ accumulation is negatively regulated by YAP abundance (39). The present results suggest that concurrent suppression of YAP and TAZ inhibits cell proliferation and that TAZ and YAP may serve distinct roles in cancer progression.

In the present study, four gastric cancer cell lines were used; one was TAZ-dominant and the other three were YAP-dominant. Kato III and NUGC-4 are signet ring carcinoma cell lines; however, YAP/TAZ dominance patterns do not depend on histological classification. VP decreases YAP and TAZ expression and subsequently reduces the expression of downstream genes, predominantly Survivin. Survivin is a new member of the inhibitor of apoptosis (IAP) family and is selectively upregulated in most human cancers but not in normal tissues (40). Moreover, it exhibits anti-apoptotic activity via caspase-3 and caspase-7 inhibition (41) and is a prognostic factor in gastric cancer associated with lymphatic metastasis (42-44). A correlation between YAP and Survivin in gastric cancer has been demonstrated for tumorigenesis and lymph node metastasis (45). The current results indicate that VP inhibits YAP/TAZ and induces apoptosis by increasing caspase-3 and caspase-7 activity via decreasing Survivin expression in gastric cancer cells.

Vascular mimicry (VM) is a concept proposed by Maniotis et al (46) in 1999. Tumors with vigorous VM are often aggressive and associated with a poor prognosis (47). Evaluating the therapeutic potential of inhibiting VM has critical clinical importance. Sun et al (48) demonstrated that VM is correlated with the differentiation, stage and metastatic potential of tumors, in addition to less favorable prognoses in clinical samples of gastric cancer. VP has been shown to suppress VM by disrupting the YAP-TEAD complex (49,50). CD31-positive cancer cells are reported to display VM (51), and the cell lines used in the present study were CD31-positive (Fig. S1B). Therefore, VM was evaluated in VP-treated cells using tube formation assays and it was revealed that VP significantly suppresses VM in MKN-45 and MKN-74 cells. Consequently, VP may represent a promising target for anti-cancer therapy in YAP-dominant or TAZ-dominant gastric cancers with poor prognostic outcomes.

Kang et al (52) demonstrated that the anti-proliferative effects of VP on gastric cancer cell lines are associated with suppression of FAT1, another marker associated with poor prognoses. Notably, more recent studies report that VP can potentially treat
chemo-resistant gastric cancer stem cells (53,54). However, these studies did not distinguish between YAP and TAZ and did not evaluate the effects of VP on Survivin expression and vascular mimicry. In the present study, characteristics that reveal underlying mechanisms of VP in the suppression of gastric tumor cell growth were investigated.

One limitation of the present study was the high concentration of VP required to achieve suppression. The plasma concentration in human patients receiving liposomized VP (Visudyne®) is ~2 µg/ml. The in vitro assays used 10 and 15 µM VP treatments, which equate to 5 and 7.5 µg/ml, respectively, or 2-4 times more than a recommended clinical dose. Liposomization can increase VP tissue uptake, leading to a high concentration in the gastric cancer tissue.

In conclusion, VP has the potential to suppress different types of gastric cancers via suppressing Survivin. Further
in vivo studies will be required to confirm the potential of VP for use as a therapeutic agent.

Acknowledgements

Not applicable.

Funding

The resources and facilities of the Faculty of Medicine, Tottori University were used to conduct this study. This research did not receive any specific grant from funding agencies in the public, commercial or not-for-profit sectors.

Availability of data and materials

The datasets used and analyzed in this study are available from the corresponding author upon reasonable request.

Authors’ contributions

TH, TS, YH, RT, YM, and TK designed and implemented the experiments. TS, TT, TN, and TM searched the literature and analyzed and interpreted the data. TS, TT, TN, and TM were also involved in writing, reviewing, and editing the manuscript. HI made substantial contributions to the conception and design of the project and gave final approval of the
version to be published. TH and TS confirmed the authenticity of all the raw data. All authors read and approved the final manuscript.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References


