Abstract. Radiation therapy is one of the most promising therapeutic strategies in unresectable esophageal squamous cell carcinoma (ESCC). The histone deacetylase (HDAC) inhibitor has been shown to enhance radiosensitivity. Valproic acid (VPA) is a well-known drug used to treat seizure disorders and epilepsy, and has been shown to inhibit HDACs. We recently reported that a clinically safe dose of VPA enhances radiation-induced cytotoxicity in human ESCC cells. However, the mechanism of radiosensitizing effect of VPA has not yet been confirmed. The present study examined the effect of VPA on DNA double-strand break (DSB) repair after radiation in the human ESCC cell lines KES, TE9 and TE11 by examining H2AX phosphorylation (γH2AX) levels as a marker of radiation-induced DSBs. The present study also examined whether VPA inhibited radiation-induced DNA DSB repair by suppressing non-homologous end joining (NHEJ), focusing particularly on the acetylation of Ku70. VPA was shown to prolong γH2AX levels after irradiation in all three ESCC cell lines. Moreover, prolonged γH2AX foci formation after irradiation was also observed by immunocytochemistry following VPA pretreatment in KES and TE9 cells. VPA was shown to induce Ku70 acetylation after irradiation in all three ESCC cell lines. Our results suggest that VPA prolonged radiation-induced DSBs by inhibiting NHEJ in DSB repair pathways in ESCC. VPA could therefore be used as an effective radiosensitizer in ESCC radiotherapy.

Introduction

Esophageal squamous cell carcinoma (ESCC) is a malignant disease with a poor prognosis. The standard treatment for advanced ESCC patients has been surgical resection, although patients with unresectable ESCC are usually treated by definitive chemoradiotherapy (CRT) (1). Various anticancer agents have been used to enhance the therapeutic effects of radiotherapy, including cisplatin and 5-fluorouracil (2,5,6). Although favorable outcomes after CRT compared with radiation therapy alone have been reported (1,7), adverse events are also noted, such as pulmonary dysfunction, leukocytopenia, radiation pneumonitis, pericardial effusion, pleural effusion, perforation and stenosis (8,9). Thus, there is a need to both enhance therapeutic effects and reduce radiation-induced adverse events when performing CRT for esophageal cancer.

Histone acetyltransferases (HATs) and histone deacetylases (HDACs) regulate the acetylation and deacetylation of histones, respectively. Core histone acetylation relaxes the chromatin structure and facilitates transcription (10,11). HDACs are involved in the deacetylation of chromatin histone proteins as well as non-histone proteins, which regulate cell differentiation, apoptosis and growth arrest (12,13). HDAC inhibitors act additively or synergistically with conventional cancer therapies such as radiotherapy, and numerous studies have shown that they enhance the radiation sensitivity of various cancer cell lines both in vitro and in vivo (14-23). These effects have been explained as the modulation of cell cycle regulation, particularly G1-phase arrest, the inhibition of DNA synthesis, the suppression of DNA repair pathways, apoptosis and the alteration of chromatin structures by histone hyperacetylation (16,19,20,24).
Much attention has been given to the suppression of the double-strand break (DSB) repair pathway by HDAC inhibitors, which enhance the radiation sensitivity and accumulation of the sensitive marker of DSBs, H2AX phosphorylation (γH2AX) (19). Homologous recombination (HR) and non-homologous end joining (NHEJ) are the main pathways by which radiation-induced DSBs are repaired (25-27). Rad51 is a key protein of HR, while NHEJ involves the Ku70/Ku86 heterodimer which binds with strong affinity to broken DNA ends, and recruits the catalytic subunit of DNA-dependent protein kinases (DNA-PKcs). Ku70 functions are regulated by HATs and HDACs, and an increase in Ku70 expression in cancer cells was found to enhance the DNA DSB repair ability and reduce Bax-mediated apoptosis (28). Ku70 is also a target of some class I, II and III HDACs (29), while the use of HDAC inhibitors led to increased Ku70 acetylation accompanied by a reduced DNA binding affinity, with no disruption to Ku70/Ku80 heterodimer formation (30). Treatment of a myelogenous leukemia cell line with HDAC inhibitors before irradiation resulted in increased and prolonged expression of γH2AX repair foci (24).

Valproic acid (VPA), an 8-carbon branched-chain fatty acid, has a well-established efficacy for the treatment of epilepsy and other seizure disorders. VPA was also shown to be an effective inhibitor of HDACs (31,32), and to enhance radiosensitivity in various cancer cell lines (18,33-35). We recently reported a clinically safe dose of VPA that can enhance radiation-induced cytotoxicity in human ESCC cells by chromatin decondensation with histone hyperacetylation and the downregulation of Rad51 (36). For certain HDAC inhibitors, the acetylation of proteins in NHEJ has been demonstrated to be involved in the inhibition of radiation-induced DSBs repair (37). However, the mechanism underlying the inhibitory effect of VPA with respect to NHEJ in esophageal cancer has not yet been confirmed. The present study therefore examined whether VPA inhibits radiation-induced DNA DSB repair by suppressing NHEJ in human ESCC cells, focusing particularly on the acetylation of Ku70 by VPA.

Materials and methods

Cell lines and cell culture. Three human ESCC cell lines were used in the present study. The poorly differentiated ESCC cell line TE9, and the moderately differentiated TE11 cells were provided by Dr Tetsuro Nishihira (Higashimatsuyama Medical Association Hospital, Saitama, Japan). The highly differentiated ESCC cell line KES was established in our laboratory from endoscopic biopsy specimens obtained from a patient carrying highly differentiated ESCC. These cells were seeded in 75-cm² dishes (Becton-Dickinson, Tokyo, Japan) and cultured in 10 ml of medium consisted of RPMI-1640 (Gibco®, Life Technologies Corp., Tokyo, Japan) supplemented with 10% heat-inactivated fetal bovine serum (Nichirei Biosciences, Inc., Tokyo, Japan), 100 IU/ml penicillin and 100 μg/ml streptomycin (Life Technologies Corp.) at 37°C in a humidified atmosphere of 5% CO₂. Cells were grown to confluency and harvested by trypsinization with 0.25 mg/ml trypsin/EDTA (Life Technologies Corp.), then resuspended in culture medium before use.

Reagents and antibodies. VPA was purchased from Sigma-Aldrich Co. (Tokyo, Japan). Rabbit anti-phospho-histone H2AX (Ser 139) (γH2AX) (cat. #9718) and anti-acetylated-lysine monoclonal antibodies (cat. #9814) were obtained from Cell Signaling Technology Co. Ltd. (Tokyo, Japan). Rabbit monoclonal anti-Ku70 antibody was purchased from Abcam (cat.#ab92450; Cambridge, UK), and goat anti-rabbit IgG horseradish peroxidase (HRP)-conjugated were purchased from Cell Signaling Technology (cat. #7074). β-actin and TATA-binding protein (TBP) were detected as control proteins using rabbit polyclonal anti-β-actin antibody (cat. #4967) and rabbit polyclonal anti-TBP antibody (cat. #8515) (both from Cell Signaling Technology), respectively.

Irradiation. Cells were irradiated by MBR-1520R-3 (Hitachi Medicotechnology, Hitachi, Japan) at a dose rate of 1 Gy/min. The X ray-irradiation power output was 125 kV, 20 mA. Forward-scattered radiation was filtered using a 0.5 mm Al and 0.2 mm Cu filter.

Immunofluorescent cytochemistry. Cells were cultured on Lab-Tec chamber slides (Nalge Nunc International, Rochester, NY, USA), treated with or without VPA (0.5 mM) for 24 h, then irradiated (6 Gy). Treated cells were fixed in methanol and acetone (1:1) for 1 min after irradiation at designated time-points (0/4/8 h). After washing in phosphate-buffered saline (PBS), the slides were immersed in methanol containing 0.3% H₂O₂ for 10 min, blocked with 3.3% normal goat serum in PBS and 0.2 mm Cu filter. Nuclei were stained with Hoechst 33258 for 5 min, and slides were observed under an immunofluorescence microscope (Olympus BX50/BX-FLA; Tokyo, Japan).

Quantification of γH2AX levels after irradiation. To assess changes in radiation-induced DNA DSB repair, we compared γH2AX levels in irradiated cells treated with or without VPA (0.5 mM) using western blotting. Cells treated with combination therapy were pre-treated with VPA for 24 h before irradiation. Cell lysates were collected after irradiation at designated times (0/10/30 min/1/2/4/8 h). The protein concentration in each lysate was measured using the Pierce BCA protein assay kit (Thermo Fisher Scientific, Rockford, IL, USA). A sample volume equivalent to 20 μg protein from each lysate was loaded on a 10% polyacrylamide gel (Mini-Protean TGX Precast Gels; Bio-Rad Laboratories Inc., Berkeley, CA, USA) and separated by electrophoresis. Proteins were transferred to polyvinylidene difluoride (PVDF) membranes (Trans-Blot Turbo mini-PVDF Transfer Pack; Bio-Rad Laboratories Inc.), blocked with Tris-buffered saline with Tween-20 Tablets, pH 7.6 (Takara Bio, Inc., Shiga, Japan), then incubated with the rabbit monoclonal anti-phospho-histone H2AX (Ser 139) antibody (1:200) at 4°C overnight. Following three washes in PBS, slides were incubated with an anti-rabbit IgG antibody conjugated with Alexa Fluor® 594 (cat. #A11012; Molecular Probes/Invitrogen, Carlsbad, CA, USA) for 1 h at room temperature in the dark. Nuclei were stained with β-actin antibody (cat. #4967) and rabbit polyclonal anti-TBP antibody (cat. #8515) (both from Cell Signaling Technology), respectively.

β-actin was detected as a control protein. Secondary antibody incubation using goat α-rabbit IgG antibody conjugated with Alexa Fluor® 594 (cat. #A11012; Molecular Probes/Invitrogen, Carlsbad, CA, USA) for 1 h at room temperature in the dark. Nuclei were stained with Hoechst 33258 for 5 min, and slides were observed under an immunofluorescence microscope (Olympus BX50/BX-FLA; Tokyo, Japan).

The X-ray irradiation power output was 125 kV, 20 mA. Forward-scattered radiation was filtered using a 0.5 mm Al and 0.2 mm Cu filter.
anti-rabbit IgG HRP-conjugated at a 1:4,000 dilution was carried out according to the manufacturer's instructions for 10 min at room temperature (cat. SNAP2BASE; SNAP i.d. 2.0; Millipore, Billerica, MA, USA). Membranes were treated with PVDF Blocking Reagent for Can Get Signal® (Toyobo Co., Ltd., Osaka, Japan). The antibody-antigen complex was detected using enhanced chemiluminescence by an ECL western blotting detection kit (General Electric Healthcare Japan Co., Ltd., Tokyo, Japan) and the Light-Capture system (ATTO Co., Ltd., Tokyo, Japan). Quantification was performed by the CS analyzer program (ATTO Co., Ltd.) and normalized to β-actin levels.

Assessment of Ku70 acetylation. Cells were incubated with or without VPA (0.5 mM) for 24 h, then irradiated (6 Gy) and harvested promptly. Nuclear proteins were extracted using a Nuclear Extract kit (Active Motif, Tokyo, Japan), and protein concentrations were determined using a Pierce BCA protein assay kit as before. A rabbit monoclonal anti-Ku70 antibody at 1:80 dilution was used for immunoprecipitation.
Magnetic beads (Dynabeads® Protein G Immunoprecipitation kit; Life Technologies, Carlsbad, CA, USA) were used for immunoprecipitation of the target antigen according to the manufacturer’s instructions. The target antibody was incubated with Dynabeads® with rotation for 10 min to enable binding to occur. After adding the extracted nuclear protein to the beads-antibody complex, the sample was incubated at 4°C overnight with rotation to form an antigen-beads-antibody complex. The target antigen was dissociated from the beads-antibody complex with elution buffer. Ku70 acetylation was then examined by western blotting using a rabbit monoclonal anti-acetylated-lysine primary antibody at a 1:300 dilution. To detect TBP as a control nuclear protein, a rabbit polyclonal anti-TBP antibody (Cell Signaling Technology) at 1:300 dilution was used. Goat anti-rabbit IgG-HRP conjugates at 1:4,000 dilution were used as secondary antibodies.

Results
Changes of radiation-induced $\gamma$H2AX levels by VPA. $\gamma$H2AX levels in cells pre-treated with VPA were higher than in untreated cells in all three ESCC cell lines (Fig. 1). $\gamma$H2AX levels were detected by western blotting and quantified by measuring the ratio of $\gamma$H2AX to $\beta$-actin. The time course of $\gamma$H2AX is displayed in Fig. 2. VPA was shown to prolong $\gamma$H2AX levels after irradiation in all three ESCC cell lines. Moreover, prolonged $\gamma$H2AX foci formation after irradiation was also observed by immunocytochemistry following VPA pretreatment in KES and TE9 cells (Fig. 3).

Acetylation of Ku70 by VPA. The acetylation of Ku70 by VPA was estimated by western blotting following immunoprecipitation of the nuclear extract with an anti-Ku70 antibody and the detection of acetylated-lysine. In all three ESCC cell lines, increased Ku70 acetylation after irradiation was observed following pretreatment with 0.5 mM VPA for 24 h (Fig. 4).

Discussion
In the present study, we showed that VPA inhibited the radiation-induced DNA DSB repair in ESCC cell lines, and that this was accompanied by acetylation of the DSB repair protein, Ku70.

Previously, $\gamma$H2AX has been described as a highly sensitive method to detect irradiation-induced DSBs (38,39). In response to this, serine 139 within the conserved C-terminal region of the H2AX protein becomes phosphorylated by one of the DNA damage signaling kinases of the phosphatidylinositol-3-OH-
Figure 3. Changes over time in the H2AX phosphorylation foci formation after irradiation, as assessed by immunofluorescent cytochemistry. Micrographs of γH2AX foci (red) in nuclei (blue) of KES and TE9 cells pre-treated with or without VPA (0.5 mM) for 24 h were obtained (0 h/4 h/8 h) after irradiation (IR). VPA was shown to prolong γH2AX foci formation after irradiation. Scale bar, 10 µm. VPA, valproic acid.
kinase-like family. This phosphorylation rapidly spreads to a region of chromatin surrounding the DSBs, resulting in the formation of γH2AX foci. The main role of H2AX phosphorylation in the DNA damage response was reported to be the recruitment of DNA repair proteins to DSB sites and the preparation of a platform for signaling and repair (38,40). In the present study, we analyzed H2AX phosphorylation levels by western blotting, and examined γH2AX foci by immunocytochemistry as a marker of DNA DSB after irradiation.

More than 18 mammalian HDAC enzymes (HDAC 1-11 and sirtuins 1-7) have been identified, which are grouped into four classes according to their homology (41). HDAC inhibitors are an emerging class of drugs that have shown promise as anticancer agents when used alone or in combination with conventional therapies (42,43). VPA has been used for the treatment of seizure disorders and epilepsy. Previous studies found that it induces apoptosis and the accumulation of hyperacetylated histones H3 and H4, and also inhibits class I HDACs (HDACs 1, 2 and 3) and class II subclass I HDACs (HDACs 4, 5 and 7) (31,32,44,45). Therapeutic plasma concentrations used clinically for epilepsy treatment range from 50-100 µg/ml, which is equivalent to 0.3-0.6 mM. Therefore, 0.5 mM VPA is a clinically safe concentration (46) that we used in the present study based on our previous growth inhibition assay in which 0.5 mM VPA did not inhibit the growth of ESCC cells (36).

We recently showed that VPA enhanced the radiosensitizing effect by chromatin decondensation with histone hyperacetylation and downregulation of Rad51 (36). Various mechanisms are responsible for this VPA-mediated enhancement in radiosensitivity. Firstly, HDAC inhibitors lead to the acetylation of histone tails, which induces reduction of the electrostatic charge interactions between DNA and histones, permitting access to the chromatin structure in association with chromatin decondensation (47,48). Additionally, Harikrishnan et al (49) reported that euchromatin was more sensitive to radiation-induced DSBs than heterochromatin, and that histone modifications contributed to the radiosensitivity effects of HDAC inhibitors. Thus, they concluded that heterochromatin is more resistant to histone modification and DNA damage. Secondly, HDAC inhibitors inhibit the radiation-induced DNA DSB repair process. In the present study, we confirmed that VPA prolonged radiation-induced DNA DSB repair by measuring γH2AX level changes overtime after irradiation. This lengthening of DNA DSBs can be attributed to the VPA-induced inhibition of the DSB repair process. While we previously found that Rad51 expression was downregulated by VPA, and Adimoolam et al (37) reported that suberoylanilide hydroxamic acid (PCI-24781) also suppressed Rad51 expression, the mechanism of its suppression by HDAC inhibitors has not been fully clarified. Further studies are therefore necessary to understand this.

The functional inhibition of non-histone proteins by HDAC inhibitors has recently been reported to involve acetylation (30). However, it is difficult to examine the acetylation of non-histone proteins by western blotting due to the lack of an appropriate primary antibody. Recent studies reported that the acetylation of non-histone proteins by HDAC inhibitors induced the functional inhibition of DNA repair activity (20,30). Furthermore, Chen et al (30) found that HDAC inhibitors reduced the DNA end-binding affinity of Ku70 by acetylation. In the present study, we focused on Ku70 as a key protein of NHEJ in the radiation-induced DNA DSB repair pathway. We confirmed that VPA increased Ku70 acetylation levels in three ESCC cell lines.

In DNA DSB repair pathways, ataxia telangiectasia mutated and ataxia telangiectasia related proteins are important components of the DSB signaling cascade. Their activation results in the phosphorylation of downstream substrates, including p53 (50). HDACs can regulate the expression of various genes by direct interaction (13). For instance, p53 is downregulated by HDACs 1, 2 and 3 (35,51) and acetylated by VPA (45), which induces cell cycle regulation, apoptosis and inhibition of DNA DSB repair (45,50). Several proteins inhibit DNA DSBs repair pathways by mediating the resection process of DNA ends (52,53). Thus, VPA inhibits HR in the repair pathway by acetylating Sae2 and Exo1, which are two nucleases involved in resecting DNA ends (54). Our present findings and those of earlier studies showed that VPA has the potential to synergistically inhibit DNA DSB repair pathways by acetylating p53 in the DSB signaling cascade, nucleases involved in DNA end resection and Ku70 in NHEJ, as well as suppressing Rad51 expression levels in HR. VPA may
therefore be a suitable agent to combine with radiotherapy due to its multiple effects in DNA DSB repair inhibition. Further studies are needed to clarify the precise mechanisms of DNA DSB repair inhibition by VPA in ESCC.

Tomiita et al observed a difference in the proportion of HR and NHEJ in DSB repair in chicken B lymphoid cell lines according to the irradiation dose rate. Although both HR and NHEJ were activated following high-dose irradiation (0.9 Gy/min), the activation of NHEJ dominated after low-dose irradiation (1.0 mGy/min) (55). This suggests that the primary inhibition of NHEJ may be a more effective means of enhancing radiosensitivity, regardless of the radiation dose. Moreover, Karagiannis and El-Osta (50) reported a protection effect associated with HDAC inhibitors. These not only act as radiosensitizers in response to irradiation, but also reduce radiation-induced dermatitis and esophagitis by suppressing the expression of transforming growth factor-β and tumor necrosis factor-α. Therefore, the concomitant use of VPA with radiotherapy against esophageal cancer has the potential to reduce various radiation-induced adverse effects.

In conclusion, VPA treatment at clinically safe concentrations enhances radiosensitivity by inhibiting both HR and NHEJ in the DNA DSB repair process, and prolonging DNA DSBs. Therefore, VPA has considerable potential as a therapeutic agent for esophageal cancer.

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References


