HSV-1 ICP27 induces apoptosis by promoting Bax translocation to mitochondria through interacting with 14-3-3

Ji Ae Kim*, Jin Chul Kim †,*, Jung Sun Min, Inho Kang, Jeongho Oh & Jeong Keun Ahn*
Department of Microbiology & Molecular Biology, College of Biological Science and Biotechnology, Chungnam National University, Daejeon 34134, Korea

The subcellular localization of Bax plays a crucial role during apoptosis. In response to apoptotic stimuli, Bax translocates from the cytoplasm to the mitochondria, where it promotes the release of cytochrome c to the cytoplasm. In cells infected with HSV-1, apoptosis is triggered or blocked by diverse mechanisms. In this study, we demonstrate how HSV-1 ICP27 induces apoptosis and modulates mitochondrial membrane potential in HEK 293T cells. We found that ICP27 interacts with 14-3-3 which sequesters Bax to the cytoplasm. In addition, ICP27 promotes the translocation of Bax to the mitochondria by inhibiting the interaction between 14-3-3 and Bax. Our findings may provide a novel apoptotic regulatory pathway induced by ICP27 during HSV-1 infection.

INTRODUCTION

In mammals, apoptosis plays critical roles during normal development, maintenance of homeostasis, aging, and defense (1). Apoptosis is triggered and regulated by diverse stimuli and conditions. The extrinsic signaling pathway of apoptosis is induced by ligation of transmembrane death receptors and death ligands (2, 3), whereas the intrinsic signaling pathway is triggered by non-receptor-mediated stimuli such as oxidative stress, DNA damage, and ER stress. All these stimuli disrupt the mitochondrial membrane integrity, and consequently release cytochrome c into the cytoplasm (4). Bcl-2 family proteins regulate the apoptotic mitochondrial events. Bcl-2 family members are comprised of anti-apoptotic members (Bcl-2, Bcl-w, and Bcl-xL), multidomain pro-apoptotic members (Bax and Bak), and BH3 domain-only pro-apoptotic members (Bad, Bim, Bik, Noxa, and PUMA) (5, 6). A pro-apoptotic Bcl-2 family member, Bax, has an essential role in inducing apoptosis in response to stress stimuli (7-9). Bax is mainly localized in the cytoplasm by the 14-3-3 proteins (11).

Herpes simplex virus type 1 (HSV-1) is a neurotropic virus, replicating lytically during acute infection and establishing latency in peripheral neurons. During lytic infection, the HSV-1 genes are expressed in a tightly regulated temporal cascade, involving the sequential expression of immediate early (IE), early (E), and late (L). IE proteins include ICP0, ICP4, ICP22, ICP27, and ICP47, and are implicated in regulating the viral early gene expression and host cellular proteins (12, 13). ICP27 is a multifunctional protein essential for viral replication, and plays pivotal roles in the switch from early to late gene expression. ICP27 is also implicated in regulating the viral early gene expression and host cellular proteins (12, 13). ICP27 is a multifunctional protein essential for viral replication, and plays pivotal roles in the switch from early to late gene expression. ICP27 is also implicated in regulating the viral early gene expression and host cellular proteins (12, 13).

This study confirms that ICP27 induces apoptosis in 293T cells. Furthermore, we demonstrate that ICP27 promotes the mitochondrial translocation of Bax, via inhibition of the cytoplasmic sequestration of Bax by 14-3-3. These findings imply that ICP27 plays a critical role in the apoptotic induction of HSV-1 infected cells.

RESULTS

ICP27 induces apoptosis

To date, the effect of ICP27 on apoptosis in HSV-1 infected cells remains unclear. In order to clarify whether ICP27
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induces apoptosis, we first expressed ICP27 in HEK 293T cells and monitored apoptosis by using the Hoechst 33258 staining assay, to visualize the apoptotic nuclear morphological alterations. Evidently, the chromatin condensation in ICP27 expressing cells (Fig. 1A). Next, we performed flow cytometric analysis with Annexin V-FITC/PI dual staining to quantify the extent of apoptosis. As shown in Fig. 1B, the percentage of the apoptotic cells in the upper right quadrant, which were positive for both Annexin V-FITC and PI, increased in the presence of ICP27. By Western blotting, we further analyzed the effect of ICP27 on caspase 3 in 293T cells. ICP27 elevates the process of procaspase 3 to active caspase 3, an executor in apoptosis. Moreover, ICP27 increased the cleavage of poly (ADP-ribose) polymerase (PARP), which is a major substrate of caspase 3 (Fig. 1C). Collectively, these data indicate that ICP27 induces apoptosis in 293T cells.

ICP27-induced apoptosis is associated with mitochondrial dysfunction
The mitochondrial membrane potential (MMP) was evaluated in order to determine how ICP27 induces apoptosis. Dysfunction of the MMP is an important feature of mitochondrial damage. Flow cytometric assay with Rhodamine 123 staining showed that ICP27 increased the percentage of cells with low MMP (Fig. 2A). This result provided the evidence that ICP27-induced apoptosis in 293T cells is associated with mitochondrial dysfunction. The MMP is tightly regulated by the Bcl-2 family proteins. Therefore, we next investigated the effect of ICP27 on the localization of Bax, a representative apoptotic Bcl-2 family protein, which, when activated, translocates to the mitochondria from the cytoplasm. The localization of Bax was assessed by staining the cells expressing GFP-Bax and ICP27 with Mitotracker, a mitochondrial-specific red fluorescent dye. Microscopic data revealed that in the presence of ICP27, GFP-Bax was localized on mitochondria (Fig. 2B). The Bax translocation was also confirmed by a subcellular fractionation assay. As expected, ICP27 increased the Bax translocation from the cytoplasm to the mitochondria (Fig. 2C). Further evaluation confirmed that ICP27 induces cytochrome c release from the mitochondria to the cytoplasm (Fig. 2D).

ICP27 interferes with the interaction between Bax and 14-3-3
To delineate the mechanism underlying ICP27-induced translocation of Bax, we speculated that ICP27 might affect the sequestration of Bax by 14-3-3, because 14-3-3 is not only an

Fig. 1. HSV-1 ICP27 induces apoptosis. (A) Chromatin condensation of 293T cells induced by ICP27. Cells were transfected with ICP27 plasmid and stained with Hoechst 33258. Arrows indicate cells showing chromatin condensation. The graph represents three independent experiments, in which 200 cells were counted and scored for chromatin condensation. (B) Flow cytometry analysis of ICP27-induced 293T cell apoptosis. At 48 h after transfection, cells were stained with Annexin V-FITC/PI, and analyzed by flow cytometry. (C) Effect of ICP27 on the expressions of the apoptotic indicators caspase 3 and PARP. In 293T cells. At 48 h after transfection with ICP27 plasmid, cell extracts were prepared and analyzed by Western blotting with specific antibodies.

Fig. 2. ICP27 induces translocation of Bax and release of cytochrome c from the mitochondria. (A) Effect of ICP27 on the mitochondrial membrane potential in 293T cells. Cells were transfected with ICP27 expressing plasmid for 48 h, and stained with Rhodamine 123 for 30 min. The fluorescence intensity was measured by flow cytometry. (B) Subcellular localization of Bax by ICP27. Cells were transfected with plasmids expressing GFP-Bax and Flag-ICP27. At 48 h after transfection, cells were stained with Mitotracker for 30 min. Subcellular distribution of GFP-Bax and mitochondria (red) were determined by immunofluorescence microscopy. (C) Subcellular distribution of Bax by ICP27 in 293T cells. At 48 h after transfection with ICP27 plasmid, cells were fractionated into cytoplasmic and mitochondrial fractions. Fractionated proteins were immunoblotted with anti-Bax antibody. β-actin was used as a cytosolic marker. (D) Cytochrome c release from the mitochondria by ICP27. At 48 h after transfection with ICP27 plasmid, cells were fractionated into cytoplasmic and mitochondrial fractions. Fractionated proteins were immunoblotted with anti-cytochrome c antibody. α-tubulin was used as a cytosolic marker and COX IV was used as a mitochondrial marker.
important regulator for cytosolic sequestration of Bax but is also one of cellular proteins screened in the yeast two-hybrid assay using ICP27 as a bait (data not shown) (11). First, we checked the physical interaction between ICP27 and 14-3-3 by the pull-down assay. Among the seven 14-3-3 isoforms, 14-3-3\(\sigma\) was selected for evaluation, since it has been reported that 14-3-3\(\sigma\) is involved in subcellular localization of Bax (11). As seen in Fig. 3A, ICP27 interacts directly with 14-3-3\(\sigma\). Further examination revealed that the interaction between 14-3-3\(\sigma\) and Bax was inhibited by ICP27 (Fig. 3B). The mechanism how ICP27 inhibits the interaction between 14-3-3\(\sigma\) and Bax was investigated by examining whether ICP27 regulates the acetylation of 14-3-3\(\sigma\). The interaction between 14-3-3\(\sigma\) and its target protein is regulated by the phosphorylation and acetylation state of 14-3-3\(\sigma\) (16-18). As shown in Fig. 3C, ICP27 decreases the acetylation level of 14-3-3\(\sigma\), which is regulated by SIRT2, a member of the class III HDAC family (16). Therefore, we speculate that ICP27 elevates the interaction between 14-3-3\(\sigma\) and SIRT2. In order to test this hypothesis, 293T cells expressing ICP27 were immunoprecipitated with 14-3-3\(\sigma\) antibody and immunoblotted with the SIRT2 antibody. The result revealed that SIRT2 is co-immunoprecipitated with 14-3-3\(\sigma\), and ICP27 increased the binding between 14-3-3\(\sigma\) and SIRT2 (Fig. 3D). Taken together, these data suggest that ICP27 inhibits the interaction between Bax and 14-3-3\(\sigma\) by enhancing the 14-3-3\(\sigma\) deacetylation through SIRT2.

In HSV-1 infection, ICP27 inhibits the interaction between Bax and 14-3-3

We next investigated whether ICP27 affects Bax translocation during the early period of HSV-1 infection. 293T cells were infected with HSV-1 at MOI 10. At 6 h post infection, 293T cells were harvested and immunoprecipitated with 14-3-3\(\sigma\) antibody. In HSV-1 infected cells, there was decreased interaction between Bax and 14-3-3\(\sigma\), compared to the mock-infected cells (Fig. 4A). To further clarify the interaction between Bax and 14-3-3\(\sigma\) affected by ICP27, 293T cells were infected with the HSV-1 wt virus and \(\Delta 27\) virus. In HSV-1 wt-infected cells, the interaction between Bax and 14-3-3\(\sigma\) was significantly decreased when compared with mock and \(\Delta 27\) infected cells (Fig. 4B). Thus, in HSV-1 infection, ICP27 inhibits the interaction between Bax and 14-3-3\(\sigma\), by interacting with 14-3-3\(\sigma\).

**DISCUSSION**

Previous reports indicate that ICP27 is associated with apoptosis in HSV-1 infected cells (19, 20). However, it remains unclear whether ICP27 induces or inhibits apoptosis. Aubert et al. showed that ICP27 is required for the prevention of apoptosis using ICP27 deletion mutant virus (21, 22). Conversely, other studies showed that IE gene expression is required for apoptosis induction, and ICP27 induces apoptosis through activating the p38 mitogen-activated protein kinase (MAPK) signaling (19, 23). ICP27 regulates viral gene expression, and it is possible that other viral proteins regulated by ICP27 may also affect apoptosis. Hence, it would be inappropriate to investigate the role of ICP27 in regulating apoptosis using...
ICP27 deletion virus exclusively, and it is more suitable to assess the role of ICP27 in apoptosis by using the protein itself.

Apoptosis is mediated by diverse stimuli, which induce the disruption of the mitochondrial membrane permeability. The consequent release of cytochrome c from the mitochondria to the cytoplasm in response to apoptotic stimulation is a critical event in apoptosis. The mitochondrial membrane integrity is tightly regulated by Bcl-2 family proteins. Particularly, Bax oligomerizes and translocates to the mitochondrial outer membrane, where it forms supramolecular openings under apoptotic stimuli (24, 25). Many studies have shown that diverse viral proteins induce apoptosis by regulating the Bcl-2 family proteins. For example, hepatitis B virus X protein induces apoptosis via enhancement of the Bax translocation (26). Our data support the reports that viral protein induces apoptosis through modulation of the Bcl-2 family protein. We showed that ICP27 induces apoptosis in 293T cells through mitochondrial dysfunction. We also found that ICP27 enhances the translocation of Bax to the mitochondria.

The subcellular localization of Bax is regulated by the 14-3-3 proteins. The 14-3-3 proteins interact with Bax, sequestering it to the cytoplasm. Recent studies show that post-translational modification of 14-3-3 proteins is important for its binding activity with target proteins (27). Phosphorylation of 14-3-3 proteins leads to self-dimerization, and affects the interaction with target proteins (28). SDK1 and JNK phosphorylate the 14-3-3 proteins, leading to the dissociation of Bax from the 14-3-3 proteins (29, 30). Lysine acetylation of 14-3-3 proteins also affects the interaction with target proteins, but the effect remains ambiguous. We showed that ICP27 decreases the acetylation level of 14-3-3 by enhancing the interaction between SIRT2 and 14-3-3, and subsequently inhibits the interaction between 14-3-3 and Bax. These results suggest new molecular mechanisms of apoptosis modulated by the HSV-1 ICP27.

MATERIALS AND METHODS

Cells and reagents

HEK 293T, Vero, and Vero 3-3 (HSV-1 ICP27 expressing Vero) cells were maintained in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum (FBS), at 37°C in 5% CO2. Cells were transfected using PEI (Sigma-Aldrich), according to the manufacturer’s instruction. Trichosertatin A (TSA) was purchased from Sigma-Aldrich. Plasmids expressing pEGFP-SUMO-1 and HA-UBC9 were provided by Dr. H. Ariga (35), respectively. pEGFP-Bax was provided by Dr. Robert W. Nickells (36).

Viruses and virus infection

HSV-1 wild-type (wt) strain KOS and ICP27 deletion mutant (Δ27) virus (31) were propagated and titrated on Vero cells and 3-3 cells, respectively. For virus infection, cell monolayers were infected with either KOS or mutant virus, at a multiplicity of infection (MOI) of 10 plaque forming units (PFU)/cell.

Plasmids

ICP27 plasmids have been described previously (32). pBH-ICP27 was provided by Dr. Stephen A. Rice (33). Plasmids expressing pEGFP-SUMO-1 and HA-UBC9 were provided by Dr. Hsiu-Ming Shih (34) and Dr. H. Ariga (35), respectively. pEGFP-Bax was provided by Dr. Robert W. Nickells (36).

Detection of apoptosis (Hoechst 33258 staining)

At 48 h after transfection, cells were washed with phosphate-buffered saline (PBS), fixed with 4% paraformaldehyde for 15 min, and then permeabilized with 0.05% Triton X-100. The fixed cells were washed with PBS and stained with 1 mg/ml Hoechst 33258 (Sigma-Aldrich) for 10 min (37). Apoptosis of the cells was determined by chromatin condensation, under a fluorescent microscope (Olympus).

Annexin-V apoptosis assay

At 48 h after transfection, the level of apoptosis was detected using the Annexin V-FITC Apoptosis detection kit (BD Pharmingen), according to the manufacturer’s manual (38). Flow-cytometric analysis was performed using a Becton Dickinson FACS Calibur flow cytometer and Cell Quest software (Becton Dickinson).

Detection of the MMP

The MMP was measured by using rhodamine-123 (Rh123) (39). Cells were harvested by trypsinization, and re-suspended in medium at a density of 5 × 104 cells/ml. The cells were then incubated with 5 μM Rh123 at room temperature for 20 min, washed twice with PBS, re-suspended in tissue culture medium, and analyzed by a Becton Dickinson FACS Calibur flow cytometer.

Subcellular fractionation assay

Briefly, cells were suspended in fractionation buffer (250 mM sucrose, 20 mM HEPES pH 7.4, 10 mM KCl, 1.5 mM MgCl2, 1 mM EDTA, 1 mM EGTA, 1 mM DTT, 1 mM PMSF, 1 mM NaF, 1 mM Na3VO4, 1 μg/ml aprotinin, 1 μg/ml leupeptin, and 1 μg/ml pepstatin). The cell lysate was passed 10 times through a 25 ga needle using a 1 ml syringe, and stored on ice for 20 min. The lysates were centrifuged at 3000 rpm at 4°C for 5 min, and the supernatants were collected as cytosolic fractions. The nuclear pellets were re-suspended in the nuclear buffer (250 mM sucrose, 20 mM HEPES pH 7.4, 10 mM KCl, 1.5 mM MgCl2, 1 mM EDTA, 1 mM EGTA, 1 mM DTT, 1 mM PMSF, 1 mM NaF, 1 mM Na3VO4, 1 μg/ml aprotinin, 1 μg/ml leupeptin, and 1 μg/ml pepstatin). The nuclear pellets were washed with 0.05% Triton X-100, fixed with 4% paraformaldehyde for 15 min, and then permeabilized with 0.05% Triton X-100. The fixed cells were washed with PBS and stained with 1 mg/ml Hoechst 33258 (Sigma-Aldrich) for 10 min (37). Apoptosis of the cells was determined by chromatin condensation, under a fluorescent microscope (Olympus).

GST pull-down assay

Transiently transfected 293T cells were lysed in modified RIPA buffer (50 mM Tris-HCl pH 7.4, 1% NP-40, 0.25% Nonidet P-40, 150 mM NaCl, 1 mM EDTA, 1 mM PMSF, 1 mM NaF, 1 mM Na3VO4, 1 μg/ml aprotinin, 1 μg/ml leupeptin, and 1 μg/ml pepstatin) at 4°C for 30 min. The cell lysates were incubated overnight at 4°C with Glutathione-Sepharose 4B
beads (Incospharm). The bead complexes were washed three times with modified RIPA buffer, and the bound proteins were analyzed by Western blotting.

**Immunoprecipitation and Western blot analysis**

Cell lysates were prepared in modified RIPA buffer and incubated overnight at 4°C, with specific antibodies. The immunocomplexes were isolated using protein-A Sepharose (Incospharm), resolved by SDS-PAGE, and transferred to nitrocellulose membrane. Immunoblotting was performed with various antibodies. Anti-ICP27, anti-Bax, anti-14-3-3, anti-cytochrome c, anti-COX IV, anti-α-tubulin, anti-β-actin, anti-acetylated lysine, anti-SIRT2, goat anti-rabbit IgG-HRP, rabbit anti-goat IgG-HRP, and goat anti-mouse IgG-HRP antibodies were purchased from Santa Cruz. Anti-Flag and anti-GST antibodies were purchased from Sigma-Aldrich.

**Immunofluorescence assay**

Vero cells were plated on sterile glass coverslips and transfected with GFP-Bax and GST-ICP27. At 48 h after transfection, cells were incubated with 20 ng/ml of a mitochondrial-specific dye (Mitotracker red CMXRos; Cambrex BioScience) for 30 min. Cells were fixed with 4% paraformaldehyde, at room temperature for 10 min, and permeabilized for 5 min with PBS containing 0.2% Triton X-100. Fluorescence microscopy was performed using an Olympus microscope.

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**CONFLICTS OF INTEREST**

The authors have no conflicting financial interests.

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