Survivin Ser81 Plays An Important Role in PI3K/Akt/mTOR Signaling Pathway

Ferry Sandra¹,²

¹Department of Biochemistry and Molecular Biology, Division of Oral Biology, Faculty of Dentistry, Universitas Trisakti, Jakarta, Indonesia
²BioCORE Laboratory, Faculty of Dentistry, Universitas Trisakti, Jakarta, Indonesia

Background: Survivin, a member of the inhibitor of apoptosis protein family, has been associated with protection from cell apoptosis and regulation of mitosis. Phosphorylated-Survivin at Ser81 was reported to provide cytoprotection against tumor necrosis factor (TNF)-related apoptosis-inducing ligand (TRAIL) in L929 cells by inducing a backloop activation of phosphatidylinositol 3-kinase (PI3K). Therefore Akt as a possible substrate of PI3K was investigated.

Materials and Methods: L929 cells were pretreated with/without 50 μM LY294002 or 10 μM Perifosine, and infected with viral particle of Survivin, anti sense of Survivin, Ser81Ala mutated Survivin or vector only. Cells were then harvested, lysed and subjected to immunoblot assay to detect Akt, phosphorylated Akt (Ser473), mammalian target of rapamycin (mTOR), phosphorylated-mTOR (Ser2448).

Results: Survivin induced Akt and mTOR phosphorylations in a viral particle concentration dependent manner. Pretreatment of LY294002 or Perifosine prior to Survivin infection, attenuated Akt or mTOR phosphorylations, respectively. Low Akt or mTOR phosphorylations were observed when L929 cells were infected with Ser81Ala mutated Survivin.

Conclusions: Ser81 phosphorylation site of Survivin played an important role in activating Survivin/PKA/PI3K/Akt/mTOR signaling pathway.

Keywords: survivin, Ser81, Akt, mTOR, LY294002, perifosine

Introduction

Survivin, a member of the inhibitor of apoptosis protein family, has been associated with protection from cell apoptosis and regulation of mitosis.¹,² In most finally differentiated adult tissues, Survivin expression is low to undetectable.¹,² However, Survivin is overexpressed in advanced cancers pertaining to poor prognosis, high recurrence and resistance to therapy.¹,² Survivin was reported to be involved in all tumor stages, started from initiation, maintenance, until development of tumor.¹ Therefore, antisurvivin has been suggested as a new approach for cancer therapy.¹,²

Survivin has been shown to play role in multiple² and back-loop³,⁴ signaling pathways. Survivin could be activated due to its phosphorylation at Thr34⁵ and Ser8.⁶,⁷,⁸ Among many other molecules, cyclin B1 and p34cdc2 were reported in activation of Survivin at Thr34⁶, while protein...
kinase A (PKA) and phosphatidylinositol 3-kinase (PI3K) were reported in activation of Survivin at Ser81. Survivin phosphorylation can also occur at Thr346, Thr117 and Ser81. Inhibition of Survivin phosphorylation by PH domain leucine-rich repeat protein phosphatase (PHLPP) induces cell apoptosis and exerts anticancer activity in gallbladder cancer.

To date, Survivin has been shown to induce vascular endothelial growth factor (VEGF)/PI3K/Akt, PKA/PI3K, and PI3K/Akt/hypoxia-inducible factor (HIF)-1α signaling pathways. However phosphorylation of Survivin as the key role of Survivin activation has not been intensely investigated. Phosphorylation of Survivin at Thr346 was the mostly investigated, other important phosphorylation sites were Thr117 and Ser81. Survivin Ser81 was reported to provide cytoprotection against tumor necrosis factor (TNF)-related apoptosis-inducing ligand (TRAIL) in L929 cells. Survivin Ser81 was reported to be important in back-loop Survivin/PKA/PI3K signaling pathway. Akt was reported as the substrate of PI3K in overexpressed-Survivin model, current study was initiated to investigate the possibility of Akt as the downstream of PKA/PI3K in Survivin phosphorylation at Ser81-modulated L929 cells.

Materials and methods

Survivin constructs, viral production and L929 cells infection

Preparation of Survivin, Antisense Survivin (Survivin-AS) and Ser81Ala mutants (Survivin-S81A) were described in our previous report. Briefly, Survivin, Survivin-AS, Survivin-S81A mutants cDNAs were polymerased, inserted into vector and transfected into BOSC23 cells. BOSC23 cells-produced viruses were harvested and tittered. L929 cells were cultured in α-DMEM containing 10% horse serum. Infection was carried out using viral product of BOSC23 cells (Survivin, Survivin-AS, Survivin-S81A or vector only) for 48 hours.

Cell treatment and lysate preparation

L929 cells were pretreated with/without 50 µM LY294002 (Cell Signaling, Danvers, MA, USA) or 10 µM Perifosine (Cell Signaling), and infected with viral particle of Survivin, Survivin-AS, Survivin-S81A or vector only. Cells were then harvested and lysed in a cold lysis buffer (20 mM Tris- HCl buffer pH 8.0, 500 mM NaCl, 1 mM EDTA, 1 mM EGTA, 10 mM β-glycerophosphate, 10 mM NaF, 10 mM pNPP, 0.4 mM Na3VO4, 1 mM benzamidine, 2 mM phenylmethylsulfonyl fluoride, aprotinin, 1 mg/ml leupeptin, 1 mM dithiothreitol and 10% Nonidet P-40). Immuno blot assay

Samples were separated by SDS-polyacrylamide gel electrophoresis and transferred to a polyvinylidene difluoride sheet. After blocking with 5% skim milk in Tris-buffered saline, the sheet was incubated with rabbit polyclonal anti-Akt (Cell Signaling), rabbit polyclonal anti-phospho-Akt (Ser473) (Cell Signaling), rabbit polyclonal anti-mammalian target of rapamycin (mTOR) (Cell Signaling) or rabbit polyclonal anti-phospho-mTOR (Ser2448) (Cell Signaling) antibody. The secondary antibody was horseradish peroxidase-conjugated goat anti-rabbit IgG antibody (Cell Signaling). The bound antibodies were visualized using Clarity Western ECL (Bio-Rad, Hercules, CA, USA) and captured using Alliance 4.7 (UVItech, Cambridge, UK).

Results

Survivin induced Akt phosphorylation

In Figure 1, basal Akt phosphorylation (Ser473) band of L929 cells (upper lane 1 from left) was seen. Upon Survivin infection in concentration of 7.5x10⁷ viral particle/ml, density of Akt phosphorylation (Ser473) band was increased (upper lane 3 from left). Density of Akt phosphorylation (Ser473) band was increased markedly when the cells were infected with Survivin in concentration of 75x10⁷ viral particle/ml (upper lane 4 from left). Similar band densities of Akt were observed for all lower lanes, showing that the same volume of proteins were electrophorated and detected.

![Figure 1. Survivin induced Akt phosphorylation](image)

Survivin induced Akt phosphorylation via PI3K

In Figure 2, high density bands of Akt phosphorylation (Ser473) were observed when L929 cells were infected with 75x10⁷ viral particle/ml retrovirus of Survivin with/
Survivin induced Akt phosphorylation via PI3K.

L929 cells were pretreated with/without 50 μM LY294002 for 2 hours prior to infection with/without 7.5x10^7 viral particle/ml retrovirus of Survivin for 48 hours as indicated in the panel. Cells were lysed and immunoblotted using anti-Akt or anti-phospho-Akt (Ser473) antibody. DMSO was used as negative control. Detailed procedures are described in "Materials and Methods". Each panel shows the typical result of 3 independent results.

Survivin Ser81 in Akt phosphorylation

Low density bands of Akt phosphorylation (Ser473) were observed when L929 cells were not infected, infected with vector merely, infected with 7.5x10^7 viral particle/ml retrovirus of Survivin-AS and infected with 7.5x10^7 viral particle/ml retrovirus of Survivin-S81A, as shown in Figure 3 (upper lane 1, 2, 4 and 5 from left, respectively). Meanwhile, a high density band of Akt phosphorylation (Ser473) was observed when L929 cells were infected with 7.5x10^7 viral particle/ml retrovirus of Survivin (upper lane 3 from left). Similar band densities of Akt were observed for all lower lanes, showing that the same volume of proteins were electrophorated and detected.

Survivin Ser81 in mTOR phosphorylation

Low density bands of mTOR phosphorylation (Ser2448) were observed when L929 cells were not infected, infected with vector merely, infected with 7.5x10^7 viral particle/ml retrovirus of Survivin-AS and infected with 7.5x10^7 viral particle/ml retrovirus of Survivin-S81A, as shown in Figure 5 (upper lane 1, 2, 4 and 5 from left, respectively). Meanwhile, a high density band of mTOR phosphorylation (Ser2448) was observed when L929 cells were infected with 7.5x10^7 viral particle/ml retrovirus of Survivin (upper lane 3 from left). Similar band densities of mTOR were observed for all lower lanes, showing that the same volume of proteins were electrophorated and detected.
Current results showed that Survivin induced mTOR phosphorylation at Ser2448 in a concentration dependent manner. By pretreatment of Perifosine, an Akt inhibitor, the mTOR phosphorylation was diminished, suggesting that Akt was the upstream of Survivin-induced mTOR phosphorylation. In Survivin S81A-infected L929 cells, the mTOR phosphorylation bands were not clearly seen as well.

Previously we reported that Survivin induced PI3K/PI3K. Hence, by combining with current results, we suggest signaling pathway of Survivin/PKA/PI3K/mTOR. This signaling pathway could serve as a survival pathway in L929 cells that attenuated TRAIL’s potential in inducing apoptosis. In addition Ser81 phosphorylation site of Survivin played an important role in activating the signaling pathway. Taken together, our current results suggest that Ser81 Survivin play an important role in inducing PKA/PI3K/mTOR survival signaling pathway. Further investigation is necessary to disclose other potential downstream of Survivin/PKA/PI3K-regulated Akt.

Discussion

Current results showed that Survivin induced Akt phosphorylation at Ser473 in a concentration dependent manner. By pretreatment of LY294002, a PI3K inhibitor, the Akt phosphorylation was diminished, suggesting that PI3K was the upstream of Survivin-induced Akt phosphorylation. These results were in accordance to previous reports, that Survivin induced PI3K/Akt signaling pathway. In vector-infected and Survivin-AS-infected L929 cells, the Akt phosphorylation bands were not clearly seen, showing that the phosphorylation was induced specifically by Survivin. In addition, Akt phosphorylation was not clearly seen in Survivin S81A-infected L929 cells, showing that Ser81 phosphorylation site of Survivin is important in this signaling pathway.

PI3K/Akt constitute an important pathway regulating the signaling of multiple biological processes such as apoptosis, metabolism, cell proliferation and cell growth. Components related to PI3K/Akt pathway have been reported widely as causal factors in cancer. Activation of PI3K and Akt are reported to occur in breast, ovarian, blood, pancreatic, esophageal, gallbladder and other cancers.

mTOR, one of the downstream of Akt, has been widely reported. PI3K/Akt/PTEN/mTOR signaling pathway plays a crucial role in regulating a broad range of cellular functions including cell growth, proliferation, cell survival, angiogenesis, invasion and migration, apoptosis, autophagy, cell cycle, DNA repair, chemoresistance and radioresistance in cancer cells. PI3K convertsPIP2 into PIP3, then Akt will be attracted by PIP3 and phosphorylated by phosphoinositide-dependent protein kinase (PDK), which subsequently causes alteration of numerous cell functions including the activation of mTOR and its substrates.

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