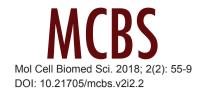
### **RESEARCH ARTICLE**



# Survivin Ser81 Plays An Important Role in Pl3K/Akt/mTOR Signaling Pathway

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**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  $\mu$ M LY294002 or 10  $\mu$ M 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.<sup>1,2</sup> In most finally differentiated adult tissues, Survivin expression is low to undetectable.<sup>1,2</sup> However, Survivin is overexpressed in advanced cancers pertaining to poor prognosis, high recurrence and resistance to therapy.<sup>1,2</sup> Survivin was reported

to be involved in all tumor stages, started from initiation, maintenance, until development of tumor. Therefore, antisurvivin has been suggested as a new approach for cancer therapy. 1,2

Survivin has been shown to play role in multiple<sup>2</sup> and back-loop<sup>3-5</sup> signaling pathways. Survivin could be activated due to its phosphorylation at Thr34<sup>6</sup> and Ser8<sup>13,7,8</sup>. Among many other molecules, cyclin B1 and p34<sup>cdc2</sup> were reported in activation of Survivin at Thr34<sup>6</sup>, while protein

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kinase A (PKA) and phosphatidylinositol 3-kinase (PI3K) were reported in activation of Survivin at Ser8<sup>13,7,8</sup>. Survivin phosphorylation can also occur at Thr53, Thr117 and Ser20.<sup>9</sup> Inhibition of Survivin phosphorylation by PH domain leucine-rich repeat protein phosphatase (PHLPP) induces cell apoptosis and exerts anticancer activity in gallbladder cancer.<sup>9</sup>

To date, Survivin has been shown to induce vascular endothelial growth factor (VEGF)/PI3K/Akt, 4 PKA/PI3K, 3 and PI3K/Akt/hypoxia-inducible factor (HIF)-1α signaling pathways<sup>10</sup>. However phosphorylation of Survivin, as the keyrole of Survivin activation has not been intensely investigated. Phosphorylation of Survivin at Thr346,11 was the mostly investigated, other important phosphorylation sites were Thr11712 and Ser813,7,8. Ser81 of Survivin 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.<sup>3</sup> Since Akt was reported as the substrate of PI3K in overexpressed-Survivin model<sup>4</sup>, 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  $\alpha$ -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  $\mu$ M LY294002 (Cell Signaling, Danvers, MA, USA) or 10  $\mu$ M 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  $\beta$ -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).

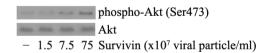
#### Immunoblot assay

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 7.5x10<sup>7</sup> 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.** L929 cells were infected with/without 1.5-75x107 viral particle/ml retrovirus of Survivin for 48 hours. Infected cells were then lysed and immunoblotted using anti-Akt or anti-phospho-Akt (Ser473) antibody. Detailed procedures are described in "Materials and Methods". Each panel shows the typical result of 3 independent results.

#### Survivin induced Akt phosphorylation via PI3K

In Figure 2, high density bands of Akt phosphorylation (Ser473) were observed when L929 cells were infected with 75x10<sup>7</sup> viral particle/ml retrovirus of Survivin with/

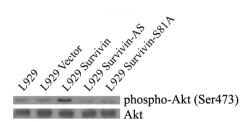
without pretreatment of DMSO (upper lane 4 and 2 from left, respectively). Meanwhile, pretreatment of LY294002 prior to infection with 75x10<sup>7</sup> viral particle/ml retrovirus of Survivin, resulted a low density band of Akt phosphorylation (Ser473) (upper lane 3 from left). Similar band densities of Akt were observed for all lower lanes, showing that the same volume of proteins were electrophorated and detected.

- - - + DMSO - - + - LY294002 - 75 75 75 Survivin (x10<sup>7</sup> viral particle/ml) phospho-Akt (Ser473)

Figure 2. Survivin induced Akt phosphorylation via PI3K. L929 cells were pretreated with/without 50  $\mu$ M LY294002 for 2 hours prior to infection with/without  $75x10^7$  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  $75x10^7$  viral particle/ml retrovirus of Survivin-AS and infected with  $75x10^7$  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 75x107 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.



**Figure 3. Survivin Ser81 in Akt phosphorylation.** L929 cells were infected with 75x10<sup>7</sup> viral particle/ml retrovirus of Survivin, Survivin-AS, Survivin-S81A or vector merely for 48 hours. Infected cells were then lysed and immunoblotted using anti-Akt or anti-phospho-Akt (Ser473) antibody. Detailed procedures are described in ""Materials and Methods". Each panel shows the typical result of 3 independent results.

#### Survivin induced mTOR phosphorylation via Akt

In Figure 4, density of mTOR phosphorylation (Ser2448) band was increased upon Survivin infection in concentration of 7.5x10<sup>7</sup> viral particle/ml (upper lane 3 from left). Density of mTOR phosphorylation (Ser2448) band was increased markedly when the cells were infected with Survivin in concentration of 75x10<sup>7</sup> viral particle/ml (upper lane 4 from left). A high density band of mTOR phosphorylation (Ser2448) was also observed when L929 cells were infected with 75x10<sup>7</sup> viral particle/ml retrovirus of Survivin with pretreatment of ethanol (upper lane 6 from left). Meanwhile, pretreatment of Perifosine prior to infection with 75x10<sup>7</sup> viral particle/ml retrovirus of Survivin, resulted a low density band of mTOR phosphorylation (Ser2448) (upper lane 5 from left). Similar band densities of mTOR were observed for all lower lanes, showing that the same volume of proteins were electrophorated and detected.

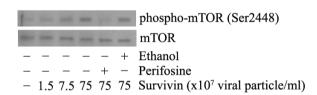
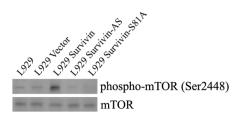


Figure 4. Survivin induced mTOR phosphorylation via Akt. L929 cells were pretreated with/without 10  $\mu$ M Perifosine for 1 hour prior to infection with/without 1.5-75x10<sup>7</sup> viral particle/ml retrovirus of Survivin for 48 hours as indicated in the panel. Cells were lysed and immunoblotted using anti-mTOR or anti-phospho-mTOR (Ser2448) antibody. Detailed procedures are described in "Materials and Methods". Ethanol was used as negative control. Each panel shows the typical result of 3 independent results.

#### 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 75x10<sup>7</sup> viral particle/ml retrovirus of Survivin-AS and infected with 75x10<sup>7</sup> 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 75x10<sup>7</sup> 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.



**Figure 5. Survivin Ser81 in mTOR phosphorylation.** L929 cells were infected with 75x10<sup>7</sup> viral particle/ml retrovirus of Survivin, Survivin-AS, Survivin-S81A or vector merely for 48 hours. Infected cells were then lysed and immunoblotted using anti-mTOR or anti-phospho-mTOR (Ser2448) antibody. Detailed procedures are described in ""Materials and Methods". Each panel shows the typical result of 3 independent results.

#### **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, 4.10 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. 13,14 Components related to PI3K/Akt pathway have been reported widely as causal factors in cancer. 13-15 Activation of PI3K and Akt are reported to occur in breast, ovarian, blood, pancreatic, esophageal, gallbladder and other cancers. 13,14

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 converts PIP<sub>2</sub> into PIP<sub>3</sub>, then Akt will be attracted by PIP<sub>3</sub> and phosphorylated by phosphoinositide-dependent protein kinase (PDK), which subsequently causes alteration of numerous cell functions including the activation of mTOR and its substrates. Page 16.

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 PKA/PI3K.<sup>3,8</sup> Hence, by combining with current results, we suggest signaling pathway of Survivin/PKA/PI3K/Akt/mTOR. This signaling pathway could serve as a survival pathway in L929 cells that attenuated TRAIL's potential in inducing apoptosis.<sup>7</sup> 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/Akt/mTOR survival signaling pathway. Further investigation is necessary to disclose other potential downstream of Survivin/PKA/PI3K-regulated Akt.

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