Actein induces apoptosis in leukemia cells through suppressing RhoA/ROCK1 signaling pathway

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Abstract. Actein is a tetracyclic triterpenoid compound, extracted from the rhizome of *Cimicifuga foetida*, exhibiting anticancer activities as previously reported. However, the effects of actein on human leukemia have not been explored before. In this study, the role of actein in regulating apoptosis induction in human leukemia cells was investigated. Actein administration significantly enhanced apoptosis, especially in human leukemia cell line of U937 and the primary human leukemia cells. The promotion was accompanied by caspase-9, caspase-3 and poly(ADP-ribose) polymerase (PARP) cleavage, and cytochrome *c* (Cyto-*c*) release. Additionally, translocation of Bax into mitochondria was increased by actein, while anti-apoptotic signals of myeloid cell leukemia-1 (Mcl-1) and B cell CLL/lymphoma 2 (Bcl-2) were decreased, accompanied by reduced phosphorylated Bcl-2-associated death promoter (Bad). Furthermore, protein kinase B (AKT) activation was downregulated by actein treatment in U937 cells. RhoA, but not caspase-3, regulated Rho kinase 1 (ROCK1) expression induced by actein. Suppression of RhoA and ROCK1 reduced ROCK1 expression, caspase-9, caspase-3 and PARP cleavage. In contrast, AKT inactivity enhanced apoptosis levels, as well as caspase signaling pathway expression. The anticancer role of actein was potentiated by inactivating AKT. *In vivo*, U937-bearing tumor growth was suppressed by actein, which was related to ROCK1 suppression, AKT dephosphorylation and apoptosis induction. These results indicated that actein has a

suppressive role in human leukemia progression through inactivating RhoA/ROCK1 and inducing caspases.

Introduction

Presently, acute promyelocytic leukemia is characterized by the differentiation arrest during the promyelocytic stage and elevation of the hematopoietic stem cells (1,2). Chemotherapy is the initial treatment of choice. Due to the abnormal collection of immature precursors, as well as the suppression of normal hemopoiesis, acute promyelocytic leukemia represents as a medical emergency, which causes a high level of early fatalities from the massive hemorrhage (3-5). Most patients with acute promyelocytic leukemia will receive a combination of medications. There are no surgical options owing to the body-wide distribution of malignant cells (6,7). Therefore, finding effective and new therapeutic strategies and revealing the underlying molecular mechanisms regulating leukemia cells are necessary and could be beneficial and useful for patients with acute promyelocytic leukemia.

Actein is a tetracyclic triterpenoid compound, isolated from the rhizome of *Cimicifuga foetida* (8). *Cimicifuga* species has a long history for medicine to protect people suffering from rheumatism, sore throat, and diarrhea in North America (9). In Asia, *Cimicifuga* species are used owing to its various bioactivities, such as antidiabetic, anti-osteoporosis and antiviral (10). In addition, the extracts from rhizome of *Cimicifuga* species have been applied to prevent female-related diseases clinically (11). Hence, *Cimicifuga* species could be considered as a natural medicinal herb with promising medicinal values. Moreover, actein was selective for human breast tumor cells, which could synergize with other chemotherapy agents to inhibit tumor growth (12). Moreover, p53 signaling pathway was revealed to be modulated by actein (11). P53, as a tumor suppressor, plays an essential role in apoptosis induction through regulating caspases (13-15). However, the effects of actein on modulating tumor growth, including human leukemia, is poorly understood. Herein, we attempted to explore if actein could suppress human leukemia development though inducing apoptosis and to reveal the underlying molecular mechanism.

Rho-associated kinase (ROCK) is reported as a serine/threonine kinase and one of the major downstream effectors of the small GTPase RhoA (16). The RhoA/ROCK signaling pathway is closely associated with the pathogenesis of various

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Abbreviations: PARP, poly(ADP-ribose) polymerase; Cyto-*c*, cytochrome *c*; Mcl-1, myeloid cell leukemia-1; Bcl-2, B cell CLL/lymphoma 2; Bad, Bcl-2-associated death promoter; AKT, protein kinase B; ROCK1, Rho kinase 1; PTEN, phosphatase and tensin homolog; FBS, fetal bovine serum; TUNEL, terminal deoxynucleotidyl transferase (TdT) dUTP nick-end labeling; MTT, 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide; Bax, Bcl-2-associated X protein; BCA, bicinchoninic acid

Key words: actein, human leukemia, apoptosis, RhoA, Rho kinase 1

disorders and is also involved in a number of aspects of tumors, including human leukemia (17,18). Rho could modulate the cell actin cytoskeleton through its down-stream effective factor of ROCK, which is highly included in the biological processes of cell movement, cell migration, gene transcription, nerve regeneration, and apoptosis (19,20). Also, elevation of RhoA/ROCK has been reported in tumors and associated with cancer development (21). Therefore, targeting RhoA/ROCK signaling pathway might be a potential therapeutic strategy for human leukemia treatment. Though it has been reported in leukemia progression, further study is still required to further reveal the underlying molecular mechanism. In this study, we attempted to explore the role of actein in modulating human leukemia cell progression through apoptosis induction, which relied on regulation of RhoA/ROCK1. *In vivo*, the U937-bearing tumor growth was inhibited by actein treatment. The suppressive role of actein in human leukemia included its effects on AKT dephosphorylation, phosphatase and tensin homolog (PTEN) activation, pro-apoptotic signal promotion as well as anti-apoptotic molecule reduction. The results suggested that actein could be an effective candidate for human leukemia.

Materials and methods

Cells and culture. Human leukemia cell lines, U937, K562 and NB4, were purchased from American Type Culture Collection (ATCC, USA). Human hepatocyte cell line L02 and human tubular epithelial cells HK2 were purchased from KeyGen Biotech Co., Ltd. (Nanjing, China). All cells were cultured in RPMI-1640 medium, which is supplemented with 10% fetal bovine serum (FBS) (Gibco, Invitrogen, USA), 100 U/ml penicillin and 100 µg/ml streptomycin at 37°C in a 5% CO₂ humidified environment. The peripheral blood samples in our studies were isolated from 10 patients with acute promyelocytic leukemia after acquiring the informed consent: two patients are M2, four are M4, and four are M5 following the French-American-British (FAB) classification system. Approval for the study was obtained from Huai'an First People's Hospital, Nanjing Medical University (Jiangsu, China). Acute promyelocytic leukemia blasts were extracted using Histopaque-1077 density gradient centrifugation (Sigma-Aldrich, USA) for 15 min at 600 g. The isolated mononuclear cells were then suspended in RPMI-1640 medium at 8x10⁵/ml. Peripheral blood mononuclear cells from blood samples collected from the healthy volunteers were also isolated by Histopaque 1077 density gradient centrifugation.

As for the gene knockdown, the cells were transfected with (100 nM) nonsense control, siRNA against caspase-3 (#1:5'-UGU AGG AGA GUU GAG GUC GAG GU) and siRNA against ROCK1 (#1:5'-GAU UAU AGA GUG GUG GUG ACG GGU A) for 24 h using Lipofectamine 2000 (Invitrogen) according to the manufacturer's protocol. All small interfering RNAs were synthesised by GenePharma (Shanghai, China). After various treatments, further experiments were conducted. Actein (CAS:18642-44-9, 98% HPLC) used in our study was purchased from Shanghai Yuanye Bio-Technology Co., Ltd. (Shanghai, China). AKT inhibitor, LY294002, and caspase-3 inhibitor, Z-VAD-FMK, were purchased from Sigma-Aldrich, and ROCK1 inhibitor, Y-27632 dihydrochloride was obtained

from Tocris Bioscience (Bristol, UK). RhoA inhibitor, C3 exoenzyme, was purchased from Alexis Biochemicals (USA).

Flow cytometric analysis. After treatment under various conditions, flow cytometric analysis was used to determine apoptosis levels in U937 cells using Annexin V/PI staining kit (Roche, Switzerland) following the manufacturer's protocol. In order to analyze the mitochondria injury, 2x10⁵ U937 cells were cultured with 3,3-dihexyloxacarbocynine at the dose of 40 nM (DiOC₆, Sigma-Aldrich) in PBS for 20 min at 37°C. The results acquisition and analysis were performed using a Becton-Dickinson FACSCalibur flow cytometer with Cell Quest software.

Mitochondrial and cytosolic fractions. Mitochondria/cytosol fractionation kits (ab65320, Abcam, USA) were used to isolate the mitochondrial fraction from cytosolic fraction. Then, the enriched mitochondrial and cytosolic fractions were used for western blot analysis.

Hoechst 33258 staining. Hoechst 33258 staining of the cells was performed to evaluate the apoptosis induced by actein. The cells were seeded at a concentration of 1x10⁶ cells/ml in 6-well plates and treated with the indicated concentration of actein. The cells were harvested, washed twice with PBS, fixed with 4% formaldehyde for 10 min and stained with Hoechst 33258 (Sigma-Aldrich) staining solution following the manufacturer's instructions. The images were immediately photographed under a fluorescence microscope (Olympus, Japan).

Evaluation of apoptosis. After induction of actein for 24 h, leukemia cell apoptosis was measured using a commercial single-stranded DNA (ssDNA) enzyme-linked immunosorbent assay kit (Millipore Chemicon, USA), which detects ssDNA, corresponding to the most specific apoptosis end-product.

3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide (MTT) analysis. MTT (Beyotime, Nanjing, China) was used to calculate cell viability. Cells (2x10³/well) were seeded on 96-well plates and treated under different conditions as indicated and incubated at 37°C. MTT solution (300 µl/well) was added after incubation. Following incubation at 37°C for an additional 4 h, the supernatants were removed and 200 µl dimethyl sulfoxide (DMSO, Sigma-Aldrich) was added into each well to dissolve the formazan crystals. The 96-well plates were then placed in a microplate reader to determine the absorbance at 490 nm. Each test was carried out in triplicate.

Animal treatments. Twenty, 5-week-old male nu/nu mice (15-18 g) were injected with 5x10⁶ U937 cells subcutaneously. All mice were purchased from the Animal Experiment Center of Nanjing Medical University (Nanjing, China). Before the experiments, all mice were allowed to adapt to the environment for a week. All protocols were in line with the Regulations of Experimental Animal Administration issued by the Ministry of Science and Technology of the People's Republic of China. Mouse care and usage were performed according to the ethical guidelines of Huai'an First People's Hospital, Nanjing Medical University. The mice were raised in air-conditioned

pathogen-free rooms ($25\pm 2^{\circ}\text{C}$, $50\pm 10\%$ humidity) under controlled lighting (12 h light/day) and fed with water and standard laboratory food. When the tumors were visible, the mice were randomly divided into two groups (10 mice per group). The control group received the vehicle (PBS) injection i.p., and the treatment group was administered with actein i.p. every day at a dose of 15 mg/kg (11). Sixty days later, all mice were sacrificed, tumors were excised and measured, and tumor tissues were fixed in 10% formalin. After embedding in paraffin, immunohistochemical analysis was performed.

Immunofluorescent analysis. U937 cells were treated with or without 40 μM actein for 24 h. Cells were then harvested through centrifugation, and fixed with 4% paraformaldehyde for 10 min, permeabilized using 0.1% Triton X-100 for 10 min, and finally blocked with 1% bovine serum albumin for 30 min at room temperature. Cells were further incubated with a primary antibody (Cyto-*c*, 1:200, Abcam) at 4°C overnight, and followed by a secondary goat anti-mouse IgG H&L (Alexa Fluor® 488) (Abcam) for 1 h at room temperature. After washing with PBS, the images were captured with a confocal microscope (Olympus, Japan).

Immunohistochemical assays. Paraffin-embedded tumor sections were used for the blinded assessment of caspase-3, Ki-67, ROCK1 and apoptosis levels, respectively. Mouse tumors were sectioned at 3 μM thickness, and terminal deoxynucleotidyl transferase (TdT) dUTP nick-end labeling (TUNEL) assay was carried out using light and electron microscopy-based kits (R&D Systems, USA) for detecting DNA fragments. For staining of ROCK1 (1:200, Abcam, UK), caspase-3 (1:200, Abcam) and Ki-67 (1:200, Abcam), the tumor sections were analyzed using a microscope. Images were arranged using the TissueFAXs (Tissue-Gnostics) software.

Western blot analysis. For western blotting, cells and tissue samples after various treatments were lysed in RIPA lysis buffer (150 mM NaCl, 0.1% Triton X-100, 0.5% sodium deoxycholate, 0.1% SDS, and 50 mM Tris-HCl, pH 8.0) to yield a homogenate. Also, the final supernatants were obtained by centrifugation at 12,000 g for 15 min. The protein concentration was calculated using bicinchoninic acid (BCA) protein assay kit (Thermo Scientific, USA) with bovine serum albumin as a standard. The total protein extract was later used for western blot analysis. Total protein (40 μg) was loaded and proteins were separated using SDS-PAGE and electrophoretically transferred to polyvinylidene difluoride membranes (Millipore, USA). The membranes were then blocked with 5% non-fat dry milk in Tris buffered saline (20 mM Tris, pH 7.6, 137 mM NaCl) with 0.1% Tween-20, washed, and then incubated with primary antibody. The primary antibodies were as follows: rabbit anti-p-AKT (1:1,000, Cell Signaling Technology), rabbit anti-AKT (1:1,000, Cell Signaling Technology), rabbit anti-Mcl-1 (1:1,000, Abcam), rabbit anti-Bcl-x1 (Abcam, USA), rabbit anti-Bax (1:1,000, Abcam), rabbit anti-PTEN (1:1,000, Abcam), rabbit anti-p-PTEN (1:1,000, Abcam), rabbit anti-caspase-3 (1:1,000, Abcam), mouse anti-Bcl-2 (1:1,000, Cell Signaling Technology), rabbit anti-PARP (1:1,000, Cell Signaling Technology), rabbit anti-p-Bad (1:1,000, Abcam), rabbit anti-Bad (1:1,000, Abcam), rabbit anti-RhoA (1:1,000,

Abcam), rabbit anti-ROCK1 (1:1,000, Abcam), rabbit anti-Cox IV (1:1,000, Abcam), mouse anti-Cyto-*c* (1:1,000, Abcam) and anti-GAPDH (1:500, Santa Cruz Biotechnology, Inc.). Immunoreactive bands were visualized by ECL Immunoblot Detection system (Pierce Biotechnology, Inc., Rockford, IL, USA) and exposed to Kodak (Eastman Kodak Co., USA) X-ray film. Each protein expression level was defined as grey value (Version 1.4.2b, Mac OS X, ImageJ, National Institutes of Health, USA) and standardized to housekeeping gene of GAPDH and expressed as a fold of control.

Statistical analysis. Results are represented as the mean $\pm$ SEM of triplicate experiments. Statistically significant values were compared by use of the ANOVA and the Dunnett's post hoc test, and P-values of <0.05 were considered to indicate a statistically significant result.

Results

Actein triggers apoptosis and mitochondrial damage in human leukemia cells. Actein has been investigated in human breast cancer though apoptosis induction (10,11). Thus, here we first examined the role of actein in apoptosis and the mitochondrial injury in human leukemia cells, U937. Treating U937 cells with different concentrations of actein for 24 h significantly enhanced apoptosis (Fig. 1A, left) and mitochondrial injury in a dose-dependent manner (Fig. 1B, left). Furthermore, U937 cells were treated with 40 μM actein for the indicated time, ranging from 0 to 24 h. Flow cytometric analysis indicated that apoptosis was highly induced (Fig. 1A, right). Also, the mitochondria injury was observed in actein-treated cells, which was time-dependent (Fig. 1B, right). Following, the proteins isolated from total cell and cytoplasm, respectively, were used for western blot analysis. The results indicated that caspase-9, caspase-3 and PARP cleavage was dramatically increased after actein administration dose-dependently. Similarly, Cyto-*c* was found to be upregulated with the increasing of actein treatment in cytoplasm, indicating actein administration promoted Cyto-*c* release into the cytoplasm (Fig. 1C, left). In line with the results of flow cytometry, caspase-9, caspase-3 and PARP activation was significantly improved after actein treatment, which was shown in a time-dependent manner (Fig. 1C, right). Furthermore, Hoechst 33258 staining indicated that the number of apoptosis was significantly high in U937 and K562 cells after actein treatment (Fig. 2A). Cell apoptosis was also evaluated using ssDNA detection kit. Untreated leukemia cells served as the control. As shown in Fig. 2B, we found that treatment of actein significantly increased the apoptosis proportion in leukemia cell lines, and consistently, U937 cells were more sensitive to actein treatment. Also, western blot analysis indicated that caspase-9, caspase-3, and PARP cleavage and Cyto-*c* in cytoplasm were apparently induced by actein in 937 and K562 cells (Fig. 2C). The results here indicated that U937 and K562 are likely to be more sensitive to actein. Taken together, the data above indicated that actein could potentiate apoptosis in human leukemia cells.

Actein induces apoptosis response in primary human leukemia cells. We attempted to evaluate if actein could induce apoptosis in human primary leukemia cells. The

Figure 1. Actein triggers apoptosis and mitochondrial damage in human leukemia cells. (A) Left, the human leukemia cells of U937 were exposed to actein at the indicated concentrations, ranging from 0 to 40 μ M as indicated for 24 h. After treatments, all cells were harvested for apoptotic analysis using flow cytometry. Right, U937 cells were cultured with 40 μ M actein for, respectively, 0, 3, 6, 9, 15, 18 or 24 h. Then, all cells were collected for calculating the number of apoptotic cells using flow cytometry. (B) Left, U937 cells were treated at different doses of actein as described for 24 h and stained with DiOC₆. Then, flow cytometric analysis was used to evaluate the number of cells experiencing mitochondrial injury. Right, U937 cells were treated with 40 μ M actein for the indicated time, followed by DiOC₆ staining. Next, all cells were harvested for flow cytometric assays. (C) All U937 cells were treated as described. The extracts from the whole cells and the cytoplasm were used for western blot analysis using primary antibodies of caspase-9, caspase-3, PARP and Cyto-c. Data are analyzed as mean $\pm$ SEM, n=8. *P<0.05, **P<0.01 and ***P<0.001 versus the Con group in the absence of any treatments.

Figure 2. Actin induces apoptosis selectively in human leukemia cell lines. (A) Hoechst 33258 staining was used to calculate the effects of actein on different human leukemia cell lines treated with 40 μ M actein for 24 h. (B) Cell apoptosis was evaluated using ssDNA detection kit. (C) Human leukemia cell lines, including U937, NB4 and K562, were exposed to 40 μ M actein for 24 h, followed by western blot analysis of caspase-9, caspase-3, PARP and Cyto-c in the whole cell or the cytoplasm as indicated. Data are analyzed as mean $\pm$ SEM, n=8.

Figure 3. Actein induces apoptotic response in primary human leukemia cells. (A) Primary human leukemia cells were derived from peripheral blood of 10 patients. Immediate after treatment with actein (40 μ M) for 24 h, the percentage of apoptotic cells was calculated through flow cytometry. (B) The lysates from the whole cell and the cytosolic fractions in leukemia cells extracted from two patients with acute myeloid leukemia were collected for western blot analysis of caspase-9, caspase-3, PARP and Cyto-c. (C) Peripheral blood mononuclear cells isolated from normal human were cultured with 40 μ M actein for 0, 3, 6, 9, 15, 18 and 24 h. Then, flow cytometry was carried out to evaluate the number of apoptosis. (D) Peripheral blood mononuclear cells obtained from normal healthy people were exposed to 40 μ M actein for 24 h, followed by assessment of caspase-9, caspase-3, PARP and Cyto-c using western blot analysis. (E) Peripheral blood mononuclear cells, human hepatocyte cell line L02 and human tubular epithelial cells HK2 were exposed to actein for 24 h at the indicated concentrations. Then, all cells were harvested for MTT analysis. Data are analyzed as mean $\pm$ SEM, n=8. ***P<0.001 versus the Con group in the absence of any treatment.

human primary leukemia cells were extracted from 10 patients with acute myeloid leukemia. Then, actein was administered to cells. As shown in Fig. 3A, we found that actein significantly induced apoptosis in human primary leukemia cells. Western blot analysis further indicated that the activation of caspase-9, caspase-3 and PARP was markedly induced in actein-treated primary leukemia cells obtained from two patients. Also, Cyto-c releasing in the cytoplasm of human primary leukemia cells was also induced by using actein (Fig. 3B). Next, the effects of actein on peripheral blood mononuclear cells obtained from normal healthy people were explored. As shown in Fig. 3C, cells exposed to 40 μ M actein for various times showed no significant difference on apoptosis induction. In addition, western blot analysis in peripheral blood mononuclear cells and cytoplasm indicated that caspase-9, caspase-3, PARP and Cyto-c were not alternative either in the group with actein exposure or not (Fig. 3D). Peripheral blood mononuclear cells, human hepatocyte cell line L02 and human tubular epithelial cells HK2 were exposed to actein for 24 h at the indicated concentrations. Then, all cells were harvested for MTT analysis. From the results in Fig. 3E, we found that there

was no significant difference between the groups of cells. The results indicated that actein at the indicated concentrations used in our study showed no cytotoxicity to normal cells. The findings above indicated that actein could trigger apoptosis in human primary leukemia cells without any changes in peripheral blood mononuclear cells from normal people.

Actein causes apoptosis through regulating pro- and anti-apoptotic signals. Anti-apoptotic and pro-apoptotic signals are well characterized in regulating apoptosis (22,23). Thus, in order to explore the underlying molecular mechanism regarding to actein-induced apoptosis in human leukemia. Protein levels in total cellular, cytoplasm and mitochondria were calculated using western blot analysis. As shown in Fig. 4A, we found that Bcl-2, Bcl-xl and Mcl-1 expressed highly in actein-treated U937 cells. Bad phosphorylation, suppressing apoptosis, was reduced by actein. In contrast, Bad was observed with increased levels after actein administration in whole cells. Bax in cytoplasm was discovered with reduced protein abundance in actein-treated cells dose-dependently. Accordingly, in mitochondria, Bax was elevated, which indicated the ability

Figure 4. Actein causes apoptosis through regulating pro- and anti-apoptotic signals. (A) U937 cells were treated with actein (0, 5, 10, 20 and 40 μ M) for 24 h, followed by western blot analysis of Bcl-2, Bcl-xl, Mcl-1, p-Bad and Bad in the total cells. Also, protein extracted from cytoplasm and mitochondria was used for Bax using western blot assays. (B) U937 cells were treated with 40 μ M actein for the indicated time. Next, western blot analysis was used to assess the whole cellular Bcl-2, Bcl-xl, Mcl-1, p-Bad and Bad. Protein extracted from cytoplasm and mitochondria was used for Bax abundance from protein levels through western blot assays. (C) Actein (40 μ M) was exposed to U937 cells for 24 h. After treatments, all cells were performed for immunofluorescent analysis. (D) Acute myeloid leukemia blasts and (E) peripheral blood mononuclear cells from normal people were exposed to 40 μ M actein for 24 h, followed by western blot analysis of the whole cellular Bcl-2, Bcl-xl, Mcl-1, p-Bad and Bad, as well as Bax in cytoplasm and mitochondria. Data are analyzed as mean $\pm$ SEM, n=8. ***P<0.001 versus the Con group in the absence of any treatment.

of actein in inducing apoptosis. The results above were also triggered by actein in a time-dependent manner in U937 cells (Fig. 4B). Furthermore, immunofluorescent analysis confirmed that the expression of Bax, proved by the fluorescent intensity, in cytoplasm was suppressed by actein (Fig. 4C). Also, primary leukemia cells isolated from patient indicated that anti-apoptotic molecules of Bcl-2, Bcl-xl and Mcl-1 were suppressed by actein administration, as well as phosphorylated Bad, while Bad was significantly triggered. Additionally, actein reduced Bcl-2-associated X protein (Bax) expression in cytoplasm, and elevated its levels in mitochondria (Fig. 4D). Finally, western blot analysis of peripheral blood mononuclear cells extracted from normal people showed no difference of these signals in expression (Fig. 4E).

Actein treatment suppresses AKT and RhoA/ROCK1 signaling pathway. Referring to previous studies, PTEN and PI3K/AKT signaling pathway is involved in regulating apoptosis (24). PTEN is reported to negatively modulate the activation of PI3K/AKT (25,26). In this regard, we found that phosphorylated PTEN was highly induced by actein treatment in a dose- and time-dependent manner, subsequently reducing AKT phosphorylated levels (Fig. 5A and B). Similar

Figure 5. Actein treatment suppresses AKT and RhoA/ROCK1 signaling pathway. (A) U937 cells were exposed to actein at the described concentrations (0-40 μ M) for 24 h or (B) treated for the indicated time, ranging from 0 to 24 h, with 40 μ M actein. Then, the cells were harvested for western blot analysis using primary antibodies of p-PTEN, PTEN, p-AKT, AKT, RhoA and ROCK1. (C) Acute myeloid leukemia blasts and (D) peripheral blood mononuclear cells from normal people were treated with 40 μ M actein for 24 h. Next, western blot analysis was carried out to evaluate the protein levels of p-PTEN, PTEN, p-AKT, AKT, RhoA and ROCK1. Data are analyzed as mean $\pm$ SEM, n=8.

Figure 6. Caspase-3-independent ROCK1 activation is involved in actein-induced apoptosis. (A) U937 cells were pre-treated with 20 μ M caspase-3 inhibitor, ZVAD, for 2 h and then exposed to 40 μ M actein for 24 h, followed by flow cytometric analysis. (B) U937 cells were transfected with caspase-3 siRNA for 24 h to knockdown caspase-3 expression, followed by exposure to 40 μ M actein for 24 h, and then flow cytometric analysis was used to calculate apoptotic cells. (C) U937 cells were treated with caspase-3 inhibitor for 2 h to suppress caspase-3 expression, followed by exposure to 40 μ M actein for another 24 h, and then western blot analysis was performed to calculate cleaved caspase-3 and ROCK1 expression. (D) Caspase-3 was knocked down using specific siRNA sequence for 24 h, followed by exposure to 40 μ M actein for another 24 h. Then, all cells were harvested for western blot analysis. Data are analyzed as mean $\pm$ SEM, n=8. ***P<0.001 versus the actein group.

findings were observed in primary human leukemia cells but not found in peripheral blood mononuclear cells from normal human (Fig. 5C and D). Thus, we supposed that PTEN phosphorylation and AKT dephosphorylation were related to actein-induced apoptosis in leukemia cells. Following this, RhoA/ROCK1 signaling pathway was investigated in our study. ROCK1 is suggested to be an important target for RhoA, which modulates PTEN activation and regulates apoptotic response. As shown in Fig. 5A and B, actein induced RhoA and ROCK1 downregulation dose- and time-dependently. In agreement with the results above, in primary human leukemia cells, marked reduction of RhoA and ROCK1 were discovered in actein-treated groups, while no significant difference was observed in peripheral blood mononuclear cells extracted from normal people (Fig. 5C and D). In conclusion, the results above indicated that RhoA/ROCK1 was also involved in actein-modulated apoptosis.

Actein-induced apoptosis has a close relationship with RhoA/ROCK1 signaling pathway. Caspase-3 is a key during

apoptosis induction, having a close relationship with ROCK1 expression (27). Thus, in this regard, we attempted to explore the role of caspase-3 in actein-induced apoptosis, and its association with ROCK1. Fig. 6A and C indicated that U937 cells treated with caspase-3 inhibitor, ZVAD, dramatically reduced apoptosis and diminished caspase-3 activation. Also, caspase-3 knockdown with specific siRNA significantly eliminated apoptosis and the cleavage of caspase-3 (Fig. 6B and D). Of note, we found that both suppression of caspase-3 and knockdown of caspase-3 showed no effects on ROCK1 expression via western blot analysis (Fig. 6C and D). Together, the data above indicated that caspase-3 was not in our study included in the regulation of ROCK1 expression during the actein-triggered apoptosis. Thus, the expression of ROCK1 regulated by RhoA possibly was explored using RhoA inhibitor. U937 cells with RhoA inhibitor treatment enhanced apoptosis and cleaved caspase-3, caspase-9 and PARP expression, especially in actein-cotreatment group. In contrast, ROCK1 was markedly reduced, indicating that ROCK1 suppressed by actein was dependent on RhoA

Figure 7. Actein-induced apoptosis has a close relationship with RhoA/ROCK1 signaling pathway. (A) U937 cells were pre-treated with RhoA inhibitor, C3 exoenzyme ($10 \mu\text{M}$), for 2 h, followed by 24-h treatment of actein ($40 \mu\text{M}$). Then, flow cytometry and western blot assays were used to evaluate the apoptosis levels. (B) The inhibitor of ROCK1, Y-27632 dihydrochloride ($10 \mu\text{M}$), was used in U937 cells for suppressing ROCK1 expression, and actein was used to treat cells for another 24 h. Apoptosis was calculated using flow cytometry and western blot analysis. (C) ROCK1 was knocked down using specific siRNA for 24 h, which was then exposed to actein for another 24 h. After various treatments, all cells were harvested for apoptosis analysis using flow cytometry and western blot assays. (D) The inhibitor of AKT, LY294002 ($5 \mu\text{M}$), was used to treat U937 cells for 2 h, and then cells were exposed to actein for another 24 h, followed by flow cytometry and western blot analysis. Data are analyzed as mean $\pm$ SEM, $n=8$. *** $P<0.001$ versus the actein group.

expression (Fig. 7A). Then, ROCK1 expression was suppressed using its specific inhibitor and siRNA with the specific sequence. As shown in Fig. 7B and C, we found that suppression of ROCK1 apparently augmented apoptotic response, as well as accelerated caspase-9, caspase-3 and PARP cleavage, which was further upregulated by co-culture with actein. AKT phosphorylated levels could be diminished by using actein. Therefore, its inhibitor was further used here to calculate its effects in apoptosis induced by actein. Fig. 7D suggested that AKT suppression using its inhibitor significantly elevated apoptosis in U937 cells, and its combination with actein further improved apoptotic response, accompanied with upregulated caspase-9, caspase-3 and PARP cleavage. The results above suggested that ROCK1 expression was dependent on RhoA to modulate apoptosis induced by actein in human leukemia cells.

Actein suppresses U937-tumor growth and triggers apoptosis in xenograft mouse models. *In vitro*, actein was evidenced to be important in suppression of human leukemia cells. In order to further confirm our data above, *in vivo* experiments were conducted. The male nu/nu mice were subcutaneously injected with U937 cells. After inoculation, half of the total number of mice was administered with actein intraperitoneally for two months. As shown in Fig. 8A, we found that actein administration apparently prolonged the survival rate of mice bearing U937 tumors, which was comparable to the control

group. The tumor volume of mice was also dramatically reduced by actein treatment (Fig. 8B). However, no significant difference of body weight was observed between the two groups (Fig. 8C). Next, TUNEL assay and Ki-67 levels were performed to further reveal the role of actein *in vivo*. Ki-67-positive areas were reduced by actein treatment. In contrast, the number of TUNEL-positive cells was upregulated by using actein, indicating apoptosis induced by actein *in vivo* (Fig. 8D). Furthermore, actein reduced ROCK1 expression in the tumor sections, while caspase-3 cleavage was found to be enhanced (Fig. 8E). Finally, western blot analysis indicated that RhoA and phosphorylated AKT had low expression in actein-treated group, and caspase-3 cleavage was elevated after actein administration, which were indicative of apoptotic response (Fig. 8F). Thus, the results indicated that actein apparently suppressed U937 tumor growth through triggering apoptosis *in vivo*.

Discussion

According to previous studies, RhoA/ROCK1 signaling pathway plays an essential role in cell survival, which has been investigated in various functional states (18,28,29). For instance, several reports indicate evidence that inhibition of the Rho/ROCK signaling pathway induces hepatocellular cancer (HCC) cell apoptosis and endothelial cells via a mito-

Figure 8. Actin suppresses U937-tumor growth and triggers apoptosis in xenograft mouse models. (A) The survival rate was assessed in xenograft mouse models treated with or without actein. (B) The tumor volume was measured. (C) The body weight was assessed. (D) The immunochemical analysis of Ki-67 and TUNEL is exhibited. (E) ROCK1 and caspase-3 expression levels in tumor tissue sections were analyzed using immunohistochemical assays. (F) Western blot analysis of RhoA, caspase-3 and AKT in the tumor samples isolated from mice treated under various conditions. Data are analyzed as mean $\pm$ SEM, n=10. ***P<0.001 versus the Con group.

chondrial apoptosis pathway (30,31). Thus, suppression of RhoA/ROCK1 signaling pathway affects the transformation and proliferation of progenitor cells (32). In addition, as previously reported, RhoA/ROCK1 signaling pathway suppression in human primary leukemia cells enhances cell death, revealing that targeting RhoA/ROCK1 might be of potential value as therapy to prevent human leukemia. In our study, we found that, in line with previous studies, RhoA/ROCK1 pathway was involved in apoptosis of acute promyelocytic leukemia. Apoptosis is an evolutionarily conserved process, which irreversibly eliminates injured or potentially harmful cells in order to protect the organism. The caspases could be structurally divided according to the absence or presence of an N-terminal pro-domain (33). The caspases containing long pro-domains are the first to be activated responding to various apoptotic stimuli (34). The activated caspases damage the cellular architecture and eventually lead to cell death. Caspase-3 is reported as the most essential member of caspase family (35,36). According to studies before, caspase-3 activation has an association with ROCK1 expression (37). Caspase-3 suppression reduced ROCK1 expression (38). However, there

was also a study, which indicated that caspase-3-independent regulation of ROCK1 was observed during apoptosis induction (39). In the present study, U937 cells were exposed to caspase-3 inhibitor of ZVAD and actein diminished caspase-3 cleavage, and apoptosis, while ROCK1 showed no difference, which illustrated that ROCK1 expression was caspase-3-independent, at least in our study, and that other factors might be involved in regulating ROCK1 expression. Attributing to the role of RhoA in modulating ROCK1, it was investigated in our study. Actein exposure induced RhoA downregulation. Also, pre-treatment with RhoA inhibitor in U937 cells, obviously eliminated apoptosis and caspases cleavage triggered by actein. Thus, from the data of our study, RhoA, but not caspase-3, stimulated the expression of ROCK1 in actein-triggered apoptosis.

In this study, we found that actein-induced apoptosis in human leukemia cells was observed, which was associated with PTEN and AKT activation. PTEN and AKT phosphorylation play an important role in modulating apoptosis responding to actein treatment. Accumulating evidence elucidated that PTEN, a ROCK1 substrate, is suggested to be a negative modulator of PI3K/AKT signaling pathway (40).

Consistent with previous studies, PTEN negatively regulated AKT activation during actein-induced apoptosis in our study. AKT phosphorylation was decreased by actein treatment, and its suppression using specific inhibitor abrogated apoptosis and caspase activation.

Bcl-2 family proteins, including anti-apoptotic members (Bcl-2, Mcl-1, and Bcl-xl) and pro-apoptotic members (Bax), play a crucial role in apoptosis (41,42). Bcl-2 has been known to form a heterodimeric complex with the pro-apoptotic member Bax, neutralizing its pro-apoptotic effects. Thus, the Bcl-2/Bax ratio is a decisive factor and plays a significant role in determining if cells are likely to undergo survival or death (43). Similarly, in our study, Bcl-2, Bcl-xl and Mcl-1 were highly reduced by using actein, indicating apoptosis induction and contributing to cell death. Also, the high expression of Bax in mitochondria contributes to apoptosis formation through releasing Cyto-*c* into cytoplasm (44). Also, we found that Bax was expressed highly in actein-treated U937 cells. In addition, accordingly, the release of Cyto-*c* into the cytoplasm was observed, which was in line with a previous study (45).

In conclusion, this study indicated that actein selectively triggered apoptotic response and mitochondrial damage in human leukemia cell lines, as well as in human primary leukemia cells, and showed no effects on peripheral blood mononuclear cells isolated from normal healthy humans. Furthermore *in vivo*, actein suppressed the growth of U937 tumor through activating PTEN/caspases and inactivating RhoA/ROCK1/AKT signaling pathways. Thus, we supposed that actein could be applied as a safe and effective candidate to treat human leukemia through inducing apoptosis.

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