**Astragalus** saponins modulate mTOR and ERK signaling to promote apoptosis through the extrinsic pathway in HT-29 colon cancer cells

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**Abstract.** We have previously demonstrated that the total saponins of *Astragalus membranaceus* (AST) possess potential anti-tumorigenic effects in human colon cancer cells and tumor xenografts. In the present study, the proapoptotic effects of AST were investigated in native and cytokine-induced HT-29 cells to further unveil its mechanism of action. Growth-inhibitory action of AST (60 μg/ml) was demonstrated in native HT-29 cells, which was exaggerated in tumor necrosis factor (TNF) (5 ng/ml)-induced cells. These were accompanied by caspase 3 activation, cleavage of poly(ADP-ribose) polymerase and a subsequent increase in apoptotic cell numbers. Furthermore, activation of procaspase 8 indicates that the extrinsic apoptotic pathway was involved, while cleavage of Bid into t-Bid implicates cross-talk with the intrinsic apoptotic pathway. Alternatively, AST caused G2/M phase arrest, while in cytokine-induced cells S phase arrest was predominant. Further adding to our recent suggestion on its correlation with phosphatidylinositol 3-kinase–Akt signaling, we have now revealed that AST has the potential for treating human cancers (3). The main constituents of *Radix Astragali* include flavonoids (isouqueretin, calycosin, odoratin), cycloartane triterpene saponins (astragalosides I–VIII, acetyl-astragaloside I, soyasaponin I, isoastragalosides I–II) and polysaccharides (4–7). We have recently discovered that the total *Astragalus* saponins (AST) exhibit anti-tumorigenic activities in both colon cancer cells and tumor xenografts (8). Nonetheless, the precise mechanism of AST action remains unexplored.

**Introduction**

*Radix Astragali* is the dried root of *Astragalus membranaceus* Bunge var. *mongholicus* (Dge.) Hsiao, which is found in the Shansi, Inner Mongolia, Jilin and Hebei provinces of mainland China. Many pharmacological studies have been conducted in *Radix Astragali*, among these its immunomodulatory and cardioprotective effects were most widely studied. It can suppress hypersensitivity and enhance the production of IgM antibodies (1,2). There is growing evidence that *Radix Astragali* has the potential for treating human cancers (3). The main constituents of *Radix Astragali* include flavonoids (isouqueretin, calycosin, odoratin), cycloartane triterpene saponins (astragalosides I–VIII, acetyl-astragaloside I, soyasaponin I, isoastragalosides I–II) and polysaccharides (4–7). We have recently discovered that the total *Astragalus* saponins (AST) exhibit anti-tumorigenic activities in both colon cancer cells and tumor xenografts (8). Nonetheless, the precise mechanism of AST action remains unexplored.

Failure in undergoing programmed cell death can lead to development of many diseases, including cancers (9). Mutation and deletion of apoptotic genes (e.g. p53, Bcl-2) play important roles in human carcinogenesis. As a result, apoptosis has been suggested as a potential target for cancer treatment (10). The induction of apoptosis occurs via two major pathways, the death receptor-activated (extrinsic) pathway through binding of ligands and the mitochondrion (intrinsic) pathway which is induced by several factors such as free radical or DNA damage (11). The extrinsic pathway plays an important role in the elimination of unwanted cells by the immune system, while the intrinsic pathway is important in the elimination of damaged cells that have undergone intracellular stress. Ligand binding of tumor necrosis factor (TNF)-α, TNF-related apoptosis-inducing ligand (TRAIL) or Fas-ligand to their receptors in association with adaptor molecules Fas-associated death domain (FADD) and TNF receptor-associated death domain (TRADD) proteins in the cytoplasm lead to the formation of death-inducing signaling complex (DISC) and summon (inactive pro-enzyme form of caspase 8 and sometimes...
the initiator caspase 10). Finally, executioner caspases 3, 6 and 7 are cleaved and activated (12). Poly(ADP-ribose) polymerase (PARP) is a cleaved product of caspase 3 that can be used as an indicator of apoptosis (13). It has been reported to play a pivotal role in the DNA repair mechanism (14). Besides, caspase 8 can cleave Bid after the death receptor ligation, which facilitates its post-translational N-myristoylation (15). Truncated Bid (t-Bid) is then translocated from cytosol to mitochondria, where cytochrome c is released by interaction with another proapoptotic protein Bax (in the intrinsic pathway). The cleavage of Bid is also believed to be responsible for the amplification of downstream effector events in the extrinsic apoptotic pathway (10).

In the present study, we aimed to elucidate the proapoptotic mechanism of AST by modulating a mammalian target of rapamycin (mTOR) and extracellular signal-regulated protein kinase (ERK) signaling, and proposed nuclear factor-κB (NF-κB) as the mediator involved in the apoptotic process in native or cytokine-induced HT-29 colon cancer cells.

Materials and methods

Cell culture and drug treatments. HT-29 human colon adenocarcinoma cells (ATCC, Manassas, VA) were cultured in Dulbecco’s modified Eagle medium (DMEM) supplemented with 10% heat-inactivated fetal bovine serum (FBS) and 1% penicillin-streptomycin (Gibco, Carlsbad, CA) at 5% CO2 and 37°C. Cells were seeded at a density of 3-4x10^4 overnight, and then pre-incubated with 60 μg/ml of AST for 1 h, with or without subsequent addition of 5 ng/ml TNF-α (cytokine-induced cells). All cells were harvested at 12, 24, 48 or 72 h after AST treatment. Extraction of AST from crude Radix Astragali was described previously (8). Chemicals were obtained from Sigma-Aldrich (St. Louis, MO) unless specified otherwise.

MTT cell viability assay. Cells were seeded at a density of 3x10^3 in 96-well plates. MTT assay was conducted at 12-72 h after drug treatment. 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazoliumbromide (MTT) solution was added to wells and incubated for 3 h. Absorbance was determined spectrophotometrically at 540 nm.

Western immunoblotting. Cells were harvested and centrifuged. Cell pellets were collected and washed with ice-cold PBS (137 mM NaCl, 2.68 mM KCl, 8.1 mM Na2HPO4, 1.47 mM KH2PO4, pH 7.4). They were then lysed in RIPA buffer [0.5% Nonidet P-40, 0.1% SDS, 0.1% Triton X-100, 10% glycerol, 50 mM Tris, pH 7.5, 150 mM NaCl, 2 mM EDTA, with 1 mM phenylmethyl-sulfonyl fluoride (PMSF) and 0.5 μg/ml aprotinin] on ice for 15 sec. Supernatant containing cytosolic protein was removed. Solution in wells was aspirated and dimethyl sulfoxide (DMSO) was added. Absorbance was measured spectrophotometrically at 540 nm.

Immunofluorescence nuclear staining. Cells were seeded in 35-mm dishes and treated as stated above. They were washed with PBS twice and fixed in 4% paraformaldehyde for 10 min. After washing with PBS three times, cells were permeated with ice-cold 75% methanol at -20°C for another 5 min. Cells were then stained with DNA binding dye bis Benzimide H33342 trihydrochloride for 10 min. Nuclear morphology was observed under a fluorescence microscope (NanoDrop Technologies, Inc., Wilmington, DE) with magnification of x400. For detection of NF-κB nuclear translocation, cells were washed with PBS twice and blocked with 0.2% BSA at room temperature. After three washes with PBS, rabbit polyclonal antibody against p65 (1:100 dilution) were added and incubated overnight at 4°C. Following three washes of PBS, anti-rabbit secondary antibody was added and incubated for 1 h at room temperature. Immunofluorescence images were observed under fluorescence microscope with magnification of x400.

Electrophoretic mobility shift assay (EMSA). Drug-treated cells were harvested and resulting pellets were collected and washed with ice-cold PBS twice. They were then re-suspended in 700 μl of Buffer A [10 mM HEPES, pH 7.9, 10 mM KCl, 0.1 mM EDTA, 0.1 mM EGTA, 1 mM dithiothreitol (DTT) and 0.5 mM PMSF] and incubated on ice for 15 min. One percent Nonidet P-40 (5 μl) was then added and vortexed for 15 sec. Supernatant containing cytosolic protein was removed by centrifugation at 15,000 x g for 15 min. Buffer C (20 mM HEPES, pH 7.9, 0.4 M NaCl, 1 mM EDTA, 1 mM EGTA, 1 mM DDT and 1 mM PMSF) was then added to re-suspend pellets and incubated on ice for 30 min to extract nuclear proteins. Supernatant was collected by centrifugation at 15,000 x g for 15 min. Protein concentration was measured according to the protein assay mentioned above. For labeling the NF-κB double-stranded oligonucleotides, a probe with binding sites for NF-κB was used, with the following sequence, 5’-AGTTGAGGGGACTTTCCCAGG-3’. The probe was incubated with T4 polynucleotide kinase buffer, γ-32P-ATP (3,000 Ci/mmol at 10 μCi/ml) and T4 polynucleotide kinase (Invitrogen, Carlsbad, CA) at 37°C for 10 min. Afterwards, 0.5 M EDTA was added followed by adding TE buffer. The probe was then purified by passing through the column (MicroSpin™ G-25 column, UK) twice and centrifuged at 1,000 x g for 2 min at 4°C, before being stored at -20°C. To perform the gel retardation assay, 10 μg of the extracted nuclear protein was incubated with the reaction buffer for 15 min, followed by addition of the 32P end-labeled oligonucleotide. Reaction mixture was separated in a non-denaturing 6% polyacrylamide gel. The gel was dried and exposed to radiographic film at -70°C.
Flow cytometry. After drug treatment for 12, 24, 48 or 72 h, cells seeded at a density of 4x10^4 were collected by trypsinization with ice-cold PBS. Cells were then fixed with ice-cold 75% ethanol overnight at -20°C. Cell pellets were collected by centrifugation at 1,000 x g for 10 min at 4°C the following day. Cells were then incubated with the master mix [0.1% Triton X-100, 100 μg/ml RNase A and 50 μg/ml propidium iodide (PI) in PBS] for 1 h on ice for flow cytometry (FACS Canto™) (BD Biosciences, Franklin Lakes, NJ). Cell distribution at different phases of the cell cycle with different treatments was determined by using PI staining.

Statistical analysis. Data are presented as mean ± SEM and analyzed by using one-way analysis of variance (ANOVA), followed by an LSD post-hoc test (SPSS version 10.0; Chicago, IL). The level of statistical significance was established as P<0.05.

Results

**AST induces growth inhibition and apoptosis in native and cytokine-induced HT-29 cells.** Survival rate of cells can be determined by the percentage of cell viability. There was a decrease in percentage of viable cells when cells were treated with AST in native or cytokine-induced cells (Fig. 1). There was no effect on cell viability when cells were treated with TNF-α alone. AST also exhibited a time-dependent growth inhibition in HT-29 cells, which was more pronounced in cytokine-induced cells.

Morphological change is an important feature that indicates apoptotic cell death. During apoptosis, the cell shrinks while the nucleus condenses and fragments, which can be observed by conducting immunofluorescence staining. Following AST treatment, cells became less rounded, shrunken in size and sometimes fragmented, with the observation of nuclear chromatin condensation (Fig. 2A). There was a significant increase in the number of cells with condensed nuclei caused by AST in cytokine-induced cells (Fig. 2B).

**AST modulates the extrinsic apoptotic cascade in native and cytokine-induced HT-29 cells.** Procaspase 8 is the inactive pro-enzyme form of caspase 8. After the ligation of death receptor, procaspase 8 cleaves to activate caspase 8 as manifested by reduced procaspase 8 expression. Our results show that AST induced downregulation of procaspase 8 expression after 48 and 72 h of treatment (Fig. 3A). This effect became more prominent in cytokine-induced cells, of which the phenomenon was observed as early as 12 h. Cleavage of procaspase 3 was designated to indicate the activation of caspase 3, the key executioner caspase. The expression of procaspase 3 was shown to be decreased by
AST treatment. The effect was more obvious after 48 h and intensified in cytokine-induced cells. These events then resulted in increased PARP cleavage (Fig. 3B).

Bid is a pro-apoptotic BH-3 protein that belongs to the Bcl-2 family. After activation of caspase 8, Bid is cleaved into truncated Bid (t-Bid) in order to amplify downstream extrinsic apoptotic pathway. The expression of intact Bid is thereby decreased during apoptosis. Our results have demonstrated that formation of t-Bid began after 48 h of AST treatment, while a clear increase in expression of t-Bid was observable at 72 h. Such reduction became more significant in cytokine-induced cells (Fig. 3C). In addition, the time of t-Bid formation actually matches that of the AST-evoked caspase 8 activation.

AST induces phase-specific cycle arrest in native and TNF-treated HT-29 cells through modulation of mTOR signaling.

HT-29 colon cancer cells were treated with AST for 12, 24, 48 or 72 h. Cells were accumulated at S phase and arrested at G2/M phase after AST treatment for 24 and 48 h (Fig. 4). Also, S phase arrest resulted when treated with TNF-α alone. AST further increased the percentage of cells accumulated at the S phase in cytokine-induced cells (when compared to treatment of TNF-α alone). Nevertheless, cell cycle arrest was not sustained after drug treatment for 72 h, when apoptosis was predominant.

mTOR is an intracellular nutrient sensor that controls protein synthesis, cell growth and metabolism. Inactivation of mTOR can induce autophagy. When mTOR is inactivated, its protein expression decreases. Our data show a reduction in p-mTOR expression following 12-72 h of AST treatment, with no alteration of mTOR expression (Fig. 5A). A similar observation was found in cytokine-induced cells, with no obvious synergistic/additive effect. AST could have inhibited mTOR to suppress growth and survival of colon cancer cells. PTEN is a tumor suppressor gene that is involved in the mTOR signaling pathway. Loss or mutation of PTEN can result in the development of many types of cancers. It has been reported that overexpression of PTEN stimulates autophagy. Our results indicate that expression of PTEN increased after AST treatment, which was slightly increased in cytokine-induced cells (Fig. 5A).

AST inhibits NF-κB/DNA binding activity in native and TNF-treated HT-29 cells that are associated with ERK activation.

The NF-κB signaling pathway was found to be correlated with different types of cancer including that of the colon. It is also an important transcription factor that participates in colonic inflammation. We were interested to determine if AST affects NF-κB signaling in HT-29 cells. NF-κB/DNA-binding activity in HT-29 cells was decreased after treatment of AST for 3-6 h, with further reduction in cytokine-induced cells. Nevertheless, the reduction began to level off after 24 h of drug treatment (Fig. 6A). In addition, both bis-benzimide H33342 stain and anti-p65 antibodies were added to HT-29 cell culture to investigate the intracellular localization of p65 following drug treatments. Under an unstimulated state, p65 was localized mainly in the cytoplasm with a visible red color. Upon stimuli
Figure 4. Cell distribution of HT-29 colon cancer cells in different phases of the cell cycle after 12, 24, 48 or 72 h of drug treatments. The percentage of cells in the S and G2/M phases was shown to be increased after AST treatment for 24 or 48 h (with concurrent reduction in the percentage of cells in the G1 phase). Cells treated with TNF-α caused cell arrest in the S phase, which became more prominent when AST was co-treated. However, cells were unable to remain arrested after 72 h. Results are expressed as percentage of cells in each phase of the cell cycle in respect to the untreated control. Bars represent means ± SEM of 2-3 experiments. *P<0.05; **P<0.01, significantly different from the corresponding control.

Figure 5. AST regulates mTOR, PTEN and ERK in HT-29 cells. HT-29 cells were treated with AST (A), TNF-α (T) or AST + TNF-α (A+T) for 12, 24, 48 or 72 h. (A) AST downregulated phosphorylated mTOR expression without affecting mTOR and caused PTEN protein overexpression. These phenomena were not intensified in cytokine-induced cells. (B) Activation of ERK-1 and ERK-2 occurred after 12-72 h exposure to AST. The effect was boosted in cytokine-induced cells after 72 h of AST incubation.
Figure 6. AST reduced NF-κB/DNA-binding activity in native and cytokine-induced HT-29 cells. (A) Effects of AST and/or TNF-α on NF-κB/DNA-binding activity after 3, 6, 12, 24, 48 and 72 h of drug treatments in HT-29 cells were determined by EMSA. Concentrations of AST (A) and TNF-α (T) were 60 μg/ml and 5 ng/ml, respectively. Control cells (C) were treated with culture medium alone without any drug treatment. NF-κB/DNA-binding activity in HT-29 cells decreased prominently after incubated with AST for 3-12 h in both native and cytokine-induced cells, and the effects began to diminish thereafter. (B) HT-29 cells were treated with either 60 μg/ml of AST or 5 ng/ml of TNF-α or both for 12 h. Cells were fixed and the intracellular location of the NF-κB p65 subunit was determined by immunofluorescence staining using anti-p65 antibody. Pictures were taken with a fluorescence microscope (x400 magnification). Results show p65 nuclear translocation following AST treatment, which would be intensified in cytokine-induced cells. Similar effects were also observed at other time points but were most apparent at 12 h.
such as proinflammatory cytokine p65 would be translocated to the nucleus. The co-localization of red and blue (H33342) colors indicates the presence of apoptotic bodies with p65 nuclear translocation after 3 h of AST treatment. This phenomenon was intensified in cytokine-induced cells. The effect was most obvious after 12 h of AST incubation (Fig. 6B). This conflicts with the reduced NF-xB/DNA-binding activity that must be clarified later.

Increasing evidence has indicated that NF-xB activation is related to MAPK family members such as ERK. The effects of AST and TNF-α on the activation of ERK was examined as a possible signaling pathway. The expression of total ERK remains unchanged, while expression of both isoforms of phosphorylated ERK was upregulated after treatment with AST for 12-72 h, which was exaggerated in cytokine-induced cells (Fig. 5B). The continuous activation of ERK could have contributed to the proapoptotic activity of AST in colon cancer cells, which may be facilitated by earlier decrease in NF-xB/DNA-binding activity.

Discussion

*Radix Astragali* is a tonifying herb commonly used in traditional Chinese Medicine. It has been shown to possess anti-proliferative effect on different cancer cell lines, including that of the colon (16). *Astragalus* saponins are one of the major groups of chemical constituents. We have demonstrated in a recent report that the total *Astragalus* saponin AST exerts antitumorigenic and proapoptotic actions in colon cancer cells and tumor xenografts (8). In the present study, we investigated the possible involvement of the mammalian target of rapamycin (mTOR) and extracellular signal-regulated protein kinase (ERK) signaling pathways in the proapoptotic effects of AST in HT-29 colon cancer cells, focusing on the extrinsic apoptotic pathway. Recent studies have reported that caspase 8 is involved in the anti-tumor mechanism of SC-1 in Hep 3B cells, propolin A and B in human melanoma cells and RRR-α-tocopherol in human breast cancer cells (14,17,18). Furthermore, another study also showed that colon carcinoma cells with mutations of caspase 8 are less sensitive to drug-induced apoptosis, of which restoration of caspase 8 expression could sensitize the resistant tumor cells for apoptosis (19). All these studies have suggested that caspase 8 plays an important role in the anti-carcinogenic mechanism. Procaspase 8 cleavage and subsequent activation of the downstream executioner caspase 3 were revealed after AST treatment, which resulted in drug-induced apoptosis. In addition, cleavage of the caspase 3 substrate PARP would facilitate cellular disassembly, leading to apoptosis (20). In contrast, following the ligation of the death receptor, caspase 8 cleaves intact Bid into its active form t-Bid, which in turn triggers the release of Bax to amplify downstream effector events in the intrinsic pathway by initiating release of apoptotic factors (e.g. cytochrome c) from mitochondria into the cytoplasm. Otherwise stated, Bid serves a key role in linking the intrinsic and extrinsic pathways to strengthen the apoptotic effects. Study of Zhou et al has re-confirmed this hypothesis, by indicating that Bid is important in the andrographolide-induced apoptotic signaling pathway of human cancer cells, as it relates the cell death signaling of caspase 8 to the mitochondria (21). In fact, we have already reported that AST stimulates the intrinsic apoptotic pathway in human cancer cells (22).

mTOR is a 289-kd phosphatidylinositol-3-kinase (PI3K)-related kinase which functions as an intracellular nutrient sensor that control protein synthesis, cell growth and metabolism. Dysregulation of mTOR signaling is involved in many human cancers such as carcinomas of lung, bladder, stomach and lymphomas. This suggests mTOR as a target of novel therapy for human cancers (23,24). There are two mTOR complexes, mTOR complex 1 (mTORC1) and mTOR complex 2 (mTORC2), where mTORC1 is responsible for nutrient sensing signals and mTORC2 is implicated in actin organization and phosphorylates the Akt protein kinase B (23). Activation of Akt by mTORC2 drives several cancer-related response including increase in cell proliferation, growth and cell migration and a shift to glycolytic metabolism. Upstream regulation of mTOR will give rise to human diseases characterized by hamartomas with a variable propensity to progress into neoplasia (25). One of the essential upstream regulators for mTORC1 activation involves stimulation of Class I PI3Ks by members of the receptor tyrosine kinase family, which triggers the conversion of phosphatidylinositol-4,5-biphosphate (PIP2) to the bioactive messenger phosphatidylinositol-3,4,5-triphosphate (PIP3). This in turn causes subsequent activation of the serine-threonine kinase Akt (protein kinase B). Nevertheless, mTOR signaling has a dual role in the PI3K-Akt pathway, of which on the other hand mTORC2 is an upstream activator of Akt, while on the other hand mTORC1 is a downstream recipient of PI3K-Akt regulation. We have recently discovered that AST induces anti-carcinogenic effect at least partly by modulating PI3K-Akt signaling (26). It is of interest to see if the upstream (e.g. PTEN) and downstream (e.g. mTOR from the mTORC1 complex) factors will also be regulated by AST. In this study, expression of mTOR was shown to be decreased in AST-treated cells, which implicates that antitumor action of AST may also act through inhibition of cell proliferation which is regulated by the mTOR signaling pathway. PTEN (phosphatase and tensin homolog deleted on chromosome ten) is a tumor suppressor gene that is ubiquitously expressed in human tissues. A deficiency of PTEN leads to hyperactivation of the PI3K-Akt pathway, which is the cause for many cancers (25). It is one of the most frequently mutated tumor-suppressor genes and is impinging upon mTOR signaling. Inhibition of cell migration and invasion by directly dephosphorylating two key tyrosine-phosphorylated proteins is one of the functions of PTEN. PTEN also plays a key role in dephosphorylating a key signal transduction lipid. This lipid signal transduction pathway can protect tumor cells from apoptosis in the absence of PTEN. Thus, PTEN negatively regulates cell interactions with the extracellular matrix and maintains cell sensitivity to apoptosis. Recent studies have shown that restoration of PTEN increases the rate of apoptosis in glioma cells and induces apoptosis in breast cancer cell lines even in the presence of cell attachment (27,28). A previous study has proven that overexpression of PTEN stimulates autophagy in colon cancer cells (29). Upregulation of PTEN following AST-treatment has been demonstrated in our immunobots.

The nuclear factor-xB (NF-xB) pathway is involved in the regulation of key cellular processes such as proliferation, stress response, innate immunity and inflammation. Mutations and
misregulation of NF-κB signaling have been implicated in a variety of cancers (30). It activates proliferation and inhibits apoptosis by activating modulators of cytokines (e.g. TNF-α), growth factors, survival genes, angiogenic factors and inflammatory genes (31,32). The NF-κB/REL family of transcription factors is comprised of Rel A/p65, c-Rel, Rel B, P105/NF-κB1 and p100/NF-κB2, whereas the most common combination is the p65/p50 heterodimer (33). NF-κB has been demonstrated to inhibit TNF-α induced cell death. TRAIL-induced apoptosis in renal cancer can be prevented by activation of this transcription factor. Therefore, chemotherapeutic agents that activate NF-κB can accounted for the resistance of colon cancer cells to cytotoxic challenges (34,35). Caspases can play an indirect role in the regulation of NF-κB activation, since NF-κB is substrate for caspase cleavage. Caspase-3-mediated cleavage of p65, p50 and c-Rel can be induced upon TNF-α activation (36). Immunofluorescence staining of HT-29 cancer cells in our study showed nuclear condensation in AST- and TNF-α-treated cells, being most prominent at 72 h. However, the manifestation of p65 nuclear translocation during early inhibition of NF-κB/DNA binding activity (~12 h) cannot be explained and the discrepancy remains to be clarified. NF-κB activity is mainly controlled by IκB inhibitor proteins. Inflammatory signals initiate the degradation of IκB by stimulating phosphorylation of serines-32 and -36 (37). It has been shown that the ERK induces site-specific phosphorylation of IκB-α in HeLa cells and directly activates the IKK complex (38). In contrast, the ERK signaling pathway involves a cascade of phosphorylation events and is generally regarded as cell proliferation- and differentiation-related (39). However, the role of ERK is controversial. Some stimuli may act through the ERK pathway to cause apoptosis, although the exact mechanism of ERK-caused apoptosis has yet to be elucidated. It has been suggested that transient ERK activation may be linked to cellular proliferation while strong and persistent activation may lead to programmed cell death (40). It is also proposed that ERK will mediate NF-κB during its proapoptotic action. Our data show that protein expression of both phosphorylated ERK will mediate NF-κB signaling and the resulting apoptotic process.

Collectively, findings from our study indicate that AST induces growth-inhibition in HT-29 cells by activation of the extrinsic apoptotic cascade as well as facilitation of phase-specific cell cycle arrest. These events appear to be modulated by both mTOR and ERK signaling, of which NF-κB has played a role in the proapoptotic activities of the latter pathway. In most of the afore-mentioned processes, the effects of AST could be amplified in cytokine-induced cells.

References