# Adenine causes cell cycle arrest and autophagy of chronic myelogenous leukemia K562 cells via AMP-activated protein kinase signaling

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Abstract. AMP-activated protein kinase (AMPK) is known as a pivotal regulator of cellular metabolism. Mounting evidences have demonstrated that AMPK activation exerts tumor suppressive activity on leukemia cells. The present study reported that adenine, an AMPK activator, triggers cell cycle arrest and autophagy of human chronic myelogenous leukemia K562 cells consequently suppressing cell viability. The present findings revealed that adenine treatment (4.0-8.0 mM) significantly inhibited the viability of K562 cells to  $69.3 \pm 2.5\%$ (24 h) and  $53.4\pm2.1\%$  (48 h) of the control. Flow cytometric analysis revealed that there was a significant accumulation in G<sub>2</sub>/M phase, but not sub-G<sub>1</sub> phase K562 cells following exposure to adenine. Additional investigation demonstrated that adenine treatments significantly increased the number of acidic vesicular organelles and the level of autophagosomal microtubule associated protein 1 light chain 3  $\alpha$  (LC3) marker. By contrast, cleavage of caspase-9, caspase-3 and poly-ADPribose polymerase was insignificantly affected in K562 cells following adenine treatment. In K562 cells, adenine was able to markedly promote the phosphorylation of AMPKa and suppress the phosphorylation of mammalian target of rapamycin (mTOR), a downstream target of AMPK. In addition, inhibiting AMPK phosphorylation using dorsomorphin restored mTOR phosphorylation, inhibited the accumulation of LC3 and significantly recovered the suppressed cell viability in response to adenine. Taken together, the present results demonstrated that adenine induced  $G_2/M$  phase arrest and autophagic cell death, consequently suppressing the viability of K562 cells, which may attribute to the AMPK activation triggered by adenine. These findings provide evidence that adenine may be beneficial to chronic myelogenous leukemia therapy by suppressing excessive cell proliferation.

## Introduction

Chronic myelogenous leukemia (CML) is a chronic myeloproliferative disorder, which causes uncontrolled growth of immature myeloid cells (1). The BCR RhoGEF and GTPase activating protein (BCR)-ABL proto-oncogene 1 nonreceptor tyrosine kinase (ABL) gene rearrangement is the main characteristic of CML, which expresses the oncogenic fusion protein BCR-ABL (2). BCR-ABL is a constitutively active tyrosine kinase, which activates multiple signaling pathways and consequently promotes malignant transformation, including uncontrolled cell proliferation (3), abnormal cell adhesion (4), and resistance to typical apoptotic inducer anti-leukemic drugs (5,6). Thus, formation of the BCR-ABL fusion gene serves an essential role in the pathogenesis of CML (7). Previously, imatinib, a specific ABL kinase inhibitor, was established as the standard treatment for CML (8).

In addition to targeting BCR-ABL kinase, previous studies have revealed that several pathways are important for CML cell survival, which may be potential targets for developing novel anti-leukemia drugs (9,10). Among these pathways, AMP-activated protein kinase (AMPK) signaling has been reported to possess anti-leukemia activity (11). AMPK serves an important role in energy metabolism in response to changes in cellular fuel levels (12). Furthermore, mounting evidences have suggested that AMPK could be a target for tumor prevention and treatment (7). Previous studies have revealed that AMPK activators exhibit anti-leukemia effects in CML cells by triggering apoptosis and autophagy (3,4). Thus, the

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identification and characterization of AMPK activators is important, and beneficial for the development of potential anti-leukemia drugs.

The present study aimed to investigate whether adenine, a purine compound that induces AMPK activation, exhibits anti-leukemia effects on human CML cells. Furthermore, the underlying mechanism, with emphasis on AMPK signaling was evaluated. The results revealed that adenine suppressed cell viability of K562 cells and induced accumulation of G0/G1 phase cells. In parallel, it was observed that adenine triggered K562 cells autophagy. Finally, it was demonstrated that suppressed cell viability, accumulation of G<sub>0</sub>/G<sub>1</sub> phase cells and induction of autophagy were associated with AMPK activation in response to adenine.

#### Materials and methods

Reagents. All chemicals were obtained from Sigma-Aldrich (Merck KGaA, Darmstadt, Germany) unless otherwise specified. RPMI-1640 medium and fetal bovine serum (FBS) were purchased from Invitrogen (Thermo Fisher Scientific, Inc., Waltham, MA, USA). Antibodies against  $\beta$ -actin, caspase-3, caspase-8, phosphorylated (p)-AMPK (T172; cat. no. 2535), AMPK (cat. no. 2532), poly-ADP-ribose polymerase (PARP) (cat. no. 9532), cell division cycle (Cdc)25 (cat. no. 3652), cyclin-dependent kinase (CDK)2 (cat. no. 2546), CDK4 (cat. no. 12790), CDK6 (cat. no. 13331), cyclin B (cat. no. 4135), cyclin E (cat. no. 20808), Wee1-like protein kinase (Wee1) (cat. no. 13084), autophagy protein (Atg)5 (cat. no. 12994), beclin-1 (cat. no. 3495) and microtubule associated protein 1 light chain  $3 \alpha$  (LC3) (cat. no. 3868) were purchased from Cell Signaling Technologies, Inc. (Danvers, MA, USA). Anti-human glyceraldehyde 3-phosphate dehydrogenase (GAPDH) (cat. no. ab9485), horseradish peroxidase-conjugated anti-mouse IgG (cat. no. ab6789) and anti-rabbit IgG (cat. no. ab6721) antibodies were purchased from Abcam (Cambridge, UK).

Cell culture and experimental treatments. The human CML K562 cell line (CCL-243<sup>TM</sup>) was obtained from the American Type Culture Collection (Manassas, VA, USA) and maintained in RPMI-1640 medium supplemented with 10% FBS at 37°C in a humidified atmosphere with 5% CO<sub>2</sub>. Cells at the density of  $5\times10^5$  cells/ml were collected and were passaged twice prior to being subcultured T-75 flasks (Corning Incorporated, Corning, NY, USA) for subsequent treatments.

For treatments, cells were seeded in 6-well plates at an initial density of  $1 \times 10^5$  cells/ml and grown to ~80% confluence at 37°C. Treatments were performed by incubating cells with various concentrations of adenine (0.5, 1.0, 2.0, 4.0 or 8.0 mM) in serum-free RPMI-1640 for 16 h. AMPK inhibition was performed by pretreating cells with 5  $\mu$ M dorsomorphin (cat. no. P5499; Sigma-Aldrich; Merck KGaA) for 2 h, and then incubating the pretreated cells with 4.0 mM adenine for 24 h or 48 h. Following treatment, the cells were washed with PBS (25 mM sodium phosphate, 150 mM NaCl, pH 7.2) and collected for subsequent analyses.

*Cell viability assay.* Cell viability was assessed using an MTT assay protocol as previously described (13) in the absence or presence of adenine. After 24 or 48 h treatments, cells were



Figure 1. Adenine suppresses the viability of K562 cells. Cells were treated with various concentrations of adenine (0.5, 1.0, 2.0, 4.0 or 8.0 mM) for 24 or 48 h, and then subjected to a cell viability assay. Data are expressed as the mean  $\pm$  standard error of the mean following three independent experiments. \*P<0.05 and \*\*P<0.01 as compared with the untreated control. C, control.

harvested, and then incubated with MTT (0.5 mg/ml) at 37°C for 4 h. The cell viability was directly proportional to the production of formazan, which was dissolved in isopropanol and determined by measuring the absorbance at 570 nm using a microplate reader (SpectraMAX 360 pc; Molecular Devices, LCC, Sunnyvale, CA, USA).

Cell cycle distribution analysis. Cells were starved for 16 h in serum-free medium, and then cultured in fresh serumcontaining medium at 37°C for 24 h to allow cell-cycle progression. Following different treatments, cells were fixed and analyzed by flow cytometry to determine cell cycle distribution. At the end of each treatment, cells were collected, fixed with 1 ml of ice-cold 70% ethanol on ice for 30 min, incubated at -20°C for 24 h and centrifuged at 380 x g at 4°C for 5 min. Cell pellets were reacted with 1 ml cold staining solution containing 20 µg/ml propidium iodide, 20 µg/ml RNase A and 1% Triton X-100, incubated at 4°C for 15 min in the dark, and then analyzed using a FACS Calibur system with CellQuest<sup>TM</sup> software (version 2.0) (both BD Biosciences, Franklin Lakes, NJ, USA). Representative results were acquired from three independent experiments.

Western blot analysis. Cells were washed with PBS and then incubated with lysis buffer (50 mM Tris-HCl, pH 7.5, 1% Nonidet P-40, 1 mM phenylmethylsulfonyl fluoride and 1 mM NaF) containing complete protease inhibitor cocktail (Roche Applied Science, Mannheim, Germany) for 1 h at 4°C. The resulting supernatants were collected for protein ßquantitation using a bicinchoninic acid kit (Pierce; Thermo Fisher Scientific, Inc.). Crude proteins (20  $\mu$ g) were subjected to 12.5% SDS-PAGE, transferred to a nitrocellulose membrane (EMD Millipore, Billerica, MA, USA) and then incubated with 5% w/v skimmed milk/PBS at room temperature for 1 h to block nonspecific binding. The blocked membrane was then incubated with primary antibodies (1:1,000 dilution) at 25°C for 2 h, followed by incubation with peroxidase-conjugated



Counts



Adenine (mM)

Figure 2. Effects of adenine on the cell cycle distribution of K562 cells. Cells were treated with 1.0, 2.0 or 4.0 mM adenine for 24 h, and then subjected to flow cytometric analysis for determining cell cycle phases. The ratio of each phase is presented as a percentage. Data are expressed as the mean  $\pm$  standard error of the mean following three independent experiments. \*P<0.05 and \*\*P<0.01 as compared with the untreated control.

anti-IgG secondary antibodies (1:2,000 dilution) at 25°C for 1 h. Detailed information for antibodies were described in the reagents. The blocked membrane without incubation with primary antibodies was also incubated with secondary antibodies for specificity test, and a nonsignificant signal was detected as the control. Signal development was performed using an enhanced chemiluminescence reagent (EMD Millipore). Luminescent signals were acquired and quantified using an image analysis system (LAS-4000 with Image Reader LAS-4000 version 2.1, Fuji, Tokyo, Japan). GAPDH was used as an internal control.

Quantification of acidic vesicular organelles (AVOs) using acridine orange staining and flow cytometric analysis. Cells (1x10<sup>4</sup>) were incubated with acridine orange at 25°C for 17 min, harvested with trypsin-EDTA, and then collected in phenol red-free growth medium (Thermo Fisher Scientific, Inc.). Green (510-530 nm) and red (650 nm) fluorescence emission from cells illuminated with blue (488 nm) excitation light was analyzed using FACS Calibur system using CellQuest software as mentioned above. With acridine orange staining, bright green and faint red fluorescence was observed in the cytoplasm and nucleolus, respectively, and bright red fluorescence was observed in acidic compartments (2,14). Since the intensity of the red fluorescence is proportional to the degree of acidity, the volume of the cellular acidic compartment could be quantified (15).

Statistical analysis. Data are expressed as the mean  $\pm$  standard error of the mean following three independent experiments. Statistical analysis was performed by using SigmaStat (version 4.0, Systat Software, Inc, San Jose, CA, USA). Statistical significance analysis was determined using one-way analysis of variance followed by Dunnett's test for multiple comparisons with the control. P<0.05 was considered to indicate a statistically significant difference.



Figure 3. Adenine inhibits cell cycle regulators but not caspase effectors. Cells were treated with 1.0, 2.0 and 4.0 mM adenine for 24 h, and then lysed for determination of protein expression levels using western blot analysis. GAPDH was used as an internal control. The apparent molecular weights of the detected proteins are indicated. PARP, poly-ADP-ribose polymerase; CDK, cyclin-dependent kinase; Cdc25A, cell division cycle 25A; Wee1, Wee1-like protein kinase; C, control.



Figure 4. Adenine induces autophagy in K562 cells. Cells were treated with 1.0, 2.0 or 4.0 mM adenine for 24 h, and then subjected to (A) western blot analysis for determination of protein levels, or (B) acridine orange staining for detection of AVOs. GAPDH was used as internal control. The apparent molecular weights of the detected proteins are indicated. Quantitative data are expressed as the mean  $\pm$  standard error of the mean following three independent experiments. \*P<0.05 and \*\*P<0.01 as compared with the untreated control. AVO, acidic vesicular organelle; Atg5, autophagy protein 5; LC3, microtubule associated protein 1 light chain 3  $\alpha$ ; C, control.

### Results

Adenine suppresses the viability of human CML K562 cells. To investigate the effect of adenine on human CML K562 cells, cell viability was determined using MTT assay. As presented in Fig. 1, for 24 h treatments, the viability of K562 cells was significantly reduced to  $81.5\pm2.1$ ,  $75.4\pm2.2$  and  $69.3\pm2.5\%$  of the control in response to 2, 4 and 8 mM adenine, respectively (all P<0.05). For 48 h treatments, the cell viability was further reduced to  $61.3\pm2.3$  and  $53.4\pm3.1\%$  of the control in response to 4 and 8 mM adenine, respectively (both P<0.01). These findings revealed that adenine at concentrations of 2, 4 and 8 mM significantly inhibited the viability of K562 cells.

Adenine induces  $G_2/M$  cell cycle arrest and alters cell cycle regulators in human CML K562 cells. To explore the inhibited cell viability in response to adenine, the cell cycle distribution of K562 cells treated with adenine was determined using flow cytometric analysis. As presented in Fig. 2, the

sub-G<sub>1</sub> phase ratio was increased by  $4.1\pm0.7\%$  in K562 cells treated with 4 mM adenine compared with the control group (P<0.05). In addition, the G<sub>2</sub>/M phase ratio was significantly increased in K562 cells treated with 2 mM (32.1±2.7%) and 4 mM (43.8±2.1%) adenine compared with that in the untreated control group (both P<0.01). The G<sub>0</sub>/G<sub>1</sub> phase ratio was significantly decreased in K562 cells treated with 2 mM (44.6±2.7%) and 4 mM (33.8±2.4%) adenine compared with that in untreated cells (both P<0.05).

Since cell viability and cell cycle distribution were altered following adenine treatment, the effects of adenine on caspase signaling cascades, and cell cycle regulators were then investigated. As presented in Fig. 3A, cleavage of caspase-8, caspase-3 and PARP was insignificantly affected by adenine treatments (1.0, 2.0 or 4.0 mM). Regarding cell cycle regulators, the protein expression levels of Cyclin D1, CDK4, CDK6 and Cdc25A were markedly decreased in response to adenine treatments (Fig. 3B). By contrast, the level of Wee1 was elevated upon adenine stimulation. Notably, Cyclin E protein expression





Figure 5. AMPK activation is involved in autophagy and suppression of viability of K562 cells in response to adenine. Cells were (A) treated with adenine alone or (B) pretreated with AMPK inhibitor dm (5  $\mu$ M) for 2 h followed by treatment with adenine for 24 h, and then subjected to western blot analysis. GAPDH was used as internal control. The apparent molecular weights of the detected proteins are indicated. Quantitative data are expressed as the mean  $\pm$  standard error of the mean following three independent experiments. (C) A cell viability assay was performed to investigate the effects of dm on K564 cells. \*P<0.05 and \*\*P<0.01 as compared with the 4.0 mM adenine treatment group. AMPK, AMP-activated protein kinase; p, phosphorylated; mTOR, mechanistic target of rapamycin; Atg5, autophagy protein 5; dm, dorsomorphin; C, control.

was insignificantly affected by adenine. Taken together, these findings suggest that adenine treatment induces  $G_2/M$  cell cycle arrest without involvement of the activation of caspase signaling cascades and consequent apoptosis of K562 cells.

Adenine induces autophagy of human CML K562 cells. To determine whether adenine suppresses cell viability via autophagic cell death, induction of autophagy in K562 cells was then examined. Western blot analysis demonstrated that adenine markedly upregulated the protein expression of Beclin-1 and Atg5, and promoted the conversion of LC3-I to LC3-II in the cells (Fig. 4A). In addition, using acridine orange staining and flow cytometric quantification, adenine was identified to significantly increase the number of AVOs in K562 cells in a dose-dependent manner (Fig. 4B). These findings suggest that adenine treatment promotes the formation of autophagosomes in K562 cells.

Adenine induces the phosphorylation of AMPK $\alpha$ , contributing to inhibition of mTOR and autophagic signaling cascades in K562 cells. Adenine has been reported to exert antitumoral activity on human renal carcinoma 786-O cells via AMPK activation (16). Thus, the roles of AMPK signaling in autophagic cell death of K562 cells in response to adenine were investigated. As presented in Fig. 5A, adenine markedly enhance the phosphorylation of AMPK $\alpha$  at T172, reduced the phosphorylation of mTOR at S2448 and elevated the protein expression level of autophagic marker Atg5 compared with untreated cells. By contrast, pretreatment with the AMPK inhibitor dorsomorphin (dm) reversed the expression changes to p-AMPKa, p-mTOR and Atg5 in K562 cells following exposure to adenine (Fig. 5B). In addition, pretreatment with dm significantly restored the viability of K562 cells suppressed by 4.0 mM adenine (Fig. 5C). Collectively, these findings suggest that the AMPK/mTOR signaling pathway is involved in adenine-induced autophagy and consequent viability inhibition of K562 cells.

### Discussion

Clinically, the Philadelphia chromosome exists in  $\sim 95\%$  of adults with CML and expresses the BCR-ABL fusion

protein with uncontrolled tyrosine kinase activity, resulting in abnormal regulation of downstream signaling pathways (17). The aberrant activation of signaling pathways, including the RAS/mitogen activated protein kinase kinase/extracellular signal-related kinase and phosphoinositide 3-kinase/RAC-alpha serine/threonine-protein kinase pathways, may promote cell proliferation, diminish cell apoptosis or contribute to growth factor independence of cancer cells (18,19). Therefore, tyrosine kinase inhibitors targeting the BCR-ABL fusion protein, including imatinib, dasatinib and nilotinib, are used for CML treatment, and have demonstrated improved outcomes in patients with CML (20). However, high relapse rates, drug resistance and high mortality rates associated with transplantation remain challenges for CML treatment (21).

Previous studies have revealed that AMPK activators can suppress several types of tumors, including pancreatic, bladder and prostate cancer, through AMPK-dependent apoptosis (19-21). In addition, AMPK has been reported as a potential target that regulates various signaling pathways, subsequently exhibiting antileukemia activity (22). Among the signaling pathways regulated by AMPK, mTOR is known to perform a central role in antitumor activity (23). Recently, several compounds, including metformin and myrtucommulone A, have been reported to exert antileukemia activity via activation of the AMPK/mTOR signaling pathway (24,25). Similarly, the present results revealed that adenine suppressed the viability of K562 cells through activation of the AMPK/mTOR signaling pathway, resulting in  $G_2$ /M phase arrest and autophagic cell death.

Autophagy serves an important role in regulating cellular physiology, removing senescent organelles and degrading long-lived proteins (26,27). In starved cells, the fatty acids and amino acids produced by hydrolysis of lipids, and proteins in autophagolysosomes provide energy to maintain cell survival (28). However, elongated autophagy is proposed to trigger autophagic caspase-independent type II programmed cell death (29). Thus, the roles of autophagy in sustaining or killing cancer cells are complicated. Although cytotoxicity of antitumor drugs is diminished by autophagy to a certain extent (30), reduced expression of autophagic genes, including Atg4, Atg5 and Atg7, has been reported to promote tumor formation in genetically-engineered mice (31,32). Collectively, the findings suggest that autophagy possesses tumor suppressive activity. In the present study, it was demonstrated that adenine can trigger autophagic cell death, but not apoptosis, consequently suppressing the viability of K562 cells. Thus the results of the current study indicate that adenine may serve as a potential anti-leukemia agent.

In conclusion, the present study demonstrated that adenine treatment significantly suppresses the viability of CML K562 cells through activation of the AMPK/mTOR pathway and synergistic induction of  $G_2/M$  phase arrest, and autophagy signaling. Thus, adenine represents a promising effective antiproliferative agent against human leukemia cells.

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