The short-term effect of histone deacetylase inhibitors, chidamide and valproic acid, on the NF-κB pathway in multiple myeloma cells

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Abstract. Research regarding histone deacetylase (HDAC) inhibitors (HDACis) has garnered interest for the treatment of multiple myeloma (MM). In addition, the high expression of nuclear factor (NF)-κB in MM cells is considered an important factor in the occurrence and development of MM. The present study aimed to determine the short-term effects of HDACis, chidamide and valproic acid (VPA), on MM cells, their effects on NF-κB and the underlying mechanisms. The present study measured HDAC activity, and the proliferation and apoptosis of U266 and RPMI8226 MM cells following treatment with various concentrations of chidamide and VPA for 6 and 48 h. Western blotting was used to detect the expression levels of phosphorylated (p)-IkB kinase (IKK)α/β, NF-κB p65 and inhibitor of NF-κB (IkBo) in U266 and RPMI8226 cells at various time points following treatment with chidamide and VPA (0, 2, 4 and 6 h). The results revealed that chidamide and VPA had no significant effect on the HDAC activity, proliferation and apoptosis of cells at 6 h; however, cell HDAC activity and proliferation were inhibited, and apoptosis was induced at 48 h. Furthermore, the expression levels of IkBo were gradually increased over time, whereas the expression levels of NF-κB p65 gradually decreased. These findings indicated that long-term (48 h) treatment with the HDACis chidamide and VPA inhibited the proliferation and promoted the apoptosis of MM cells; however, these HDACis had little effect on cell proliferation and apoptosis in the short term (6 h). Notably, in the short term (2-6 h), hyperactivation of NF-κB was inhibited via the IκBα-NF-κB p65 pathway. These findings indicated that cell growth may be inhibited and drug susceptibility may be promoted by blocking the NF-κB pathway at an early stage, when HDACis are combined with other drugs in the treatment of MM.

Introduction

Multiple myeloma (MM) is a monoclonal, malignant, proliferative tumor derived from plasma cells, which is characterized by monoclonal proteins in the serum or urine, and clonal plasma cells in the bone marrow (1). MM accounts for ~13% of hematological malignancies worldwide (2). With the introduction of novel therapeutic drugs and methods, the 5-year survival rate of patients with MM has been increased to 45% (3); however, the majority of these patients experience relapse and refractory disease. The treatment of patients with relapsed/refractory MM remains a complex issue (4).

Due to the abnormal expression pattern of epigenetic genes in MM, including abnormal methylation, the overproduction of misfolded proteins and the successful application of panobinostat in the clinical treatment of relapsed and refractory MM, histone deacetylase (HDAC) inhibitors (HDACis) have become a focus of research in the treatment of MM (5,6). HDACis can be classified as members of five classes of compounds: i) Hydroxamic acids, ii) short chain fatty acids, iii) benzamides, iv) cyclic tetrapeptides and v) sirtuin inhibitors (7). HDACis bind to the catalytic region of HDACs, reduce their activity, and exert their antitumor effects by altering epigenetic modifications (8).

Chidamide is a novel benzamide class of HDACi, which was developed in China. It exerts its cell death-inducing effects on several types of tumor through its high selective inhibition of HDAC1, 2, 3 and 10 (9), and has been approved by the China Food and Drug Administration for its clinical use in the treatment of peripheral T-cell lymphoma. In vitro, negative effects on MM cells can be exerted by blocking cell cycle progression, and inducing caspase-3-mediated apoptosis and DNA damage (9). Furthermore, a previous study reported the successful treatment of a patient with relapsed/refractory MM with chidamide, bortezomib and dexamethasone in China (10).

Valproic acid (VPA) is type of clinical antiepileptic drug, which belongs to the short-chain fatty acids class of...
HDACis, which can exert antitumor effects by acting on class I and IIa HDACs (11). In vitro, it has been reported that VPA inhibits the development of MM by blocking cell cycle progression (12), inducing apoptosis (13), suppressing angiogenesis (14,15), inhibiting the Notch pathway (16) and regulating autophagy (17,18). In addition, VPA has been revealed to inhibit the nuclear factor (NF)-κB pathway in malignant myeloblasts in the short term (2-4 h), and this effect remains following denucleation of cells. It has also been reported that the inhibitory effects on the NF-κB pathway are not regulated by histone acetylation, but are a result of direct effects on NF-κB pathway-associated proteins (19).

NF-κB is an important nuclear transcription factor, which comprises RelA (p65), RelB, c-Rel, NF-κB1 (p50/p105) and NF-κB2 (p52/p100) subunits (20). In addition, NF-κB is involved in classical and non-classical pathways. The NF-κB classical pathway predominantly refers to the p50/p65 heterodimer, which is ubiquitous in mammalian cells. The p50/p65 heterodimer coexists in the cytoplasm with inhibitor of NF-κB (IκB) in an inactive state (21). IκB kinase (IKK) is phosphorylated by the action of inflammatory factors or apoptotic genes, which leads to the degradation of IκB, after which, NF-κB is released from IκB, translocates to the nucleus, induces target gene production and participates in various cellular processes (22). The high activation state of NF-κB in MM is an important cause of MM development and drug resistance (23,24). Downregulating the activity of NF-κB in MM cells can inhibit cell proliferation and drug resistance (25), thus providing novel therapeutic targets.

There are numerous proteins involved in the NF-κB pathway, and inhibition of NF-κB is predominantly an additional inhibitory effect of other targeted drugs, such as arsenic trioxide (26) and VPA (19); the latter acts rapidly on P39 MDS/AML cells, and its effects may not be regulated by histone acetylation. Therefore, the present study aimed to determine whether short-term treatment with chidamide and VPA, both of which act via epigenetic modification, can induce inhibitory effects on the NF-κB pathway in MM cells. In addition, the study aimed to determine whether these treatments could affect the proliferation and apoptosis of MM cells in the short term.

Materials and methods

Cell lines and culture. Human MM cell lines U266 and RPMI8226 were purchased from the Tumor Cell Bank of Chinese Academy of Medical Sciences (Beijing, China). U266 cells were cultured in RPMI-1640 (Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) supplemented with 10% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc.) and 1% penicillin/streptomycin, and RPMI-8226 cells were cultured in RPMI-1640 supplemented with 15% FBS and 1% penicillin/streptomycin, at 37°C in a humidified incubator containing 5% CO₂.

Cell Counting kit (CCK)-8 cell proliferation assay. Cells (5x10⁵-1x10⁶) were seeded in 96-well plates with culture medium. U266 cells were cultured with chidamide (0.5, 1, 2, 4 and 8 µM) or VPA (0.5, 1, 2 and 4 mM) for 6 and 48 h, whereas RPMI-8226 cells were cultured with chidamide (0.5, 1, 2, 4 and 8 µM) or VPA (0.25, 0.5, 1, 2 and 4 mM) for 6 and 48 h. Simultaneously, the chidamide control group was treated with an equal volume of dimethyl sulfoxide (DMSO), and the VPA control group was treated with an equal volume of culture medium. In addition, the blank control well contained no cell culture solution. Each group was treated in triplicate at the same time. Following treatment, a 10-µl aliquot of CCK-8 (Dojindo Molecular Technologies, Inc., Kumamoto, Japan) was added to each well, and the cells were incubated for a further 4 h. The plate was read at a wavelength of 450 nm using a microplate reader and cell proliferation was calculated using the following equation: Cell proliferation inhibition rate (%)=[1-(OD450 of the experimental group-OD450 of the blank control group)/OD450 of the control group-OD450 of the blank control group)] x100%. OD refers to optical density. Each experiment was independently repeated three times.

Flow cytometry and apoptosis. Cells (1x10⁵) were seeded in 24-well plates with culture medium. U266 cells were incubated with chidamide (1, 2, 4 and 8 µM) or VPA (0.5, 1, 2 and 4 mM) for 6 and 48 h, whereas RPMI8226 cells were incubated with chidamide (1, 2, 4 and 8 µM) or VPA (0.25, 0.5, 1 and 2 mM) for 6 and 48 h. The chidamide control group was treated with an equal volume of DMSO, and the VPA control group was treated with an equal volume of culture medium. Annexin V-fluorescein isothiocyanate/propidium iodide (BD Pharmingen; BD Biosciences, Franklin Lakes, NJ, USA) double staining was used to quantify apoptosis, according to the manufacturer's protocol, and cell apoptosis was analyzed by flow cytometry using BD FACSDiva software 8.0 (BD Biosciences).

Western blot analysis. U266 cells were treated with chidamide (2 µM) or VPA (1 mM), and RPMI8226 cells were treated with chidamide (2 µM) or VPA (0.5 mM) at different time points (0, 2, 4 and 6 h). For HDAC activity analysis. U266 and RPMI8266 cells were treated with chidamide (1 µM) and VPA (1 mM) for 72 h at 37°C. Subsequently, cells were resuspended in lysis buffer containing radioimmunoprecipitation assay buffer and 10 µg/ml phenylmethylsulfonyl fluoride. Protein concentrations were determined using the bicinchoninic acid protein assay kit (Pierce; Thermo Fisher Scientific, Inc. Proteins
(20 µg) were separated by 8% SDS-PAGE and were transferred onto nitrocellulose membranes, which were blocked with 5% non-fat milk in Tris-buffered saline containing 0.1% Tween-20. The membranes were then incubated with the following primary antibodies at 4˚C overnight: Anti-IκBα (1:1,000; cat. no ab32518; Abcam, Cambridge, MA, USA), anti-NF-κB p65 (1:50,000; cat. no. ab32536; Abcam), anti-IKKα/β (1:1,000; cat. no. ab178870; Abcam), anti-phosphorylated (p)-IKKα/β (1:1,000; cat. no. 2697; Cell Signaling Technology, Inc., Danvers, MA, USA), anti-acetyl-histone H3 (1:1,000; cat. no. 9649; Cell Signaling Technology, Inc.) and anti-GAPDH (1:1,000; cat. no. ab9485; Abcam) as an internal control. Subsequently, membranes were visualized using an enhanced chemiluminescence (ECL) plus reagent (Western Lightning Plus-ECL; cat. no. NEL104001EA; PerkinElmer, Inc., Waltham, MA, USA), following the addition of a horseradish peroxidase-conjugated secondary antibody at room temperature for 1 h (1:2,500; cat. no. TA140002; OriGene Technologies, Inc., Beijing, China), under a chemiluminescence and fluorescence imaging system (LumiX®/ChemiX®/FluorChemIX®; Federal Bioproducts, Inc., London, United Kingdom).

Statistical analysis. Each experiment was repeated three times. All data are expressed as the means ± standard deviation, and were analyzed using SPSS19.0 (SPSS, Inc., Chicago, IL, USA). Pairwise comparisons were conducted using Student’s t-test. Multiple comparisons were performed by one-way analysis of variance followed by Dunnett’s post hoc test. P<0.05 was considered to indicate a statistically significant difference.

Results

Chidamide and VPA enhance histone H3 acetylation and inhibit HDAC activity. To explore whether chidamide and VPA might influence histone H3 acetylation in MM cells, the present study detected histone H3 acetylation levels in cells following exposure to chidamide or VPA. Following treatment with chidamide or VPA for 48 h, the acetylation levels of histone H3 were increased compared with in cells treated for 0 or 6 h. There were no marked alterations in acetylation levels between the 0 and 6 h chidamide or VPA groups (Fig. 1A). The results of a HDAC activity analysis indicated that treatment of cells with chidamide or VPA for 48 h resulted in a greater HDAC inhibition compared with in the 0 and 6 h culture groups (Fig. 1B). These results indicated that, in the 48 h chidamide and VPA culture groups, levels of histone H3 acetylation were increased and HDAC activity was inhibited compared with in the 0 and 6 h culture groups. There was no significant difference between the 0 and 6 h culture groups when MM cells were exposed to chidamide or VPA.

Chidamide and VPA inhibit MM cell proliferation after 48 h. In order to study the effects of chidamide and VPA on the activity of U266 and RPMI8226 cells during a short time period, cell viability was measured after 6 h of drug action. In addition, cell viability was measured after 48 h as a comparison. The experimental results revealed that there was no obvious effect on cell proliferation after 6 h (P>0.05); however, chidamide and VPA inhibited cell proliferation after 48 h compared with in the 6 h group (P<0.05; Fig. 2). In addition, the inhibition of cell proliferation was increased alongside drug concentration. These results indicated that chidamide and VPA may affect the proliferation of U266 and RPMI8226 cells in the long term.

Chidamide and VPA induce MM cell apoptosis after 48 h. Cell apoptosis was measured following treatment with chidamide and VPA for 6 and 48 h using flow cytometry. The results revealed that chidamide and VPA exerted no significant effect on cell apoptosis after 6 h (P>0.05; Fig. 3). Conversely, after 48 h, chidamide (2, 4 and 8 µM) and VPA (1, 2 and 4 mM) induced apoptosis of U266 cells, and chidamide (1, 2, 4 and 8 µM) and VPA (0.25, 0.5, 1 and 2 mM) induced apoptosis of RPMI8226 cells compared with in the control group (P<0.05; Fig. 4). The apoptotic effect was increased with increasing
drug concentrations. These findings suggested that chidamide and VPA may affect the apoptosis of U266 and RPMI8226 cells in the long term.

**Chidamide and VPA inhibit the p-IKKα/β-IxBα-NF-κB p65 pathway in the short term in MM cells.** NF-κB exhibits high activity in MM cells, and promotes its occurrence and development (27). To explore the effects of chidamide and VPA on the NF-κB signaling pathway in U266 and RPMI8226 cells, western blotting was conducted to detect the expression levels of NF-κB pathway-related proteins at 0, 2, 4 and 6 h following treatment with chidamide and VPA. The expression levels of IxBα were gradually increased in response to increasing drug concentrations. Conversely, the expression levels of p-IKKα/β and NF-κB p65 were gradually decreased; however, there was some effect on total IKKα/β protein expression levels following treatment with chidamide or VPA (Fig. 5). These findings indicated that chidamide and VPA may exert their anti-MM effects through inhibiting the NF-κB pathway, and alterations in the expression levels of pathway-associated protein were observed within the 2-6 h time period.

**Discussion**

As an important nuclear transcription factor, NF-κB is involved in various biological cellular processes (28). Gene expression analysis and immunostaining revealed that the majority of MM cells undergo continuous activation of NF-κB, which can promote the development of MM by affecting cell survival, apoptosis, angiogenesis, cell migration and drug resistance (23,29).

HDACis are considered an important research focus as a potential method to treat MM. Among HDACis, VPA has been reported to negatively effect MM via various mechanisms *in vitro*. In addition, NF-κB activity of malignant myeloblasts can also be inhibited during the early stage, and this inhibition exists following denucleation (19). As the first HDACi developed in China, chidamide has successfully been used in the clinical treatment of peripheral T-cell lymphoma (30), and it has also been confirmed to kill and inhibit various tumor cells *in vitro* (31-33); however, research regarding HDACis in MM is rare. Due to the importance of NF-κB in the development of MM, and the ability of VPA to rapidly inhibit the activity of NF-κB in malignant myeloblasts, the present study hypothesized that chidamide and VPA may inhibit NF-κB, and affect MM cell proliferation and apoptosis in the short term.

The present study selected two MM cell lines, U266 and RPMI8226, to ensure the reliability of the experimental results. Exploration and selection of drug concentrations used for the cell proliferation assay in this study were derived from previous studies on chidamide and VPA in MM (9,15) or other tumors (34,35). Chidamide doses (1, 2, 4 and 8 µM) that exhibited 20-90% cytotoxicity in the cell proliferation study were selected for the treatment of U266 and RPMI8226 cells prior to apoptosis assessment. In the VPA groups, VPA (0.5, 1, 2 and 4 mM) was used to treat U266 cells, whereas VPA (0.25, 0.5, 1 and 2 mM) was used to treat RPMI8226 cells. In addition, doses close to the half maximal inhibitory
concentrations of chidamide (2.395 µM) and VPA (0.554 mM) were used to treat cells prior to western blotting. Proliferation and apoptosis of cells was detected following treatment with chidamide or VPA for 6 and 48 h. The results revealed that these treatments could inhibit cell proliferation and promote apoptosis of U266 and RPMI8226 cells after 48 h, which was consistent with the findings from previous studies in adenoid cystic carcinoma and myelodysplastic syndromes (34,36); however, no significant effects were detected after 6 h. In addition, the present results revealed that treatment with chidamide and VPA for 48 h led to higher levels of histone H3 acetylation and inhibition of HDAC activity compared with in the 0 and 6 h groups; there was no significant difference between the 0 and 6 h culture groups. Western blotting was used to detect the expression levels of NF-κB pathway-associated proteins, p-IKKα/β, IκBα and NF-κB p65, at different time

Figure 3. Effects of chidamide and VPA on the apoptosis of multiple myeloma cells after 6 h, as determined by flow cytometry using Annexin V-FITC/PI staining. Effects of various concentrations of chidamide on the apoptosis of (A) U266 and (B) RPMI8226 cells. Effects of various concentrations of VPA on the apoptosis of (C) U266 and (D) RPMI8226 cells. (E) Chidamide and VPA had no effect on the apoptosis of U266 and RPMI8226 cells after 6 h. FITC, fluorescein isothiocyanate; PI, propidium iodide; VPA, valproic acid.
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Figure 4. Effects of chidamide and VPA on the apoptosis of multiple myeloma cells after 48 h, as determined by flow cytometry using Annexin V-FITC/PI staining. Effects of various concentrations of chidamide on the apoptosis of (A) U266 and (B) RPMI8226 cells. Effects of various concentrations of VPA on the apoptosis of (C) U266 and (D) RPMI8226 cells. (E) Chidamide and VPA could promote apoptosis of U266 and RPMI8226 cells after 48 h; the apoptotic rate was increased with increasing drug concentrations. *P<0.05, **P<0.01 vs. the untreated control group. FITC, fluorescein isothiocyanate; PI, propidium iodide; VPA, valproic acid.

points (0, 2, 4 and 6 h) following treatment. The expression levels of IκBα were increased after 2 h, whereas p-IκKα/β and NF-κB p65 expression was decreased; the extent of expression alterations was increased with the prolongation of treatment time. However, there was some effect on the total expression of IκKα/β following treatment with chidamide and VPA; therefore, it is unclear as to whether the effects of chidamide and VPA on p-IκKα/β follow alterations in the total expression of IκKα/β. Further studies are required to determine the effects and mechanism of chidamide and VPA on p-IκKα/β and total IκKα/β expression. The present findings indicated that IκBα and NF-κB expression was altered following treatment with chidamide and VPA. Therefore, chidamide and VPA may inhibit the activity of NF-κB in MM cells via the NF-κB p65 pathway during the early stage, which is similar to the effects of VPA and trichostatin on malignant myeloblasts (19). Unlike the effects of VPA on Notch signaling in MM cells at 48 h (16), autophagy of lesioned cortices 24 h after traumatic brain injury (TBI) and HDAC3 expression 72 h after TBI (37), and the effects of chidamide on Mcl-1 expression in pancreatic
cancer cells at 24 h (38), the effects of chidamide and VPA on the NF-κB pathway were detected in the short term in the present study.

The present study hypothesized that HDACi-induced inhibition of the NF-κB pathway in tumor cells depended on non-histone acetylation in the cytoplasm, rather than the epigenetic reprogramming caused by alterations in the nuclear transcriptome. However, the specific targets of HDACis in the cytoplasm require further analysis.

Previous studies have revealed that chidamide and DNA damaging agents can synergistically promote apoptosis in relapsed and refractory acute myeloid leukemia cells and reduce tumor burden in patients (39), and that VPA and bortezomib cooperate to induce pancreatic cancer cell death (40). Although chidamide and VPA did not inhibit MM cell proliferation and promote apoptosis, as expected, in the short term (6 h), which may be associated with the complexity of cell proliferation and development mechanisms, the rapid inhibition of NF-κB may inhibit the continued growth of MM cells in the early stages. It remains to be determined as to whether chidamide or VPA, combined with other drugs, may increase the cell death-inducing effects of these other drugs and increase drug sensitivity. In addition, whether the findings of the present study are associated with the successful treatment of patients with relapsed and refractory MM with chidamide combined with bortezomib and dexamethasone is unclear. Further studies are required regarding whether the combination of HDACis and other drugs can increase anti-MM effects by blocking NF-κB in the early stages.

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Availability of data and materials
All data generated or analyzed during this study are included in this published article.

Authors' contributions
ZL and RF designed the research. ZL and QJ contributed to cell culture and western blotting. YL, FM and CX contributed to RT-qPCR and flow cytometry. All authors read and approved the final manuscript.

Ethics approval and consent to participate
Not applicable.

Patient consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

References
1. Kohgo Y: Recent advances in multiple myeloma. Int J Clin Oncol 20: 411-412, 2015.
2. Palumbo A, Anderson K. Multiple myeloma. Current Opinion in Oncology 24(Suppl): S1, 2011.
3. Siegel R, Ma J, Zou Z and Jemal A: Cancer statistics, 2014. CA Cancer J Clin 64: 9-29, 2014.
4. Moreau P, Attal M and Fagon T: Frontline therapy of multiple myeloma. Blood 125: 3076-3084, 2015.
5. Laubach JP, San-Miguel JF, Hungria V, Hou J, Moreau P, Sezer O and Heider U: The effects of the histone deacetylase inhibitor valproic acid on cell cycle, growth suppression and apoptosis in multiple myeloma. Haematologica 91: 248-251, 2006.
6. Greig SL: Panobinostat: A review in relapsed or refractory cancer. Haematologica 91: 248-251, 2006.
7. Ceccacci E and Minucci S: Inhibition of histone deacetylases 6. PLoS One 7: 229-237, 2012.
8. Eckschlager T, Plch J, Stiborova M and Hrabeta J: Histone deacetylase inhibitors: An advance in myeloma therapy? Expert Rev Hematol 10: 229-237, 2017.
9. Gu et al: Chidamide inhibits the NF-κB pathway in multiple myeloma. Mol Med Rep 10: 832-838, 2014.
10. Alkalay I, Yaron A, Hatzubai A, Ammon A, Chechonov A and Ben-Neria Y: Stimulation-dependent IκBα phosphorylation marks the NF-κB/xB family for degradation via the ubiquitin-proteasome pathway. Proc Natl Acad Sci USA 92: 10599-10603, 1995.
11. Bhatti AC, Shishodia S, Reuben JM, Weber D, Alexanian R, Raj-Vadhan S, Estrov Z, Talpaz M and Aggarwal BB: Nuclear factor-xB inhibitors: An advance in myeloma therapy? Expert Rev Hematol 10: 229-237, 2017.
12. Xu L, Tang HL, Gong X, Xing XL, Dong Y, Gao GX, Gu HT, Shu MM and Chen XQ: Valproic acid represses autophagy and enhances the anti-myeloma activity of DNA-damaging drugs. Zhongguo Shi Yan Xue Ye Xuezazhi 23: 718-721, 2015 (In Chinese).
13. Wang YY, Zhang ZH, Zhao RJ, Li H, Wang TR, Yan LN, Gu CH, Zhao L and Hao CL: Valproic acid activates autophagy in multiple myeloma cell lines RPMI2652 and U266. Zhonghuaxue Ye Xuezazhi 37: 478-483, 2016 (In Chinese).
14. Dong XF, Song Q, Li LZ, Zhao CL and Wang LQ: Histone deacetylase inhibitor valproic acid inhibits proliferation and induces apoptosis in MM3 cells via downregulating VEGF receptor. Neuro Endocrinol Lett 28: 775-780, 2007.
15. Kitazoe KI, Abe M, Hisa M, Oda A, Amou H, Harada T, Nakano H, Takeuchi K, Hamashimo T, Ozaki S and Matsumoto T: Valproic acid exerts anti-tumor as well as anti-angiogenic effects on myeloma. Int J Hematol 89: 45-57, 2009.
16. Qiao L, Yang Y, Zhao RJ, Wang LH, Zhao L, Yan LN, Zhang ZH and Hao CL: Effects of VPA on the expression of notch signaling pathway in multiple myeloma. R PBM 8226 cell line. Zhongguo Shi Yan Xue Ye Xuezazhi 24: 1449-1453, 2016 (In Chinese).
17. Jin YL, Dong BX, Xu L, Tang HL, Gao GX, Xiong Y, Gao GX, Shu MM and Chen XQ: Valproic acid represses autophagy and enhances the anti-myeloma activity of DNA-damaging drugs. Zhongguo Shi Yan Xue Ye Xuezazhi 23: 718-721, 2015 (In Chinese).
18. Wang JN, Zhang YY, Zhang ZH, Zhao RJ, Li H, Wang TR, Yan LN, Gu CH, Zhao L and Hao CL: Valproic acid activates autophagy in multiple myeloma cell lines RPMI2652 and U266. Zhonghuaxue Ye Xuezazhi 37: 478-483, 2016 (In Chinese).
19. Fang C, Grosjean J, Taillier M, Bocher S, Adès L, Perfettini JL, de Botton S, Fenaux P and Kroemer G: A novel effect of DNA methyltransferase and histone deacetylase inhibitors: NFκB activation in malignant myeloblasts. Cell Cycle 7: 2193-2198, 2008.
20. Dutta J, Fan Y, Gupta N, Fan G and Gélinas C: Recent insights into the regulation of programmed cell death by NF-xB. Oncogene 25: 6800-6816, 2006.
21. Oeckinghaus A and Ghosh S: The NF-xB family of transcription factors and its regulation. Cold Spring Harb Perspect Biol. 4(4)00034, 2009.
22. Chen ZJ: Ubiquitination in signaling to and activation of IKK. Immunol Rev 246: 95-106, 2012.
23. Annunziata CM, Davis RE, Demchenko Y, Bellamy W, Gabrea A, Zhan F, Lenz G, Hamamura I, Wright G, Xiao W, et al: Frequent engagement of the classical and alternative NF-kappaB pathways by diverse genetic abnormalities in multiple myeloma. Cancer Cell 12: 115-130, 2007.
24. Hayden MS and Ghosh S: NF-xB, the first quarter-century: Remarkable progress and outstanding questions. Genes Dev 26: 203-234, 2012.
25. Kannayari R, Hay HS, Rajendra P, Li F, Shanmugam MK, Vali S, Abbasi T, Kapoor S, Sharma A, Kumar AP, et al: Celestrol inhibits proliferation and induces chemosensitization through down-regulation of NF-xB and STAT3 regulated gene products in multiple myeloma cells. Br J Pharmacol 164: 1506-1511, 2011.
26. Qu X, Du J, Zhang C, Fu W, Xi H, Zou J and Hou J: Arsenic trioxide exerts antimyeloma effects by inhibiting activity in the cytoplasmic substrates of histone deacetylase 6. PLoS One 7: e32215, 2012.
27. Li Z, Yang Z, Peng X, Li Y, Liu Q and Chen J: Nuclear factor-xB is involved in the protocadherin-10-mediated pro-apoptotic effect in multiple myeloma. Mol Med Rep 10: 832-838, 2014.
28. Akhalay I, Yaron A, Hatzubai A, Orian A, Chechonov A and Ben-Neria Y: Stimulation-dependent IκBα phosphorylation marks the NF-κB-xB family for degradation via the ubiquitin-proteasome pathway. Proc Natl Acad Sci USA 92: 10599-10603, 1995.
29. Bharti AC, Shishodia S, Reuben JM, Weber D, Alexanian R, Raj-Vadhan S, Estrov Z, Talpaz M and Aggarwal BB: Nuclear factor-xB inhibitors: An advance in myeloma therapy? Expert Rev Hematol 10: 229-237, 2017.
30. Shieh Y, Dong M, Hong X, Zhang W, Feng J, Zhu J, Yu L, Ke X, Huang H, Shen Z, et al: Results from a multicenter, open-label, pivotal phase II study of chidamide in relapsed or refractory peripheral T-cell lymphoma. Ann Oncol 26: 1766-1771, 2015.
31. Liu L, Chen B, Qin S, Li S, He X, Qiu S, Zhao W and Zhao H: Histone deacetylase inhibitor Chidamide induces apoptosis of human colon cancer cells. Biochem Biophys Res Commun 392: 190-195, 2010.
32. Ning ZQ, Li ZB, Newman MJ, Shan S, Wang XH, Pan DS, Zhang J, Dong M, Du X and Lu XP: Chidamide (CS055/HBI-8000): A new histone deacetylase inhibitor of the benzamide class with anti-tumor activity and the ability to enhance immune cell-mediated tumor cell cytotoxicity. Cancer Chemother Pharmacol 69: 901-909, 2012.
33. Walkinshaw DR and Yang XJ: Histone deacetylase inhibitors as novel anticancer therapeutics. Curr Oncol 15: 237-243, 2008.
34. Liu Z, Ding K, Li L, Liu H, Wang Y, Liu C and Fu R: A novel histone deacetylase inhibitor Chidamide induces G0/G1 arrest and apoptosis in myelodysplastic syndromes. Biomed Pharmacother 83: 1032-1037, 2016.
35. Saha SK, Yin Y, Kim K, Yang GM, Dayem AA, Choi HY and Cho SG: Valproic acid induces endocytosis-mediated doxorubicin internalization and shows synergistic cytotoxic effects in hepatocellular carcinoma cells. Int J Mol Sci 18: E1048, 2017.
36. Yang S, Nan P, Li C, Lin F, Li H, Wang T, Zhou C, Zhang X, Meng X, Qian H, et al: Inhibitory effect of chidamide on the growth of human adenoid cystic carcinoma cells. Biomed Pharmacother 99: 608-614, 2018.
37. Chen X, Wang H, Zhou M, Li X, Fang Z, Gao H, Li Y and Hu W: Valproic acid attenuates traumatic brain injury-induced inflammation in vivo: Involvement of autophagy and the Nrf2/ARE signaling pathway. Front Mol Neurosci 11: 1-13, 2018.
38. Mu H, Qiao Z, Wang Y, Kuai Q, Li C, Wang Y, Jiang X, Wang X, Li W, He M, et al: Chidamide inhibits aerobic metabolism to induce pancreatic cancer cell growth arrest by promoting Mcl-1 degradation. PLoS One 11: e0166896, 2016.
39. Li Y, Wang Y, Zhou Y, Li J, Chen K, Zhang L, Deng M, Deng S, Li P and Xu B: Cooperative effect of chidamide and chemotherapeutic drugs induce apoptosis by DNA damage accumulation and repair defects in acute myeloid leukemia stem and progenitor cells. Clin Epigenetics 9: 83, 2017.
40. GilardiniMontani MS, Granato M, Santoni C, Del Porto P, Merendino N, D’Orazi G, Faggioni A and Cirone M: Histone deacetylase inhibitors VPA and TSA induce apoptosis and autophagy in pancreatic cancer cells. Cell Oncol (Dordr) 40: 1-14, 2017.