Matrine-induced apoptosis in Hep3B cells via the inhibition of MDM2

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Abstract. Matrine, an alkaloid component derived from the Sophora root, can inhibit cancer cell proliferation and induce autophagy via p53 associated pathways. However, numerous tumor cells lack functional p53 and little is known about the effect of matrine on the p53-deficient/mutant cancer cells. The present study aimed to assess anticancer effects of matrine in p53-deficient human Hep3B hepatoma cells. The present results demonstrated that matrine caused Hep3B cell apoptosis by suppressing gene expression of minute double-mutant (MDM)2. Notably, it was revealed that matrine inhibited MDM2 at the transcriptional level in a time- and dose-dependent manner. This MDM2 inhibition resulted in induction of the p53 family member, p73; however, the functions of p73 were not induced since matrine-induced p73 failed to activate its target genes, p21 and p53 upregulated modulator of apoptosis. The matrine-induced downregulation of MDM2 led to an inhibition of inhibitor of apoptosis protein 3, which might serve a critical role in matrine-induced apoptosis in MDM2-overexpressing Hep3B cells. Finally, combination therapy of matrine with 100 µM epotoside successfully killed more Hep3B cells, suggesting that matrine can sensitize p53-deficient Hep3B cells to epotoside-induced apoptosis.

Introduction

Matrine is a component of the traditional Chinese medical herb Sophora flavescent ait. It performs a variety of medical effects as a κ-opioid receptor and µ-receptor agonist (1). Since Matrine has a wide range of clinical effects, including anti-inflammatory, antiviral, immuno inhibitory, antifibrotic and anti diarrhoea effects (2,3), it has been widely used in treatment of viral hepatitis, hepatic fibrosis, cardiac arrhythmia and skin diseases, including atopic dermatitis and eczema in China (4-7). Therefore, matrine can serve as a potential anticancer drug to treat various types of human cancer. A previous study reported that matrine can suppress the proliferation of hepatoma G2 (HepG2) cells, possibly by inducing apoptosis through the activation of B-cell lymphoma-2-activated X protein (8). However, its antitumor mechanism in p53 deficient hepatoma Hep3B cells remains to be elucidated.

In response to various cellular stresses, the tumor suppressor gene, p53, acts as an important safeguard mechanism by preventing cells from undergoing uncontrolled proliferation in response to DNA damage (9,10). The downstream DNA damage response (DDR) involves a series of events that lead to either cell-cycle arrest induced by p21 or apoptosis induced by p53 upregulated modulator of apoptosis (PUMA) (11). Mouse double-minuted (MDM)2, a target gene of p53 that can form a negative feedback loop with p53, it is also considered as an antiapoptotic factor that can inhibit apoptosis of tumor cells (12). MDM2 has attracted great attention following its identification as a negative modulator of p53. MDM2 can bind to p53 through its N-terminus to repress p53 anticancer functions, while its C-terminus serves as an E3 ubiquitin ligase to mediate p53 degradation, which maintains p53 at a low protein level during normal homeostasis without stress signals (12-14). In addition to interacting with and regulating p53, growing evidence suggests that MDM2 has numerous p53-independent functions. For example, MDM2 was shown to bind to and ubiquitinate Rb, resulting in Rb degradation and release of E2F1, which promotes cell cycle progression (18,19). Since MDM2 is able to bind RNA and shuttle between the nucleus and the cytoplasm, properties of most
internal ribosome entry site (IRES) trans-acting factors (ITAFs)/ribonucleoproteins (RNPs), this suggests that the dephosphorylated cytoplasmic MDM2 may act as an ITAF/RNP to regulate translation through binding of its C-terminus to specific RNAs (20). Additionally, previous studies also demonstrated that MDM2 can physically interact with the IRES of the 5'-untranslated region of inhibitor of apoptosis protein (IAP), and in turn induce translation of the latter, which allows for development of resistance to anticancer treatment (21,22).

As a negative modulator of p53, it may be suggested that MDM2 is an indirect oncogene, when overexpressed would be oncogenic by preventing the release of activated p53. This hypothesis is supported by numerous previous studies (23,24). For example, in MDM2 overexpression mice, the incidence rate of tumor formation is significantly increased. In particular, MDM2 overexpression is even observed in cancer types that lack MDM2 gene amplification, including acute lymphoblast leukemia cells (23,24). Regardless of any molecular mechanism involved, MDM2 overexpression is associated with the development of tumors and poor prognosis.

A previous study reported that matrine at concentrations of 0.25, 0.5, 1.0 and 2.0 mg/ml inhibits the growth of HepG2 cells in a dose- and time-dependent manner (8). The present study used identical concentrations of matrine with minimal modification to investigate the influence of matrine on MDM2 expression and induction of Hep3B cell death. It was revealed that matrine markedly suppressed MDM2 transcription and caused significant apoptosis of MDM2-overexpressing Hep3B cells. In addition, the present study examined the expression of the p53 family member, p73, as well as its downstream effectors, p21 and PUMA. Investigating these effects may assist with elucidating the molecular mechanism by which matrine induces MDM2 downregulation and apoptosis of the p53-deficient Hep3B cells.

Materials and methods

Reagents. Matrine was supplied by Xi’an Tianyuan Biologics Plant (Xi’an, China), with a purity of >99%. Matrine was dissolved in sterile double distilled water at a stock concentration of 40 mg/ml, stored at -20°C in the dark and was subsequently diluted in Dulbecco’s modified Eagle’s medium (DMEM) to obtain the desired concentration. Actinomycin D, benzo(a)pyrene and etoposide were purchased from Sigma-Aldrich (St. Louis, MO, USA). DMEM and fetal bovine serum (FBS) were products from Gibco (Thermo Fisher Scientific, Inc., Waltham, MA, USA). The protein isolation kit was obtained from KeyGen Biotech. Co., Ltd. (Nanjing, China). MDM2 CRISPR activation plasmid, and MDM2 (cat. no. BS1223), PUMA (cat. no. AB10418), p73α (cat. no. 4662), p73α (Try99) (cat. no. 46655) and β-actin (cat. no. 4967) antibodies were all obtained from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA, USA). IAP3 (2F1) antibody (cat. no. ab5746) was purchased from Abcam (Cambridge, MA, USA). An mRNA extraction kit and SYBR green I mix for quantitative polymerase chain reaction (qPCR) were supplied by Qiagen (Hilden, Germany) and Invitrogen (Thermo Fisher Scientific, Inc.), respectively. Activated caspase-9 (cat. no. BS7070), activated caspase-3 (cat. no. BS4301) and activated poly ADP-ribose polymerase (PARP; cat. no. BS7047) antibodies, as well as IRDye-conjugated secondary antibodies (cat. no. BA29880) were all purchased from Bioworld Technology, Inc. (St. Louis Park, MN, USA). The dilution ratio of all primary and secondary antibodies were 1:500 and 1:5,000, respectively. The annexin V-fluorescein isothiocyanate (FITC)/propidium iodide (PI) apoptosis detection kit was purchased from Multi-Sciences Biotechnology (Hangzhou, China).

Cell culture and treatment. Both Hep3B and L0-2 cells were obtained from the Cell Bank of Type Culture Collection of the Chinese Academy of Science (Shanghai, China). The two cell lines were grown in DMEM containing 10% (v/v) FBS and were cultured at 37°C in a humidified 5% CO₂ atmosphere.

Western blotting. Hep3B and L0-2 cells were collected and lysed in a traditional radioimmunoprecipitation buffer (1 M Tris-HCl, 5 M NaCl, 1% Nonidet P-40, 1% sodium deoxycholate, 0.05% SDS and 1 mM phenylmethylsulfonyl fluoride) for 15 min. The proteins were denatured at 96°C for 5 min following mixing with 5 µl SDS-loading buffer. The proteins were subsequently separated on 12% SDS-PAGE gels and were transferred onto polyvinylidene difluoride membranes (EMD Millipore, Billerica, MA, USA). The immunoblotting assay was performed as described previously (25). The protein band densities were measured using Quantity One software (Bio-Rad, Hercules, CA, USA). The data was expressed as the density normalized against that of β-actin.

Reverse transcription (RT)-qPCR. The total RNA was extracted from cells using the RNeasy Mini kit (cat. no. 75144; Qiagen), according to the manufacturer’s protocol. cDNA synthesis was performed using 1 µg total RNA sample mixed with dNTP, reverse transcriptase, random monomers, oligo-dT as primers, 10X reaction buffer and DNAse I. The samples were run in a PCR system (T100; Bio-Rad) at the following temperatures: 42°C for 55 min, 70°C for 10 min and 4°C until collected. The samples were stored at 4°C. The amplification was performed using a 7900 Real-Time PCR system (Applied Biosystems; Thermo Fisher Scientific, Inc.), according to the manufacturer’s protocol for the Quanti-Fast SYBR Green RT-PCR kit (Qiagen). All specific primers for amplification of specific genes, as well as the housekeeper gene β-actin, were designed and synthesized by Qiagen. All samples were run in triplicate.

Pulse-chase and cycloheximide (CHX) assays. The degradation rate of mRNA was assessed using a standard actinomycin D analysis. Following the addition of 5 µg/ml actinomycin D, Hep3B cells were treated with or without matrine. Subsequently, at different time points, the cells were harvested for isolation of the total RNA. The mRNA expression of MDM2 was detected by RT-qPCR.

Protein translation level was evaluated using a standard protein-synthesis inhibitor CHX assay. Briefly, the cells were pretreated with 100 µg/ml CHX for 15 min at 4°C to arrest polysome migration. The cells were subsequently treated with or without matrine, and the cells were harvested and lysed.
in a buffer containing 20 mM Tris-HCl (pH 8.0), 100 mM NaCl, 5 mM MgCl₂, 0.5% Triton X-100 and 500 U/ml RNAsin, as well as a cocktail of protease inhibitors. Fractionation was performed on a 15-45% (w/v) sucrose gradient, centrifuged at 24,000 x g for 1 h at 4˚C. The fractions were collected from each gradient tube by upward replacement and absorption monitored at an optical density of 254 nm, using a fractionator (Brandel, Gaithersburg, MD, USA). Lastly, immunoblotting was performed to observe concurrent turnover of MDM2.

MDM2 plasmid transfection. For transfection, Hep3B cells were seeded into 6-well plates at density of 2x10⁵ cells/well. Following culturing to 70-80% confluence, the cells were harvested to make a cell suspension using cell medium. Both Lipofectamine 2000 transfection reagent (Invitrogen; Thermo Fisher Scientific, Inc.) and MDM2 overexpression plasmid were diluted with the Opti-MEM I medium (Thermo Fisher Scientific, Inc.) at a final concentration of 5% and 50 nM, respectively. The two mixtures were mixed and incubated at room temperature for an additional 20 min. Following incubation, the above transfection mixture was added to new 6-well plates. The Hep3B cell suspensions (final concentration for MDM2 plasmid, 5 nM) were overlaid onto the transfection mixture. Following incubation at 37˚C for 4 h, the media was removed and the cells were cultured with fresh cell medium.

Flow cytometry. The matrine-induced cell cycle arrest and cell death were assessed by flow cytometry. For the cell-cycle sample preparation, the harvested cells were washed twice with phosphate-buffered saline (PBS) and fixed overnight with 70% ethanol at 4˚C. After washing three times with PBS, the cells were resuspended with 0.5 ml PBS containing 20 µg/ml PI and 1 µg/ml RNase A. Following incubation at 4˚C for an additional 30 min, the cells were analyzed using a FACScan (Becton-Dickinson, San Jose, CA, USA) with WinList software (Verity Software House, Topsham, ME, USA). For analysis of cell death type, the matrine-treated cells were harvested and washed twice with PBS. The cells were subsequently suspended in 400 µl binding buffer containing 10 µl PI and 5 µl Annexin V-FITC, and were incubated for 20 min at room temperature in the dark, followed by flow cytometric analysis.

Statistic analysis. The differences between the groups were examined for statistical significance using the one-way analysis of variance followed by Dunnett's post-hoc test using SPSS.
MDM2 inhibition occurred at ~4 h post-treatment and was followed by a steady-state downregulation (Fig. 1C), with a style distinct from irradiation-treatment that instead induced a rapid and transient (1-2 h) decrease of MDM2 expression, followed by a clear upregulation (Fig. 1D).

Matrine-induced decrease of the mRNA synthesis of MDM2. The present study next investigated the mechanism by which matrine suppress the expression of MDM2. As shown in Fig. 2A, the mRNA expression of MDM2 was clearly inhibited by matrine. To investigate whether the matrine-induced MDM2 mRNA decrease is associated with mRNA stability, pulse-chase and RT-qPCR reactions were performed. As shown in Fig. 2B, matrine caused no affect on the mRNA stability of MDM2. MDM2 protein stabilization was further assessed using a standard CHX pulse-chase assay. As shown in Fig. 2C, compared with the control groups, the half-life of MDM2 was not affected by matrine, suggesting that matrine decreases MDM2 via the inhibition of mRNA synthesis. Enhanced expression of p73 by matrine is a result of matrine-mediated inhibition of MDM2, since MDM2 can compete with p73 for binding to its upstream effectors (17).

The present study investigated whether the matrine-induced MDM2 decrease can affect the expression of p73α. Firstly, a possible direct influence of matrine on p73 mRNA level was investigated by comparing it with the known effect of benzo(a)pyrene on the mRNA expression of p73 (26). As shown in Fig. 3A, benzo(a)pyrene increased p73 transcription in the cells. By contrast, matrine failed to either increase or decrease the mRNA expression of p73. Whether matrine can affect p73 translation was next determined. As shown in Fig. 3B, no difference in the expression of p73 was detected in the presence or absence of the protein synthesis, as assessed by CHX treatment. Lastly, the turnover of p73 protein was measured in matrine-treated cells, as shown in Fig. 3C. Compared with the control groups, matrine notably increased the half-life of p73. These results suggested that the observed matrine-upregulated expression of p73 in Hep3B cells occurs only at the post-translational level. Additionally, the expression of p73 after matrine treatment in Hep3B cells was assessed, as shown in Fig. 3D. Matrine increased the expression of p73 following the downregulation of MDM2.

p73 cell-cycle arrest function is not activated in matrine-treated Hep3B cells. p21 and PUMA are two down-stream targets of p73. Therefore, the present study examined the expression levels of p21 and PUMA in matrine-treated Hep3B cells. As shown in Fig. 4A, benzo(a)pyrene clearly increased the expression levels of p21 and PUMA, as well as the mRNA expression of MDM2. By contrast, matrine failed to increase or decrease the expression levels of the above genes. Additionally, as shown in Fig. 4B, the matrine-induced p73 protein increase was not to the same level as that of epotoside-induced. The data in Fig. 4B also demonstrated that epotoside increased the protein expression levels of p21 and PUMA, whereas matrine decreased the expression levels. In addition, matrine failed to induce accumulation of the p73 activated form (p73α-Try99), compared with epotoside, which increased p73α-Try99 levels. The cell-cycle analysis also confirmed that p21 was not functional in the

Results

MDM2 expression in matrine-treated Hep3B and L0-2 cells. To assess whether MDM2 expression levels were influenced by matrine, western blotting analysis was performed. It was revealed that matrine notably downregulated MDM2 in the two studied cell lines (Fig. 1A). Matrine suppressed the protein expression of MDM2 in a dose-dependent manner, even at low concentrations (Fig. 1B). The matrine-induced
matrine-treated cells. As shown in Fig. 4C, matrine failed to induce G1 arrest. By contrast, epotoside clearly induced cell accumulation in G1 phase, whereas matrine decreased cell numbers in G1, S and G2/M.

Matrine inhibited the expression of IAP3 in MDM2-overexpressing Hep3B cells. IAP3 is a translational target of MDM2, which exhibits reduced expression level following MDM2 silencing (21). The present study assessed the expression levels of IAP3. As shown in Fig. 5A, matrine markedly decreased the expression levels of MDM2 and IAP3, whereas both genes were increased in the epotoside-treated cells. To investigate whether MDM2 inhibition affected the expression of IAP3 in matrine-treated Hep3B cells, an MDM2-overexpressing Hep3B cells cell line was treated with matrine. As shown in Fig. 5B, a more marked decrease of IAP3 was induced in the matrine-treated MDM2-overexpressing Hep3B cells compared with the matrine-treated negative plasmid-transfected Hep3B cells. Additionally, the IAP3 protein stability was also evaluated in the matrine-treated Hep3B cells. As shown in Fig. 5C, compared with the control group, matrine caused no effect on IAP3 protein stability. These results suggested that matrine can strongly suppress the expression of IAP3 in MDM2-overexpressing Hep3B cells at both the mRNA and protein expression levels.

Matrine sensitized Hep3B cells to epotoside-induced apoptosis. The present study assessed the effect of matrine on Hep3B and MDM2-overexpressing Hep3B cell viabilities using an MTT assay. As shown in Fig. 6A, matrine exhibited cytotoxic activity in each of the cell lines. Matrine exhibited a more marked cytotoxic effect on MDM2-overexpressing Hep3B cells that express very high level of MDM2. To investigate the mechanism by which matrine induces MDM2-overexpressing Hep3B cell death, the activation of several apoptotic effectors was assessed. As shown in Fig. 6B, cleavage of caspases-3 and -9, and PARP in MDM2-overexpressing Hep3B cells was observed 4 h post-treatment. Whether matrine has synergistic effects on epotoside-induced apoptosis was determined by treating the Hep3B cells with both. When administered alone, a 24-h treatment induced 27.4 and 34.2% apoptosis for etoposide and matrine treatment, respectively. When the same doses were administered to cells in combination, as shown in Fig. 6C,
compared with the control and either alone treated groups, the apoptosis ratio was significantly increased in the combination group (>70%; P<0.01). These results suggested that matrine sensitized Hep3B cells to epotoside-induced apoptosis.

**Discussion**

The p53 tumor suppressor gene serves a crucial roles in maintaining the integrity of the genome and the defense
against tumor metastasis and is mutated or deleted in ~50% of human cancer types (12). However, numerous tumors retain wild-type p53, suggesting that the above tumor cells contain abnormalities of other genes, including overexpression of the p53 negative regulator, MDM2. MDM2 overexpression has been reported in ~1/3 human cancer cells that contain wild-type p53 gene (27). Therefore, MDM2 can be considered as a proto-oncogene. The present study used a p53-deficient cell model, Hep3B cells, to investigate the anticancer activity of a known Chinese herbal medicine, matrine. It was found that matrine notably decreased the mRNA expression of MDM2, of its negative effector, MDM2. This p73-induced increase of p21 and PUMA may be inhibited by matrine, even though the reason why biological functions of p21 and PUMA were inhibited rather than activated in the matrine-treated Hep3B cells. Another important p73 transcriptional target for execution of mitochondria-dependent apoptosis, PUMA, was also not increased and rather decreased in the matrine-treated Hep3B cells. Although the present study did not investigate the molecular mechanism underlying the above remains to be elucidated. The most important observations of the present study were that matrine notably increased the sensitivity of the p53-deficient Hep3B cells to epotides-induced apoptosis. It is well known that epotides and numerous other anticancer alkylating drugs kill cancer cells through induction of DNA damage, the latter in turn increases immediate accumulation and activation of p53 family members through ATM-Chk2 and ATR-Chk1 pathways, which are activated by DNA double-strand breaks and single-stranded DNA, respectively (11,34). The activated p73, similar to p53, can bind to the promoters of p21 and PUMA, which in turn execute cell cycle arrest and mitochondria-dependent cell apoptosis (35). The activated p73 also triggers mRNA expression of its negative effector, MDM2. This p73-induced increase of MDM2 expression begins to compete with p73 for binding to the p300/CBP N-terminus and suppresses its biological functions (17). In addition to interacting with and inactivating p53, there is evidence to suggest that MDM2 can also interact with other molecules, including specific protein and RNA, which may serve a p53-independent role in oncogenesis (e.g. induction of IAP3) (33). Aside from regulating its downstream effectors, MDM2 itself can be modulated by various upstream signals. For example, cell growth factor-induced activation of phosphatidylinositol 3-kinase-Akt can phosphorylate MDM2-serine 166 and 186 in the cytoplasm, which in turn triggers translocation of MDM2 from the cytoplasm into the nucleus (36). For anther case, cellular stress and DNA damage can induce dephosphorylation of the central

Figure 5. Effect of epotides-increased and matrine-decreased MDM2 on the expression of IAP3. (A) Hep3B cells were incubated with either 100 µM epotides or 0.5 mg/ml matrine for the indicated duration. The protein expression levels of MDM2 and IAP3 were assessed by western blotting. (B) Normal and MDM2 overexpression Hep3B cells were treated with 0.5 mg/ml matrine for the indicated durations. The transfection efficiency of MDM2 and IAP3 in MDM2-overexpressing Hep3B cells were evaluated by western blotting. (C) The protein stability of IAP3 was detected using a pulse-chase assay following treatment with 0.5 mg/ml matrine. β-actin was used as a loading control. MDM, mouse double-minute; IAP, inhibitor of apoptosis protein; CHX, cycloheximide.
Figure 6. Apoptotic effect of matrine on Hep3B and MDM2-overexpressing Hep3B cells. (A) Hep3B and MDM2-overexpressing Hep3B cells were treated with 0.25–4.0 mg/ml matrine for 24 h. The cell viability was assessed using a 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide assay. The data are presented as the mean ± standard deviation (n=3; *P<0.05 and **P<0.01 compared with the corresponding control group; ∆P<0.05 and ∆∆P<0.01 compared with the untransfected Hep3B cells). (B) Hep3B cells were treated with 100 µM epotoside or 4.0 mg/ml matrine, alone or in combination, for 24 h. Cell apoptosis was analyzed by dual-parameter flow cytometry utilizing annexin V-FITC and propidium iodide, and the results of three independent experiments were pooled and average values are presented. Group a, control; group b, 100 µM epotoside; group c, 4.0 mg/ml matrine; group d, 100 µM epotoside + 4.0 mg/ml matrine. The data are presented as the mean ± standard deviation (n=3; **P<0.01 compared with the control group; ∆∆P<0.01 compared with the epotoside group). (C) Hep3B cells were treated with 100 µM epotoside or 4.0 mg/ml matrine. The protein expression levels of cleaved and total Caspase-3, -9 and PARP were assessed by western blotting. FITC, fluorescein isothiocyanate; PARP, poly ADP ribose polymerase.
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acidic domain of MDM2, which may result in inhibition of the nuclear translocation for MDM2, or accumulation in the cytoplasm (37). The present results suggested a functional role for MDM2 in tumor development, which extends their current understanding of resistance to chemotherapy in cancer.

In conclusion, the present study indicated that matrine causes cytotoxicity and induces apoptosis in p53-deficient Hep3B cells. The major underlying mechanism is the inhibition of the MDM2-AP3 pathway. Therefore, it is hypothesized that matrine may be an interesting candidate drug in the development of therapies against p53-defective cancer cells.

References
