Abstract. The aim of this study was to investigate the molecular mechanism of 8-chloroadenosine 3',5'-monophosphate (8-Cl-cAMP) in the inhibition of the growth and induction of apoptosis of multiple myeloma (MM) cells. Two MM-derived cell lines, RPMI-8226 and U266, were used. Cell viability, apoptosis induction and mitochondrial transmembrane potential were determined and the expression levels of cell cycle regulatory proteins (Cdk2, cyclin E, p27 and c-myc) and p38 mitogen-activated protein kinase (MAPK) protein were detected. Following treatment with 8-Cl-cAMP, the percentage of apoptotic cells increased in a concentration- and time-dependent manner and the mitochondrial transmembrane potential collapsed to reveal typical apoptotic features. Our data further demonstrated that 8-Cl-cAMP induced progressive phosphorylation of p38 MAPK and that the expression levels of p27 proteins in the MM cells were increased whereas those of c-myc were significantly decreased. Notably, the pro-apoptotic effect of 8-Cl-cAMP was largely prevented by a p38 MAPK inhibitor. Furthermore, knockdown of p27 was able to decrease the 8-Cl-cAMP-induced apoptosis in the MM cells. These results indicate that 8-Cl-cAMP-induced cellular apoptosis occurred in a concentration- and time-related manner with mitochondrial transmembrane potential collapse, increased expression levels of p27 proteins, and decreased expression levels of c-myc in MM cells. These findings demonstrate the potential of cAMP-modulating agents for use in the treatment of MM.

Introduction

Multiple myeloma (MM) is an incurable B-cell malignancy characterized by infiltrating, slow-growing plasma cells in the bone marrow that produce monoclonal immunoglobulin molecules (1). MM is more common than Hodgkin disease and acute leukemia and accounts for 1% of all cancers and slightly more than 10% of all hematological cancers. Few effective therapies have existed until recently. Alkylating agents and corticosteroids were the therapy of choice for several decades. A number of novel agents for MM have now become available, including the immunomodulatory drugs thalidomide and lenalidomide, as well as the proteasome inhibitor bortezomib (2-5). Although numerous therapeutic advances, including combined chemotherapy and hematopoietic stem cell transplantation, have been made to improve the survival of patients with MM, a high proportion of patients are not able to expect long term remission due to drug-resistant disease, minimal residual disease or infection. In order to overcome these limitations, certain new promising approaches are being widely studied (6,7).

As with unregulated proliferation, dysregulated apoptosis is also an important event in carcinogenesis. MM cells overexpress the apoptotic antagonists bcl-2 and bcl-xL, although they are not overtly hyperproliferative (8,9). The protection against apoptosis may be significant in the expansion and/or resistance of the MM cell clones to chemotherapy. Modulation of apoptosis with adjuvant therapy may have the potential to improve the clinical outcome of MM. In further studies, the safety and efficacy of a combination therapy should be investigated (10).

Cyclic adenosine monophosphate (cAMP), a well-known and ubiquitous chemical messenger, has an antiproliferative effect on the majority of cell types (11,12). The best studied of its derivatives is 8-chloroadenosine 3',5'-monophosphate (8-Cl-cAMP), which has demonstrated antiproliferative effects in vitro and in vivo and has been evaluated in phase I/II clinical trials (11,13,14). 8-Cl-cAMP is a site-selective analog of cAMP. It has been reported that 8-Cl-cAMP shows a potent growth inhibitory effect and has reverse-transforming activity in cancer cells (15-18). In the current study, we investigated the ability of 8-Cl-cAMP to induce apoptosis in two MM cell lines, RPMI-8226 and U266. Our results indicate that 8-Cl-cAMP-induced cellular apoptosis occurred in a concentration- and time-related manner with mitochondrial transmembrane potential collapse, increased expression levels of p27 proteins, and decreased expression levels of c-myc in MM cells. These findings demonstrate the potential of cAMP-modulating agents for use in the treatment of MM.
of p27 and decreased expression levels of c-myc. p27 knockdown was able to decrease the 8-Cl-cAMP-induced apoptosis of the MM cells, indicating that the apoptotic action occurred through a p27-dependent pathway.

Materials and methods

The study was approved by the Independent Ethics Committee of Shanghai Ninth People’s Hospital Affiliated to Shanghai Jiao Tong University School of Medicine, Shanghai, China.

Reagents and cell culture. 8-Cl-cAMP, propidium iodide (PI), rhodamine 123 (Rh123), SB202190 and other reagents were purchased from Sigma-Aldrich (St. Louis, MO, USA). p27 siRNA was designed by Dharmacon (Lafayette, CO, USA). Fetal bovine serum (FBS), RPMI-1640 medium and penicillin-streptomycin were obtained from Gibco-BRL (Gaithersburg, MD, USA). All antibodies were purchased from Santa Cruz Biotechnology, (Santa Cruz, CA, USA). The ECL kit was purchased from Amersham Pharmacia Biotech (Amersham, UK). An Annexin-V-FITC apoptosis detection kit, Oligofectamine and a mitochondrial membrane potential detection kit were purchased from Invitrogen (Eugene, OR, USA). The human myeloma cell lines RPMI-8226 and U266 (Shanghai Institute of Hematology, China) were cultured in RPMI-1640 medium supplemented with 10% FBS in a humidified atmosphere of 5% CO2 at 37°C.

Trypan blue exclusion assay. The effect of 8-Cl-cAMP on MM cell viability was measured by the trypan blue exclusion assay (19). RPMI-8226 and U266 cells were collected, mixed with an equal volume of PBS containing 0.4% trypan blue dye and manually counted. Actual cell numbers were calculated by multiplying by the dilution factor and were compared with the initial cell numbers. Cell viability (%) = viable cell numbers/total (viable + dead) cell numbers x 100.

Flow cytometric analysis of nuclear DNA distribution. Cells (2x10⁶) were collected, rinsed and fixed overnight with 70% cold ethanol. They were then rinsed with PBS, treated with 1 mg/ml RNase at 37°C for 30 min and stained with 250 µg/ml PI. The nuclear DNA contents were detected by flow cytometry (Beckman Coulter, Miami, FL, USA). All data were collected, stored and analyzed by MultiCycle software.

Flow cytometric analysis of mitochondrial membrane potential. After washing with PBS twice, 1-2x10⁵ RPMI-8226 cells were incubated with 10 µg/ml Rh123 at 37°C for 30 min. Subsequently, 250 µg/ml PI was injected into cells. Rh123 and PI staining intensities were determined by flow cytometry.

Western blot analysis. At appropriate time-points following treatment with 10 µmol/l 8-Cl-cAMP, the RPMI-8226 cells were collected. Protein extracts (100 µg) were loaded onto a 10% SDS-polyacrylamide gel, electrophoresed and transferred onto nitrocellulose membranes, which were subsequently stained with 0.2% Ponceau red to assure equal protein loading and transfer. After blocking with 10% non-fat milk powder, the membrane was incubated with primary antibody overnight at 4°C. The membrane was then washed with PBS and incubated with horseradish peroxidase-conjugated secondary antibody for 60 min at room temperature. The blots were again washed and the immunocomplex was visualized using the ECL kit.

Transfection of p27 siRNA and cell viability assay. The cells (1x10⁴ cells/well) were seeded in a 96-well plate, incubated for 24 h to enable them to attach to the bottom of the well, and then transfected with 80 nM p27 siRNA or control siRNA using Oligofectamine. Following transfection for 12 h, the cells were treated with 10 µmol/l 8-Cl-cAMP for 48 h. Cell growth was then determined by measuring the MTT dye absorbance of the living cells (20). Following exposure to 8-Cl-cAMP for the indicated time, 20 µl MTT solution (5 mg/ml in PBS) was added to each well and the plates were incubated for an additional 4 h at 37°C. The MTT solution in the medium was aspirated. To achieve solubilization of the formazan crystals formed in viable cells, 150 µl dimethylsulfoxide (DMSO) was added to each well and the absorbance at 570 nm was measured using a microplate reader (Bio-Rad, Hercules, CA, USA).

Statistical analysis. Each result in this study is representative of at least three separate experiments. Values represent the mean ± standard deviation (SD) of these experiments. The statistical significance was calculated with a Student’s t-test. P<0.05 was considered to indicate a statistically significant result.

Results

8-Cl-cAMP induced MM cell growth inhibition and apoptosis in a concentration- and time-related manner. To investigate...
the cytotoxic effect of 8-Cl-cAMP on RPMI-8226 and U266 cells, the cell viability following treatment with 8-Cl-cAMP at various concentrations (1, 3, 10, 30 µM) was determined by trypan blue exclusion assay. As shown in Fig. 1B, 8-Cl-cAMP reduced the cell viability of the RPMI-8226 and U266 cells in a concentration-dependent manner. Higher concentrations of 8-Cl-cAMP yielded more effective inhibition. The RPMI-8226 cells were more sensitive than the U266 cells to 8-Cl-cAMP. When treated with 30 µM 8-Cl-cAMP for 72 h, almost all RPMI-8226 cells died (Fig. 1A). The results of Annexin V-FITC and PI staining revealed that 8-Cl-cAMP induced apoptotic and necrotic cell death. The percentage of cell death was 19.27% in the RPMI-8226 cells treated with 30 µM 8-Cl-cAMP for 24 h (Fig. 2A). Since the reduction of mitochondrial transmembrane potential is often associated with apoptosis, we further investigated the effect of 8-Cl-cAMP on mitochondria. The treatment of RPMI-8226 cells with 8-Cl-cAMP resulted in an increase in the percentage of cells with a reduced mitochondrial transmembrane potential and increased apoptosis in a time-dependent manner (Fig. 2B). These features suggest that the 8-Cl-cAMP-induced MM cell death was due to apoptosis.

8-Cl-cAMP induces cell cycle arrest and apoptosis. To elucidate whether the apoptosis induced by 8-Cl-cAMP was due to cell cycle arrest, we evaluated the cell cycle distribution by flow cytometry. Flow cytometric analysis of the cellular DNA contents revealed that the sub-G1 phase ratio rose from 6.5 to 63.6% in the RPMI-8226 cells treated with 30 µM 8-Cl-cAMP for 48 h (Fig. 3A). The expression of the cyclin-dependent kinase (Cdk) inhibitor p27, which is involved in the arrest of the cell cycle in the G1 phase, was also evaluated. p27 levels, not detectable under basal conditions, were found to be increased following exposure to 10 µM 8-Cl-cAMP for 48 h (Fig. 3B). We also monitored the levels of other cell cycle-associated proteins. The levels of Cdk2 and cyclin E almost remained constant, but the expression levels of c-myc protein decreased in the RPMI-8226 cells exposed to 10 µmol/l 8-Cl-cAMP for 48 h (Fig. 3B). These results suggest that the 8-Cl-cAMP-induced cell cycle arrest was mediated by the expression of cell cycle-associated proteins. We also found that poly(ADP-ribose) polymerase (PARP) was cleaved following treatment with 8-Cl-cAMP (Fig. 3B).

8-Cl-cAMP induces apoptosis in MM cells through a p27-dependent pathway. Cell cycle associated proteins play a critical role in mediating the cytotoxicity induced by 8-Cl-cAMP, but their targets are unknown. To determine whether higher p27 levels were a critical factor contributing to the 8-Cl-cAMP-induced MM cell apoptosis, we knocked down p27 expression by RNA interference. Immunoblotting revealed that the transient introduction of p27 siRNA into the RPMI-8226 cells markedly reduced p27 expression levels compared with those in their control counterparts (Fig. 4A). The viability of the RPMI-8226 cells in the presence of 10 µM 8-Cl-cAMP was 14.7%, whereas that of the p27-knocked-down RPMI-8226 cells was 74.1% (Fig. 4B). Similarly, flow cytometric analysis revealed that the level of apoptosis of sub-G1 cells increased in the RPMI-8226 cells treated with 10 µM 8-Cl-cAMP, whereas the apoptosis of sub-G1 cells was almost undetectable in the p27-knocked-down RPMI-8226 cells (Fig. 4C). Thus, on the basis of the above results, p27 is functionally linked to the sensitivity of the RPMI-8226 cells to 8-Cl-cAMP-induced apoptosis.
cells is p27-dependent, and p27 may be the target of 8-Cl-cAMP‐induced apoptosis in MM cells is p27-dependent, and p27 may be the target of 8-Cl-cAMP-induced apoptosis in MM cells.

Figure 4. Role of p27 and p38 mitogen‐activated protein kinase (MAPK) expression. 8-Cl-cAMP has been shown to induce p38 MAPK phosphorylation in HL60 and HeLa cells (15,21). Therefore, we evaluated the effect of 8-Cl-cAMP on p38 MAPK phosphorylation in the MM cells. We identified that the treatment of RPMI-8226 and U266 cells with 8-Cl-cAMP was associated with a progressive increase of p38 MAPK phosphorylation that started after 48 h of incubation (Fig. 3B). Finally, we evaluated whether the pro‐apoptotic effect of 8-Cl-cAMP was dependent upon p38 MAPK activation. To this aim, we treated the two cell lines with 8-Cl-cAMP for 48‐72 h in the presence or absence of a selective p38 MAPK inhibitor (SB 202190) and analyzed the cell cycle distribution by flow cytometry as described previously. Notably, the inhibition of p38 MAPK largely prevented the pro‐apoptotic effect of 8-Cl-cAMP (Fig. 4D). These data strongly suggest that the pro‐apoptotic effect of 8-Cl-cAMP on RPMI-8226 and U266 cells is mediated by p38 MAPK.

Discussion

Following the identification of cAMP in 1958 by Rall and Sutherland, research focused for more than a decade on elucidating the role that this secondary messenger played in regulating metabolic pathways, as well as identifying the enzymes responsible for cAMP synthesis and catabolism (22-24). By the 1970s, however, cAMP was implicated as a regulator of cell growth (25-27), and several investigators reported that the elevation of cAMP levels induced proliferation arrest or cell death in susceptible normal or malignant lymphoid populations (28-33). The cAMP signaling pathway has emerged as a key regulator of hematopoietic cell proliferation, differentiation and apoptosis. The 8-chloro derivative of cAMP is a very potent cAMP analog that is under investigation as a potential chemotherapeutic agent. Phase I clinical studies in patients with solid tumors revealed its safety as well as evidence of clinical response (34).

In our study, we identified that 8-Cl-cAMP induced mitochondrial transmembrane potential collapse and apoptosis simultaneously in RPMI-8226 cells. Western blot analysis revealed that 8-Cl-cAMP (10 µmol/l)‐induced apoptosis was accompanied by upregulation of p27 and downregulation of c‐myc in RPMI-8226 cells. p27, also known as Cdk inhibitor 1B, is an enzyme which belongs to the Cip/Kip family of Cdk inhibitor proteins (35). It binds to and prevents the activation of cyclin E‐Cdk2 and cyclin D‐Cdk4 complexes, and thus controls the cell cycle progression at G1 (36). It is often referred to as a cell cycle inhibitor protein since its major function is to stop or slow down the cell division cycle (35-38). In the current study, we found that 8-Cl-cAMP induced the upregulation of the Cdk inhibitor p27. Therefore, we assessed the significance of p27. Fig. 4 shows that p27 knockdown inhibited the 8-Cl-cAMP‐induced decreases in cell viability, cell cycle arrest and cell death. These results indicate that the 8-Cl-cAMP‐induced apoptosis in MM cells is p27‐dependent, and p27 may be the target of 8-Cl-cAMP.

The proto‐oncogene c‐myc is involved in the control of cell proliferation, apoptosis and differentiation, and its aberrant expression is frequently observed in human cancer (39). It has been previously reported that small‐molecule inhibitors of c‐myc suppress proliferation and induce apoptosis of promyeloctytic leukemia cells via cell cycle arrest (39,40). Although our data show that 8-Cl-cAMP induced the downregulation of c‐myc, the mechanism for this action is not well defined. We also hypothesize that c‐myc is significant in the mediation of 8‐Cl‐cAMP‐induced cytotoxicity.

In addition, we identified that the pro‐apoptotic effect of 8-Cl-cAMP was accompanied by a progressive increase of p38 MAPK phosphorylation. The p38 MAPK is a key regulator of cell survival (41,42) and has been implicated in the pro‐apoptotic effect of 8-Cl-cAMP in HL60 and HeLa cells. In conclusion, our data indicate that 8-Cl-cAMP has a potent antiproliferative effect on MM cell lines through multiple pathways.

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References


