Knockdown of CDK6 enhances glioma sensitivity to chemotherapy

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Abstract. Chemotherapy is widely used for the treatment of glioma. Given the high resistance of brain neoplasm tissues to chemotherapy, it is important to find new methods to improve the effects of chemotherapy. However, the molecular mechanisms underlying glioma resistance to chemotherapy are largely unknown. Here, we demonstrate that CDK6, a cell cycle regulator, is significantly upregulated in glioma cells, and the increasing expression of CDK6 correlates well with the grades of glioma malignancy. Using shRNA-mediated CDK6 knockdown, we found that the proliferation and survival of tumor cells were dramatically inhibited. Moreover, CDK6 knockdown in the U251 glioma cell line caused significant increase in the apoptosis of U251 cells treated with temozolomide (TMZ). Furthermore, CDK6 knockdown reduced the expression level of drug resistance genes such as MRP and MDR. These data indicate that CDK6 is an important mediator of glioma resistance to chemotherapy. Our findings provide a new strategy for the development of chemotherapy sensitizer.

Introduction

Glioma is the most common tumor of the brain and accounts for 45-55% of all brain tumors. Based on histological features (nuclear atypia, mitosis, microvascular enrichment and necrosis) and malignancy, glioma is classified into four grades (1). Grade 1 is innocent and curable with surgical resection; grade 2 is a low grade diffuse glioma and can progress into higher grades; grade 3 and 4 gliomas are malignant and have increased capability to proliferate and invade. The grade 4 glioma, glioblastoma multiforme (GBM), is the most aggressive.

Surgical resection is the primary therapy to treat glioma (2). However, surgical removal is very restrictive, thus it is difficult to completely remove diffusive tumor tissues, especially higher grade gliomas. These glioma tissues invade into the white matter and may diffuse along the ventricle surface, so other strategies such as chemotherapy are very important to remove cancer cells as completely as possible (3-5). Chemotherapy increases the life of GBM patients. Clinical statistics reveals that, chemotherapy treatment leads to a 15% reduction of mortality, and a 6% increase of 1-more year survival. However, the effect of chemotherapy on glioma is not satisfactory, largely due to three problems. The first problem is the diagnosis of diffusive glioma tissues, which are tiny and grow deep in the brain tissue and are hard to access. The second problem is the blockade of chemotherapy drug by the blood-brain barrier. The third problem is that glioma is highly tolerant under chemotherapy treatment. Understanding the fundamentals, especially the molecular mechanisms, is critical to increase the susceptibility of glioma to chemotherapy.

Tumor cells originated from cells with abnormal proliferation and dysregulation of the cell cycle (6,7). Molecular profiling studies showed that the genes controlling cell growth and restricting cell division are silenced, and genes promoting cell proliferation and facilitating cell cycle are overexpressed. Many genes that promote cell cycle are found to be increased in transformed tumor cells (8). In cell cycle, CDK4 and 6 phosphorylate tumor suppressor Rb and promote G1 progression and G1/S transition (9). CDK6 is arrested and inhibited by INK4 and cyclin D (10). The CDK4/6 signaling was found to be highly activated in tumor cells, including sarcoma and leukemia (11,12). Lam et al reported that 12 in 14 tumor samples showed increased expression of CDK6 (13). Moreover, mir-125b blocked G1/S transition and increased apoptosis by inhibiting CDK6 expression (14). Another microRNA mir-34a down-regulated CDK6 expression and significantly inhibited the proliferation of glioma cells (15). However, whether CDK6 mediates the chemotherapy resistance of glioma remains unknown.

In this study, we tested the hypothesis that increased expression and activation of CDK6 in glioma contribute to its resistance to chemotherapy treatment. Using immunohistochemistry, we found that the expression level of CDK6 correlated well with the tumor grade of gliomas. We then found that knocking down of CDK6 led to significant reduction on the cell cycle progression and proliferation of glioma cells. Finally, we found that glioma
cells transfected with CDK6 shRNA showed increased apoptosis when exposed to temozolomide (TMZ) compared to cells transfected with control scRNA. Moreover, CDK6 knockdown resulted in down-regulated expression of several genes relating to drug resistance. Our findings suggested that CDK6 is an important molecular determinant contributing to chemotherapy resistance.

**Materials and methods**

**Cell culture.** The widely used glioma cell line U251 (ATCC, Manassas, VA) was cultured for our study. Cells were cultured in Dulbecco's modified Eagle's medium (DMEM) (Invitrogen, Carlsbad, CA), supplemented with 10% fetal bovine serum (Thermo Fisher Scientific, Inc., Waltham, MA) and antibiotics (penicillin and streptomycin, 50 U/ml each) (Invitrogen). Cells were grown in 60 mm dish and incubated in a humidified incubator with 5% CO₂ at 37°C. Cells were inoculated at 1x10⁵ cells/dish, and were used at a confluence of ~75%. For chemotherapy treatment, cells were treated with TMZ (10 nM) for 24, 48 or 72 h.

**Collection of human glioma and normal tissue samples.** Surgically removed human glioma tissue samples and normal brain tissues were obtained frozen or paraffin-embedded from Changhi Hospital (China Secondary Military Medical University, Shanghai, China). Gliomas were graded by the Pathology Department of Changhi Hospital according to the World Health Organization grading system, presence and absence of nuclear atypia, mitosis, microvascular enrichment, and necrosis. Human normal brain tissues (mostly from the cortex) were obtained from patients with physical injuries to the brain. These specimens were collected from the patients registered at the above-mentioned hospitals, and written informed consent was obtained from the patients. The use of human tissues was approved by the ethics committees of the hospitals.

**RT-PCR.** Total RNA from mouse brain tissue or culture was extracted using TRIzol reagent (Invitrogen). The RNA was treated with DNase I to remove genomic DNA contamination and then reverse transcribed on a 96-well temperature cycler (ABI) using an M-MLV RT kit (Invitrogen). For PCR, a SYBR Green Premix Ex Taq kit (Takara) was used to prepare PCR solution and programs were run on Mastercycler Gradient PCR machine (Eppendorf) or Rotor-Gene 3000 Realtime Cycler (Corbett Research, Inc.). The primers used are: cdk6 forward, GCCGCC TATGGGAAAGTGTTC, cdk6 reverse, TTGGGTTGCTC GAAAGTCT. Mdr-1 forward, 5-TGGTTCAGGTGGCTCTG TGAT-3, mdr-1 reverse, 5-CTGTAGACAAACGATGACCTATC ACA-3; mpr forward, 5-GGCAAGAAGAAATAAACGCAGCTG AA-3; mpr reverse, 5-GGCTGTGTGTGTCCATAGGCAAT-3. For vector based shRNA construct, an shRNA with the selected targeting sequence was cloned into pSuper-GFP or pLVTHM. For transfection of U251 cells, 4 µg of shRNA or control plasmid was dissolved in TRIzol reagent, diluted in sample loading buffer and subjected to SDS-PAGE gel electrophoresis, then transferred to PVDF membrane (0.45 µm, PALL), and blocked by 0.05% TBST with 5% milk and then blotted with primary antibodies at 4°C overnight and secondary antibodies at room temperature for 1 h. Bound antibodies were detected by the ECL immunoblotting detection reagent (GE Healthcare) and exposed to X-ray films. Band densities were quantified by Gel-Pro Analyzer, and the densitometric results are shown. The relative amount of proteins was determined by normalizing the density value of target protein to internal control and the external control.

**RNAi and transfection.** To knockdown cdk6, three siRNA targeting sequences were designed using online siRNA Target Finder (Ambion). The three sets of siRNA was synthesized and annealed to form double strand siRNA and transfected into U251 cells by Lipofectamine 2000 (Invitrogen). The knockdown efficiency was validated by RT-PCR and Western blotting, and finally the most specific and efficient targeting sequence was used for functional assay. The sequence is: 5’-CATGTCGATCAAGACTTGA-3’. For vector based shRNA construct, an shRNA with the selected targeting sequence was cloned into pSuper-GFP or pLVTHM. For transfection of U251 cells, 4 µg of shRNA or control plasmid was dissolved in Opti-MEM and coated with Lipofectamine 2000 reagent for a 60-mm dish. Six hours later, the culture medium was replaced, and 48 h later, the cells were subjected for analysis.

**MTT assay.** Cells were cultured in a 96-well plate at a density of 5x10⁵. Two hours before examination, cell-counting kit-8 reagent (10 µl/well) was added into the wells and incubated for 2 h. The reaction was terminated by adding SDS solution (0.1% final concentration). Before reading, the plate was shaken by the supplied plate shaker and then read at 450 nm.

**Flow cytometry to examine cell cycle progression.** Flow cytometry assays were used to study the effect of knocking down CDK6 cell cycle progression. U251 cells were harvested, treated with 0.25% trypsin, washed in PBS, centrifuged (1500 rpm, 5 min) and resuspended with binding buffer. The cell density was adjusted to 3x10⁵ cells/ml. Cell suspension (195 µl) and 5 µl Annexin V/FITC were combined and incubated at RT for 10 min. Next, propidium iodide was added to a final concentration of 1 µg/ml and incubated at 37°C for 30 min. In each sample,
6x10^6 cells were assayed on FACSCalibur (Becton-Dickinson, Franklin Lakes, NJ) and the cell cycle phase distribution was analyzed by CellQuest Pro software (Becton-Dickinson).

Statistical analysis. The data are presented as mean ± SD. All statistical analysis were done using One-way ANOVA and SPSS software.

Results

Significant upregulation of CDK6 in glioma. Although CDK6 expression was reported to be increased in gliomas, it remains unclear whether the malignant grade correlates with the upregulation of CDK6 expression. To examine this, we collected 34 glioma samples at different grades and 4 samples of normal brain tissue. The mRNA of CDK6 showed good correlation with different grades of gliomas (normal tissue, 1.02±0.25; grade 1, 5.50±0.25; grade 2, 11.78±5.90; grade 3, 46.74±28.81; grade 4, 78.11±14.04, Fig. 1A and B). The expression change of CDK6 protein was also studied using Western blotting, and similar upregulation pattern was found in glioma samples (Fig. 1C and D). In normal tissue, the protein level of CDK6 was 16.81±3.25, whereas in grade 1 glioma, CDK6 expression was increased to 41.33±7.38, and in grade 2 glioma to 35.11±37.54. In highly malignant glioma, CDK6 expression was increased to 41.33±7.38, and in grade 4 glioma to 35.11±37.54. In normal brain tissue (Fig. 2A). In grade 1 glioma tissue, CDK6 was kept at much low level in normal brain tissue (Fig. 2A). In grade 1 glioma tissue, CDK6 was kept at much low level in grade 2 glioma, CDK6 was highly upregulated (Fig. 2C), whereas in more malignant grade 3 or 4 glioma, higher upregulation of CDK6 expression was observed (Fig. 2D).

We also employed microarray chips to do whole-genome comparison of gene expression profiles between glioma and normal tissues. Among those highly upregulated genes in glioma tissue, CDK6 also showed good correlation with glioma grades and survival time according to follow-up visit data of patients (data not shown).

Knockdown of CDK6 mRNA in glioma inhibits glioma growth. Increased high level of CDK6 may promote the G1 progression and G1/S transition, thus leads to uncontrolled cell division and proliferation, and finally tumorigenesis and malignancy (13,16,17). Although upregulation of CDK6 expression correlated with glioma grades, whether CDK6 contributed to the higher proliferation and invasion capability requires further determination.

To test this hypothesis, we first established a culture model of CDK6-knockdown glioma. Chemotherapy is an indispensable treatment for higher grade gliomas (4,18). However, the low permeability of drugs across blood-brain barrier reduced drug concentration in cerebral-spinal fluid. On the other hand, glioma cell are resistant to chemotherapy treatment, mainly due to high activity of DNA repair mechanism, tolerance to cell cycle check points, reduced uptake of drug and quick metabolism of drug molecules (18-20). In the above studies, we found that CDK6 expression is highly upregulated in glioma cells. As CDK6 is a transducer of cytoplasm signal to nuclear transcription activity and DNA repair mechanism, we speculated that the chemotherapy-resistance of glioma is mainly mediated by increased level of CDK6.

Knocking down cdk6 inhibited cell cycle progression, thus blocking glioma proliferation and tumor growth. To examine the function of cdk6 in cell growth, we used MTT assay to analyze U251 cells after shRNA transfection (Fig. 3C). Compared to control scRNA, cdk6 shRNA expression greatly reduced the MTT absorption value (scRNA, 0.613±0.075; cdk6 shRNA, 0.417±0.065; p<0.05, Fig. 3D and E), thereby leading to a great reduction of the ratio of cells in G2/M phase (scRNA, 25.93±0.50; cdk6 shRNA, 21.85±1.32; p<0.05, Fig. 3D and E). Knocking down cdk6 inhibited cell cycle progression, thus blocking glioma proliferation and tumor growth.

Enhanced chemotherapy effect of TMZ on CDK6-knockdown glioma. Temozolomide (TMZ) is an alkylating agent and effective for the treatment of higher grade glioma (5). We thus tested the effect of TMZ on glioma in vitro (Fig. 4). Compared to control reagent, TMZ showed inhibition of U251 cell growth even at 24 h after treatment (con, 0.111±0.012; TMZ, 0.098±0.010). At 48 h, significant inhibition of U251 cell growth was achieved with TMZ.
by TMZ can be observed (con, 0.286±0.036; TMZ, 0.186±0.006; P<0.01). Next, we compared the effect of combined treatment of U251 cells with TMZ and cdk6 knockdown to single treatment with TMZ. Interestingly, combined treatment of U251 cells with TMZ and cdk6 shRNA resulted in significant decrease of U251 cell growth at 48 h after treatment (0.155±0.018, p<0.05), lower than single treatment with either TMZ (0.186±0.006) or cdk6 shRNA (0.177±0.017). These results strongly suggested that increased expression of CDK6 contributed to the resistance of glioma to chemotherapy.

Reduced drug-resistance to chemotherapy in CDK6-knockdown glioma. The facilitation of TMZ-mediated chemotherapy toxicity on U251 glioma cell growth by CDK6 knockdown indicated that aberrant cell signaling of glioma cells underlies the resistance of glioma to chemotherapy. To test whether CDK6 knockdown had an effect on drug resistance-related genes, we analyzed the expression change of several key genes by RT-PCR and western blotting. MDR-1 is a transmembrane glycoprotein of the ATP-binding cassette superfamily (21). MDR-1 acts as a bump to transport drug molecules out of the cells, thereby reducing drug resistance.
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Drug toxicity to cancer cells. In cdk6-knockdown cells, MDR-1 expression was greatly down-regulated (shRNA, 0.44±0.18; scRNA, 3.07±0.49; sham, 3.09±0.27; p<0.01, Fig. 5A). MRP is another member of the ABC transporter superfamily (22). We also found that MRP expression in cdk6-knockdown U251 cells was lower than scRNA transfected cells (shRNA, 1.93±1.40; scRNA, 7.27±1.49; sham, 8.39±1.17; p<0.01, Fig. 5B).

Drug resistance can also result from high level of DNA repair activity. PARP is a DNA polymerase which plays an important role in DNA repair (23). PARP can detect DNA single-strand break and bind to the break sequence and initiate the repair process by the synthesis of a poly(ADP-ribose) chain and recruiting other DNA repair proteins. PARP1 is highly expressed in glioma (24), and its expression can be increased by TMZ treatment. In turn, the upregulated PARP1 protects glioma cells from DNA-damage induced cell cycle retention and apoptosis. In cdk6-knockdown cells, PARP1 protein was significantly down-regulated (scRNA, 623.77±15.36; cdk6 shRNA, 237.23±28.09; p<0.05, Fig. 5C and D). Conclusively, increased level of CDK6 resulted in enhanced expression of several key genes responsible for drug resistance, and CDK6 knockdown or inhibition was effective in reducing their expression, thereby reducing chemotherapy drug-resistance of glioma cells.

Discussion

Gene mutation and their aberrant expression are very common in tumor cells. Microarray studies revealed that in each type of tumor cells, more than one hundred genes exhibited abnormal expression or mutation (25). Thus, finding the key genes that participate in or regulate the tumorigenesis process has become an important task in cancer research. CDK6 regulates cell cycle progression, especially in the transition from G1 to S phase (9). We first demonstrated that CDK6 upregulation correlated with the malignant grades of glioma, which implied that the expression level of CDK6 might determine the proliferation and invasion capability. As an upstream regulator of cancer repressor Rb, CDK6 expression level is an indicator of cell division activity (16). CDK6 phosphorylates Rb and activates transcription factor E2F1/2, thus pushing cells into S phase and triggering DNA synthesis (6).

The control of G1/S transition is the first gate to cell division, thus becomes extremely important for cell cycle control. The highly upregulated CDK6 in glioma cells make the G1/S checkpoint permissive, thereby enabling cell proliferation. We found that cdk6 knockdown resulted in significant decrease of cell growth, indicating that CDK6 is an organizer of cell cycle, and higher expression of CDK6 may underlie the chemotherapy resistance of glioma. As a kinase, CDK6 activity can easily be inhibited by small molecules, which can pass across the blood-

Figure 5. Expression of drug-resistant genes is significantly down-regulated in CDK6-knockdown glioma cells. (A) Decreased expression of MDR1 in CDK6 knockdown U251 cells. Sham, empty vector without shRNA; scRNA, scrambled shRNA; shRNA, cdk6 shRNA. (B) Down-regulated expression of MRP in CDK6 knockdown U251 cells. (C) Protein level of PARP1 was significantly reduced in CDK6 knockdown U251 cells examined by western blotting. (D) Statistical analysis of western blotting result of PARP1. *p<0.05; **p<0.01.
brain barrier. Many drug-resistant genes are upregulated in glioma cells, some of which may be downstream genes of E2F transcription activator complex (26,27). Therefore, inhibiting CDK6 may have considerable effect on enhancing the sensitivity of glioma cells to chemotherapy drugs. In our study, we observed that combined treatment of glioma cells with chemotherapy drug TMZ and CDK6 shRNA significantly inhibited cell growth, which is more effective than single treatment either with TMZ or CDK6 shRNA.

The mechanism underlying chemotherapy-resistance of glioma has been intensively studied (18,19,21,22). Many drug-resistant genes have been found, which mainly belongs to DNA repair enzymes, small-molecule transmembrane transporters and drug metabolism proteins. Given their wide divergence and differential protein structures, it may not be practicable to interfere with a single drug-resistant gene. Therefore, finding the upstream regulators maybe more effective to increase drug uptake and toxicity. CDK6 knockdown resulted in significant down-regulation of MDR1, MRP and PARP, indicating that CDK6 is an upstream activator of these drug-resistance related genes, and should be a promising target to reduce drug-resistance of glioma cells. Future work will be needed to study how CDK6 overexpression leads to the upregulation of these genes.

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References