CENPE promotes glioblastomas proliferation by directly binding to WEE1

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Background: Glioblastoma (GBM) is one of the most lethal tumors. Even though radiotherapy and chemotherapy have been greatly developed, the survival of patients with GBM is still only about 16 months.

Methods: In this assay, centromere protein E (CENPE) expression levels were measured using quantitative real-time polymerase chain reaction (qRT-PCR) and Western blot (WB) assays. 5-ethynyl-2'-deoxyuridine (EDU) assay was used to assess the effect of CENPE on cell proliferation. The propidium iodide (PI) method was used to detect the cell cycle.

Results: CENPE expression was increased in GBM tissues and was associated with clinical stage and unfavorable overall survival in glioma patients. Inhibition of CENPE expression resulted in proliferation inhibition of U251 and U87 cell. Cell cycle assay showed that CENPE mainly regulates the G0–G1 and G2/M phase transitions. Then, we found that CENPE regulated the proliferation of GBM mainly through WEE1 G2 checkpoint kinase (WEE1) pathway and can bind to WEE1.

Conclusions: CENPE promotes GBM proliferation through WEE1 pathway and binding to WEE1.

Keywords: Centromere protein E (CENPE); glioblastoma (GBM); proliferation; WEE1 G2 checkpoint kinase (WEE1)

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Introduction

Use the GLOBOCAN 2018 estimates of cancer incidence and mortality produced by the International Agency for Research on Cancer, there are 296,851 new cancer cases and 241,037 million cancer deaths in brain, nervous system in 2018. Glioblastoma (also known as GBM) is a malignant grade IV tumor in which most tumor cells multiply and divide at any given time. GBM is usually classified as an isocitrate dehydrogenase (IDH)-wild type (WT) or IDH-mutant. IDH-WT GBM accounts for the majority, is more aggressive, and has a worse prognosis. With standard treatment, the median survival of adult GBM (IDH-WT) is approximately 11–15 months. Therefore, further understanding of the proliferation mechanism of glioma, especially GBM, can play an important role in the control and treatment of GBM.

Cell proliferation is achieved by mitosis. The most important process of mitosis is the cell cycle. The separation of chromosomes is closely related to the correct assembly of microtubules. In addition, microtubules play an important role in maintaining cell shape, cell movement and intracellular organelles movement. Centromere protein E (CENPE) is one of human kinetochore proteins of 312 kDa. Studies have shown that knockdown of CENPE or knockdown of xeroderma pigmentosum group A (XPA)-binding protein 2 (XAB2) can inhibit cell proliferation, and CENPE is directly regulated by XAB2. In addition, the expression of CENPE is closely related to the proliferative capacity of tumors. In incurable
castration-resistant prostate cancer (CRPC), knockout or pharmacological inhibition of CENPE significantly inhibits tumor growth (6). In gastric cancer cells, when treated with mycophenolic acid (MPA), the expression of CENPE was significantly inhibited, and cell proliferation was inhibited (7). In human neuroblastoma cell lines, knockdown of CENPE results in inhibition of cell proliferation in vitro and in vivo (8).

Methods

Cell lines and patient samples

All of the GBM tumors and paired normal tissue (NT) samples were collected from Tongde Hospital of Zhejiang Province. The study was approved by institutional/regional/national ethics committee/ethics board of Ethics Committee of Zhejiang Institute of Traditional Chinese Medicine (No. XMSC2018048). The U251 and U87 human GBM cell lines were purchased from the American Type Culture Collection (Manassas, VA, USA) and cultured in DMEM high glucose medium (Hyclone, USA) with 10% fetal bovine serum (FBS; Hyclone, USA) and 1% penicillin/streptomycin (Gibco, USA), then, incubated at 37 °C in an atmosphere of 5% CO₂.

Small interfering RNA (siRNA)

2.5×10⁵ cells were seeded in 6-well plates, cultured overnight (about 12 hours), then, cells were transfected with siRNA (siCENPE, 50 nmol/L) or negative control siRNA (NC, 50 nmol/L) using Lipofectamine 3000 (Thermo Fisher Scientific, Waltham, MA, USA) follow the manufacturer’s protocol. The sequence of the siRNA is: CENPE, 5'-CAUCAACUUAUUACGUUAA-3', reverse, 5'-TGCCCTGGGTATACTCCCAA-3'; glyceraldehyde-3-phosphate dehydrogenase (GAPDH): forward, 5'-ACAACCTTGTTATCGTGAAG-3', reverse, 5'-GCCATCACGACAGTTTC-3'. Relative expression levels were calculated using the 2⁻ΔΔCT method.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR) analyses

TRIZol reagent (Invitrogen, Thermo Fisher Scientific, Waltham, MA, USA) was used to extract total RNA from tissue samples or cells. An RT reagent kit with gDNA Eraser (TaKaRa Biotech, Dalian, China) was used to reverse transcribe RNA into cDNA. The quantitative real-time PCR (qRT-PCR) reaction was then performed using SYBR Green (TaKaRa Biotech, Dalian, China) and detected with ABI 7500 System (Applied Biosystems, Foster City, CA, USA). Each sample were repeated three times. All PCR primers were purchased from Invitrogen (Thermo Fisher Scientific, Waltham, MA, USA): CENPE: forward, 5'-GATTCTGCCATAAGGCTACAA-3', reverse, 5'-TGCCCTGGGTATACTCCCAA-3'; glyceraldehyde-3-phosphate dehydrogenase (GAPDH): forward, 5'-ACAACCTTGTTATCGTGAAG-3', reverse, 5'-GCCATCACGACAGTTTC-3'. Relative expression levels were calculated using the 2⁻ΔΔCT method.

Western blot (WB) analysis

Cell and tissue samples were both lysed in RIPA lysis buffer (Beyotime, Haimen, China) containing 1% protease inhibitor PMSF (Beyotime, Haimen, China). Then, a total of 40 micrograms of protein was separated by 10% SDS-PAGE, and the protein was transferred to a PVDF membrane (Roche Applied Sciences, Indianapolis, IN, USA). Membranes were blocked in 5% BSA (Boster, AR0004, Wuhan, China) for 1 hour at room temperature and then incubated overnight at 4 °C with the primary antibody (anti-CENPE, anti-WEE1 anti-GAPDH). After incubation with a suitable HRP-conjugated secondary antibody (Cell Signaling Technology, Danvers, MA, USA) for 1 hour at room temperature, the membrane was visualized with an ECL kit (Beyotime, Haimen, China), and then membranes were visualized using LI-COR Odyssey system (Odyssey; LI-COR Biosciences). The proteins were visualized using a chemiluminescence method (ECL plus WB detection system; Amersham Biosciences, Foster City, CA, USA). The analysis included mouse monoclonal anti-GAPDH (Boster, BM3876, Wuhan, China), rabbit anti-CENPE (Abcam, ab133583, Cambridge, UK), mouse anti-WEE1 (Abcam, ab23354, Cambridge, UK).

Coimmunoprecipitation (co-IP)

After cells were harvested, the 200 μL cell lysate was incubated with ChIP-grade protein G magnetic beads (CST, Danvers, MA, USA) and 5 μL antibodies, shook overnight at 4 °C, wash five times with cold PBS [contain 1% PMSF (Beyotime, Haimen, China)] for 5 minutes each time, suspended in 2x SDS-PAGE loading buffer and then detected by WB with relevant antibodies. The antibodies used for IP were rabbit anti-CENPE (Abcam, ab133583, Cambridge, UK), rabbit anti-WEE1 (Abcam, ab23354, Cambridge, UK).

Cell proliferation assay

The 647-flow cytometry assay kit (GeneCopoeia, Rockville,
MD, USA) was used to detect cell proliferation. After the cells were transfected with small RNA for 24 hours, 5-ethynyl-2’-deoxyuridine (EDU) (10 μM) was added to each well for 12 hours, and then stained and flow-detected according to the manufacturer’s protocol.

Cell cycle analysis
Propidium iodide (PI) (Beyotime, Haimen, China) staining is used to determine the cell cycle. After 24 hours of cell small RNA interference in the 6-well plate, each well was separately digested and collected into an EP tube. The cells were permeabilized overnight with PBS at 75% cold ethanol, then washed twice with PBS and stained with PI solution for 15 minutes at room temperature [50 μg/mL PI in PBS, 0.1% Triton-X100, 50 μg/mL RNase A (TaKaRa Biotech, Dalian, China)]. Cells were detected within 3 hours using flow cytometer.

Analysis of data from The Cancer Genome Atlas (TCGA)
Difference expression of CENPE in normal and gliomas and survival in glioma patients was analyzed in the website Gliovis (http://gliovis.bioinfo.cnio.es/). And the Graphpad Prism 6 software was used to analyze data.

Results

Expression of CENPE in collected GBM tissues
We examined the mRNA expression levels of CENPE in GBM and paracancerous samples by RT-qPCR and analyzed protein expression by WB. mRNA assay results showed that CENPE mRNA levels were significantly elevated in GBM tissues compared to paired normal brain tissue samples (Figure 1A, P<0.0001). WB results showed that the expression level of CENPE in GBM tissues was higher than that in paired NTs, and the expression of CENPE was almost undetectable in paired normal brain tissues (Figure 1B).

Expression of CENPE in TCGA low-grade gliomas and glioblastomas (GBMLGG)
To further determine the expression of CENPE in gliomas, we analyzed data on gliomas in the TCGA database, including GBMLGG. It was found that as the grade of glioma increased, the expression of CENPE also increased (Figure 2A). At the same time, we also found that the expression of CENPE was reduced in GBM with IDH1 mutation (Figure 2B). At the same time, the O6-methylguanine-DNA methyltransferase (MGMT) promoter methylation tissue also showed low CENPE expression (Figure 2C). We then analyzed the survival curves of gliomas with high and low expression of CENPE and found that patients with high expression of CENPE had a worse prognosis (Figure 2D).

CENPE promotes proliferation of glioma cell lines
To examine the effect of CENPE on glioma cell proliferation, we performed an EDU assay on CENPE-interfering cell lines. First, we designed siRNA to knock down CENPE in glioma cell lines. CENPE mRNA and protein levels significantly reduced after RNA interference (Figure 3A,B). EDU experiments demonstrated that proliferation of glioma cells was inhibited after CENPE knockdown (Figure 3C,D) (P<0.0001).
**Figure 2** Expression of CENPE in TCGA GBMLGG tissues. (A) CENPE expression of different grade of glioma; (B) CENPE expression of different IDH1 status of glioma; (C) CENPE expression of different MGMT promoter methylation of glioma; (D) Kaplan-Meier analysis of GBMLGG RNA-seq data from the TCGA database. ****, P<0.0001. CENPE, centromere protein E; TCGA, The Cancer Genome Atlas; GBMLGG, low-grade gliomas and glioblastomas; IDH1, isocitrate dehydrogenase; MGMT, O6-methylguanine-DNA methyltransferase; WT, wild type.

**Inhibition of CENPE mainly inhibits G0–G1 transition**

To further understand the regulatory mechanisms of CENPE, we examined changes in cell cycle after interference with CENPE. We found that cells in the G0–G1 phase increased significantly, while cells in the G2-M phase were significantly reduced in siCENPE-U251 group compared with the NC-U251 group (*Figure 4A,B*). U87 cells also showed the same trend (*Figure 4C,D*). This result indicated that the knockdown of CENPE has the greatest impact on G0–G1 transition.

**CENPE affects cell cycle by affecting the expression of WEE1**

To detect what genes are downstream of CENPE, we selected genes that are thought to be involved in the cell cycle. We found that CCNA1 and CCNE1 genes did not change significantly after interference with CENPE, while the expression of WEE1 gene decreased significantly (*Figure 5A,B*). These results suggested that CENPE may affect the transformation of G1 phase to S phase by affecting the expression of WEE1. Then we further studied the RNA-seq data of CENPE and WEE1 in the TCGA database, and found that CENPE has a significant correlation with the expression of WEE1 (*Figure 5C*). At the same time, the mRNA data of the samples we collected also obtained similar results (*Figure 5D*).

**CENPE is in direct contact with WEE1**

In order to further study the relationship between CENPE and WEE1, the co-IP experiments were performed. At first, we found that endogenous WEE1 proteins could be coimmunoprecipitated by CENPE antibodies in U251 cell line.
Figure 3 CENPE silencing inhibits proliferation in GBM cells. (A) qPCR analysis of CENPE in U251 and U87 cells transfected with control siRNA or siCENPE; (B) immunoblot analysis of CENPE in U251 and U87 cells transfected with control siRNA or siCENPE; (C) cell proliferation rates as determined by EDU assays in U251 and U87 cells transfected with control siRNA or siCENPE; (D) statistical analysis of EDU results of U251 cells and U87 cells. ***, P<0.001 (mean ± SEM). CENPE, centromere protein E; GBM, glioblastoma; qPCR, quantitative polymerase chain reaction; EDU, 5-ethynyl-2'-deoxyuridine; siRNA, small interfering RNA; SEM, standard error of mean; NC, negative control.

(Figure 6A). Then, to further determine the accuracy of this result, we performed a co-IP using the WEE1 antibody and found that the CENPE protein was also detected (Figure 6B).

Discussion

Rapid proliferation is one of the important characteristics of tumor cells, and the higher the degree of malignancy, the more proliferative ability (9,10). A better understanding of the mechanism of rapid tumor proliferation can play an important guiding role in the treatment of tumors. In addition, finding biomarkers related to rapid tumor proliferation plays an important role in the diagnosis and treatment of tumors. At present, in addition to its ability to control proliferation, many studies found that it could also be used as a biomarker for many tumors. In invasive ductal carcinoma (IDC), Studies have found that CENPE is a potentially key gene associated with IDC (11). In multiple myeloma, side population (SP) cells exhibit high CENPE expression characteristics (12). In prolactin pituitary tumors, CENPE is one of 5 genes associated with the pathological classification (13). In our study, we found that the
proliferation of glioma cells was significantly inhibited after knocking down CENPE. At the same time, the expression of CENPE was not only significantly correlated with stage, but also significantly associated with IDH1 mutation. These results reflect the important characteristics of CENPE as a biomarker for tumors.

In order to ensure that the cells can be completely and normally split into two daughter cells, the cells develop three cell cycle checkpoints (G1/S phase checkpoint, G2/M phase checkpoint, spindle assembly checkpoint), allowing cell repair damaged DNA to maintain genomic integrity (14,15). In cancer cells, the G1 checkpoint mechanism of many cancer cells is defective, so, cells can proliferate rapidly and the G2 checkpoint and the spindle assembly checkpoint are far more important than normal cells (16,17). Wee1 was first identified in fission yeast, and wee1 deficiency resulted in premature mitotic entry and replication of smaller-sized yeast (18). In addition, the expression of WEE1 is also associated with the proliferation of many tumors. In glioma cells, miR-526b-3p inhibits tumor cell proliferation by inhibiting the expression of WEE1 (19). In liver cancer, upregulated DLX6-AS1 promoted cell proliferation, migration, and invasion through increasing expression of WEE1 via targeting miR-424-5p (20).

In addition, WEE1 inhibitors have been used in clinical trials. WEE1 inhibition by AZD1775 abrogates the G2-M arrest, allowing cells with a deregulated G1 into premature mitosis, resulting in mitotic catastrophe and apoptosis through unrepaired DNA damage (21). A phase I trial of AZD1775 in 25 patients with advanced solid tumors has established the maximum tolerated dose (MTD) to be 225 mg twice a day over 2.5 days per week for 2 weeks every 21 days (22). This is the first reported phase I study showed safety and efficacy of single-agent, AZD1775, especially in BRCA mutant patients (22). Furthermore, one phase II study provided clinical evidence that AZD1775 enhanced carboplatin efficacy in TP53-mutant tumors (23). Since small molecule inhibitors of WEE1 also have been used in clinical trials of glioma (24) and WEE1 is closely related to the proliferation of glioma (19). Therefore, the CENPE gene, which regulates the expression of WEE1, may also be used as a target gene for clinical treatment of glioma.

**Conclusions**

CENPE expression was increased in GBM tissues and was associated with clinical stage and unfavorable overall
Figure 5 CENPE affects cell cycle by affecting the expression of WEE1. (A) qPCR results showed WEE1 expression reduced after knockdown of CENPE (P<0.001); (B) Western blot results showed WEE1 expression reduced after knockdown of CENPE; (C) CENPE has a significant correlation with the expression of WEE1, data from the TCGA database; (D) CENPE has a significant correlation with the expression of WEE1, data from the collected sample. ***, P<0.001 (mean ± SEM); CENPE, centromere protein E; WEE1, WEE1 G2 checkpoint kinase; qPCR, quantitative polymerase chain reaction; TCGA, The Cancer Genome Atlas; SEM, standard error of mean; ns, no significant difference; NC, negative control; GAPDH, glyceraldehyde-3-phosphate dehydrogenase.

Figure 6 CENPE is in direct contact with WEE1. (A) Endogenous WEE1 was immunoprecipitated by anti-CENPE antibodies in U251 cells; (B) endogenous CENPE was immunoprecipitated by anti-WEE1 antibodies in U251 cells. CENPE, centromere protein E; WEE1, WEE1 G2 checkpoint kinase; IP, immunoprecipitation; IB, immunoblotting.
survival in glioma patients. Inhibition of CENPE expression resulted in proliferation inhibition of GBM cell. CENPE promotes GBM proliferation through WEE1 pathway and binding to WEE1.

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**Footnote**

Conflicts of Interest: The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was approved by institutional/regional/national ethics committee/ethics board of Ethics Committee of Zhejiang Institute of Traditional Chinese Medicine (No. XMSC2018048).

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