The Role of CHMP4C on Proliferation in the Human Lung Cancer A549 Cells

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Abstract

The human lung cancer has high incidence rate and mortality among the carcinoma. The research on enhancing the efficacy of therapy for lung cancer is significant. A recent research found that as a subunit of ESCRT-III, CHMP4C functioned to retard abscission timing to coordinate midbody resolution and prevent accumulation of DNA damage in the abscission checkpoint through phosphorylated by AuroraB. In the current study, we evaluated the possible mechanism of the effects of CHMP4C inhibition on cell cycle and cell survival in A549 cells. We found that CHMP4C knockdown caused lagging S phase in cell cycle through enhancing the phosphorylation of Rb, raising the expression of cyclin B1-cdc2 and suppressing the activation of cyclin A. Meanwhile, CHMP4C deletion depressed cell survival via decreasing cell viability and increasing caspase 3/7 activity. This study may promote new significant reference and advance for the mechanism underlying specific function of CHMP4C as well as further research on enhancing therapy effect on non-small lung cancer.

Keywords

The Human Lung Cancer, A549, CHMP4C, siRNA

1. Introduction

The human lung cancer is generally categorized into small cell lung cancer (SCLC) and non-small cell lung cancer (NSCLC) based on clinical. NSCLC accounts for 84% in the human lung cancer, with the 5-year survival rate of approximate 16.6% [1] [2]. Hence, how to improve efficiency in diagnostics and therapy poses a critical challenge. It is essential to find an effective therapeutic target for achieving better therapy effects.

CHMP4C was found recently a critical role in the abscission timing checkpoint which governs the cytokinetic...
abscission to monitor the final separation of the two daughter cells to insure the property abscission time [3]-[5]. Besides, CHMP4C has lower expression level in normal tissues and increased expression in cancer cells. In one research, Genome wide association studies (GWAS) meta-analysis identifies three new susceptibility loci for ovarian cancer, and CHMP4C is one of them [6]. CHMP4C also plays an important role in exosome production [7] [8].

As so far, the research on CHMP4C in cancer therapy is just beginning. In the present study, we want to explore the possible mechanism of the function of CHMP4C on cell cycle and cell survival in A549 cells. This study may promote new significant reference and advance for the specific function of CHMP4C, and further research on enhancing therapeutic effect on non-small lung cancer. CHMP4C may also be a meaningful drug target gene for lung cancer therapy.

2. Materials and Methods

2.1. Cell Culture

The human lung cancer A549 cell line (Cell resource center, Peking union medical college) was cultured in RPMI-1640 containing 10% fetal bovine serum (Invitrogen, USA), incubated in 37°C humidified incubator with 5% CO₂.

2.2. siRNAs Transfection

A549 cells grew to the concentration of 80% and were transfected with siCHMP4C (Ambion) or negative control siRNA (Ambion). All transfection of A549 cells with siRNA were performed using Lipofectamine RNAiMAX reagent. 48 h after transfection, the target gene knockdown efficiency was checked by western blotting, and the cells were harvested for testing.

2.3. Western Blot

The cells were harvested and lysed in RIPA lysis buffer (Thermo Scientific Pierce). The protein was collected at 12,000 g for 15 min at 4°C and measured by BCA protein assay kit (Pierce, USA). Equal amounts of protein were separated on 10% SDS-polyacrylamide gels and blotted on nitrocellulose membranes for western blot analysis. The membranes were blocked in 5% nonfat milk and then incubated with the primary antibodies: anti-CHMP4C antibody (Abcam), anti-Rb (Ser795) antibody, anti-cycinB1 antibody, anti-cdc2 (Tyr15) antibody, anti-cycinA antibody and anti-β-actin antibody (Cell Signaling Technology). Membranes were washed in tris-buffered saline containing 0.5% tween-20 and then incubated with goat anti-rabbit IgG (Cell Signaling Technology) or goat anti-mouse IgG (Cell Signaling Technology) conjugated to horseradish peroxidase for 1 h at room temperature. The membranes were detected using Chemiluminescence liquid (Thermo Scientific Pierce) according to the manufacturer’s protocol and analyzed by the Image J software (Bio-Rad).

2.4. Cell Cycle Assay

Cells were harvested and treated with Vibrant Dyecycle green stain (Invitrogen, USA) at 37°C for 30 min in dark place. And then cell cycle was analyzed on a flow cytometer.

2.5. Cell Viability Assay

After treatment, cells were mixed with 10% volume of alamarblue reagent (Invitrogen, USA) and incubated for 30 min at 37°C, protected from direct light. Results were recorded using fluorescence at 570/585 nm (excitation/emission).

2.6. Cell Apoptosis Assay

Cells were washed twice with cold PBS and then resuspended in 1 × binding buffer (BD) at a concentration of 1 × 10⁶ cells/ml. After stained with PE AnnexinV and 7-AAD (BD), samples were analyzed by flow cytometry.

2.7. Caspase 3/7 Activity Assay

Cells were seeded in 96 well plates at a concentration of 1 × 10⁴ cells/ml. 48 h after transfection with siCHMP4C,
100 μl of Caspase-GloR 3/7 reagent (Promega) was added to each well and incubated at room temperature for 30 min, luminescence values were read out (Microplate multimode reader, Turner Biosystems).

2.8. Statistical Analysis

Data analysis are presented as the mean ± SE of three independent experiments. Differences in mean values between groups were determined by students’ t test using Microsoft Excel (Microsoft Campus, Redmond, WA, USA) with p-value < 0.05 indicating statistical significance.

3. Results

3.1. CHMP4C Inhibition Decreased Cell Viability and Promoted Cell Apoptosis by Enhancing Caspase 3/7 Activity

To detect the reason for CHMP4C deletion depressing cell survival, we first employed cell viability and apoptosis assays in A549 cells with or without CHMP4C silencing. The results indicated that CHMP4C knockdown decreased cell viability and produced apoptosis compared to normal CHMP4C expression.

We next checked the activity of caspase 3/7 and then ascertained that silencing of CHMP4C increased the activity of caspase 3/7. We demonstrated that CHMP4C promoted cell apoptosis through activation of caspase 3/7 (Figure 1).

**Figure 1.** CHMP4C inhibition decreased cell viability and promoted cell apoptosis by enhancing caspase 3/7 activity. (a) After transfection for 48 h, siRNAs transfection efficiency of A549 cells was assayed by western blotting; (b) The A549 cells were transfected with siCHMP4C for 48 h. The cells were stained with alamarBlue. The fluorescence intensity was read out at 570/585 nm (excitation/emission); (c) Apoptosis assay of A549 cells with or without siCHMP4C transfection; (d) Relative caspase 3/7 viability of A549 cells treated by siCHMP4C for 48 h was measured. Data are presented as the mean ± SE of three independent experiments, ∗p < 0.05.
3.2. CHMP4C Silencing Retards S-Phase of the Cell Cycle through Regulation of Rb, Cyclin B1-cdc2 and Cyclin A

From the result that CHMP4C silencing delayed S-phase in the cell cycle, we wished to explore the possible mechanism underlying CHMP4C and cell cycle. A549 cells were depleted of CHMP4C for 24 h and the delayed S phase was checked (Figure 2(a)).

And then, we analyzed cell cycle relevant protein by western bolt. Rb plays a critical role in G1/S transition, and we found CHMP4C deletion enhanced the phosphorylation of Rb in the ser795 site. Meanwhile the result also showed that inhibition of CHMP4C expression upregulated CyclinB1 and cdc2 which are major in controlling G2/M checkpoint [9] [10]. However, CyclinA was observed downregulated by knockdown of CHMP4C, which is necessary to S phase progression (Figure 2(b)).

4. Discussion

Lung adenocarcinoma often occurs in women and non-smokers. Their five-year survival rate was only about 15%. Chemo-radiotherapy remains the major treatment method for lung adenocarcinoma but limited success has been achieved in the clinical. With further investigation on the treatment of lung adenocarcinoma, targeted therapy has been intensively researched to improve therapy effect [11].

CHMP4C is a subunit of ESCRT-III (Endosome sorting complex-III) and plays a major role at the cytokinetic abscission checkpoint by delaying mitosis to prevent premature and accumulation of DNA damage. One recent research reported that CHMP4C is overexpressed in cancer tissues and may have relation with poor prognosis of ovarian cancer.

Current investigations on CHMP4C mainly focus on its function endosome generation [12]. But, the mechanism of its function in cellular response is not clear. In the current study, we explored the mechanism underlying CHMP4C and cell survival as well as cell cycle.

In the apoptosis cysteine aspartic acid-specific protease (caspase) signaling pathway, caspase-8, caspase-9, and caspase-10 act at upstream, which couple cell death stimuli to the downstream caspase-3 and caspase-7. Caspase-3 and caspase-7 are both activated universally during apoptosis and play key effector roles in apoptosis.
in mammalian cells [13]. Our results demonstrated that inhibition of CHMP4C promoted cell apoptosis via enhancing the activity of caspase-3 and caspase-7.

Rb (Retinoblastoma gene) regulates cell proliferation by governing G1/S transition in the cell cycle [14]. The cyclinB1-cdc2 kinase is critical in regulating the G2/M transition. Overexpression of cyclinB1-cdc2 can promote G2/M transition [15] [16]. Cyclin A is associated with the E2F transcription factor control and is required for S-phase in cell cycle. Inhibition of cyclin A can delay the G1 to S transition and prevent DNA replication [17]. In the study, we found that CHMP4C depletion enhanced Rb phosphorylation, increased cyclinB1-cdc2 expression and reduced cyclin A expression, which in turn led to lagging S-phase.

5. Conclusion

In conclusion, our results further reveal that CHMP4C deficiency disturbed cell cycle progress and enhanced cell death as well as its mechanism, suggesting a new therapy target for the treatment of lung cancer.

References
