Cysteine-rich intestinal protein 1 silencing alleviates the migration and invasive capability enhancement induced by excessive zinc supplementation in colorectal cancer cells

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Abstract: Cysteine-rich intestinal protein 1 (CRIP1) is overexpressed in colorectal cancer (CRC) tissues and functions as an oncogene in regulating the migration and invasion of CRC cells. However, the underlying mechanism is unclear. CRIP1 has a role in zinc absorption and functions as an intracellular zinc transport protein. Here, we aimed to focus on the function of zinc and its underlying mechanism in CRC and determine whether CRIP1 promotes invasion and CRC metastasis through excessive zinc-induced epithelial-mesenchymal transition (EMT) by affecting the phosphorylated glycogen synthase kinase (GSK)-3beta. The results showed that ZnSO₄ (Zn²⁺) supplementation in medium increased the labile intracellular zinc content. Furthermore, excessive Zn²⁺ supplementation activated the GSK3/mTOR signaling pathway in both SW620 and LoVo cells, and excessive Zn²⁺ supplementation promoted migration, invasion, and EMT of SW620 and LoVo cells. This migration promotion was alleviated by the specific mTOR inhibitor rapamycin, indicating that the GSK3/mTOR signaling pathway was involved in this process. CRIP1 silencing increased the labile intracellular zinc content and inhibited EMT and GSK3/mTOR signaling pathway. CRIP1 silencing alleviated the zinc supplementation effects on migration, invasion, EMT, and GSK3/mTOR signaling pathway. In conclusion, excessive Zn²⁺ promotes migration and invasion capabilities of SW620 and LoVo cells through GSK3/mTOR signaling pathway-induced EMT.

Keywords: Excessive zinc, epithelial-mesenchymal transition, cysteine-rich intestinal protein 1, migration and invasion capabilities

Introduction

Colorectal cancer (CRC), that is, the development of cancer in the colon or rectum, is the second most commonly diagnosed cancer in women, the third most commonly diagnosed cancer in men, and the fourth most common cause of cancer death after lung, stomach, and liver cancers [1]. The morbidity and mortality rates are high in developing countries [2-4] and CRC greatly threatens human life and health. CRC cells have a strong ability to migrate and invade, leading to recurrence, which finally results in death of patients. Therefore, factors that may affect the invasion and CRC metastasis should be evaluated.
tein, CRIP1 is involved in zinc absorption [13-15]. Considering CRIP1’s important role in regulating CRC invasion, metastasis, and zinc absorption, we predicted that zinc might play a role in regulating CRC invasion and metastasis. This study aimed to demonstrate the predicted role of zinc in regulating CRC invasion and metastasis and determine its related underlying mechanism.

Zinc has been reported to induce an increase in phosphorylated (p) glycogen synthase kinase (GSK)-3beta level, which could be inhibited by rapamycin, a specific mechanistic target of rapamycin kinase (mTOR) inhibitor in SH-SY5Y neuroblastoma cells [16]. GSK-3beta can bind to and phosphorylate Snail at two consensus motifs to dually regulate the function of Snail, triggering the epithelial-mesenchymal transition (EMT) of epithelial cells by repressing the E-cadherin expression [17, 18]. However, whether zinc can affect the EMT of tumor cells remains unclear. At present, we discuss the role of zinc and CRIP1 in regulating the EMT of CRC. In addition, we discuss whether CRIP1 promotes CRC invasion and metastasis through zinc-regulated EMT by affecting the phosphorylation of GSK-3beta.

Materials and methods

Cell culture

The human colon cancer cell lines SW620 and LoVo were purchased from America Type Culture Collection (Manassas, VA, USA) and cultured in the condition according to the instruction of supplier.

siRNA interference

The siRNA oligonucleotides for targeting the CRIP1 gene (siCRIP1, sense: 5’-GCAACAAGGA-GGUGUACUUTT-3’) and negative control siRNA (siNC) were obtained from GenePharma (Shanghai, China). Cells were transfected with the 100 nM siCRIP1 or siNC using the Lipofectamine™ 2000 (Promega, Madison, WI, USA).

ZnSO₄ supplementation and experimental group

To detect the effect of ZnSO₄ supplementation on the cell survival rate, SW620 and LoVo cells were seeded on a 6-well plate and treated with 0.5, 5, 10, 50, 100, or 200 µM ZnSO₄ for 24 h. To determine the effects of ZnSO₄ supplementation on the levels of p-mTOR and p-GSK-3beta, SW620 and LoVo cells were seeded on a 6-well plate and treated with 5, 10, or 50 µM ZnSO₄ for 24 h, or with 5 µM ZnSO₄ for 12, 24, or 48 h. To determine the effects of 50 µM ZnSO₄ and/or 50 nM rapamycin on migration and invasion capabilities, SW620 and LoVo cells were seeded on a 6-well plate, treated with 50 µM ZnSO₄, 50 µM ZnSO₄ plus 50 nM rapamycin, or 50 nM rapamycin for 24 h, and harvested for Transwell assays and western blotting. To determine the effects of CRIP1 silencing on the level of p-mTOR and p-GSK-3beta, SW620 and LoVo cells were seeded on a 6-well plate and transfected with siCRIP1 and siNC for 48 h. To determine the effects of CRIP1 silencing on the function of ZnSO₄, SW620 and LoVo cells were seeded on a 6-well plate, pre-transfected with siCRIP1 and siNC for 24 h, and subsequently treated with 5 µM ZnSO₄ for 24 h.

Cell proliferation assay

After being treated with 0.5, 5, 10, 50, 100, or 200 µM ZnSO₄ for 24 h, cell proliferation was monitored using the CellTiter 96® AQueous One Solution Cell Proliferation Assay (Promega) following the manufacturer’s protocol.

Labile intracellular zinc content

At the end of different treatment, labile intracellular zinc in SW620 and LoVo cells treated as described above was monitored using Zinquin ethyl ester (Abcam, Cambridge, MA, USA). Briefly, cells were harvested and washed three times with phosphate-buffered saline (PBS) to remove extracellular Zn prior to labelling with Zinquin. Then, cells were suspended in PBS and 5 × 10⁶ cells/ml were seeded in black-bottom 96-well plates, incubated with 25-uM Zinquin at 37°C for 30 min. The fluorescence intensity was determined using a multiplate reader SuPerMax 3000FA (Shanghai Flash spectrum biological technology co. LTD) at 368 and 490 nm, respectively. The results in relative light units were obtained from raw data minus the reagent blank and expressed as the percentage of fluorescence-intensity controls.

Western blotting

The total protein isolation, protein concentration measurement, and western blot were carried out according to the method described previ-
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The primary antibodies, anti-CRIP1, anti-mTOR, anti-mTOR (phospho S2448), anti-GSK-3beta, anti-GSK-3beta (phospho Y216), were purchased from Abcam. The primary antibody of internal reference, glyceraldehyde-3-phosphate dehydrogenase (GAPDH), was purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA).

Transwell assay

After treated with different condition, transwell assays for cell migration or invasion were carried out according the method described previously [5]. Finally, the number of migrated or invasive cell was counted in five randomly selected fields under a phase contrast microscope (magnification: 200×; IX71, Olympus, Tokyo, Japan). Assays were performed in triplicate. The number of invasive or migrated cells was adjusted by dividing with the survival rate.

Statistical analysis

All results are presented as the mean ± standard deviation and were analyzed using the Statistical Package for the Social Science (SPSS) version 19.0 software (IBM SPSS, Armonk, NY, USA). Intergroup comparisons of more than two groups were conducted using one-way analysis of variance (ANOVA), followed by post hoc tests of the least significant difference for multiple pairwise comparisons. Intergroup comparisons of two groups were conducted using Student’s t-test. A P-value of < 0.05 was considered to indicate a statistically significant difference.

Results

Zn²⁺ supplementation increased the labile zinc levels in the CRC cells

The CRC cell lines SW620 and LoVo were treated with different concentrations of ZnSO₄ (Zn²⁺) for 24 h, and the cell survival rate was calculated using cell proliferation assay. The results showed that 0.5 µM Zn²⁺ supplementation did not decrease the survival rate of both SW620 and LoVo cells (Figure 1A). The inhibition rate of 5-, 10-, and 50-µM Zn²⁺ on SW620 and LoVo cell proliferation was < 5% (Figure 1A). The cell survival rate was clearly inhibited after 100-, 125-, 150-, or 200-µM Zn²⁺ supplementation (Figure 1A). Although 50-µM Zn²⁺ can significantly inhibit the survival rate of SW620 and LoVo, the survival rate remains to be > 95%. Because we investigated the effects of Zn²⁺ supplementation on migration and invasion capabilities, 5-, 10-, and 50-µM Zn²⁺ concentrations that cannot obviously suppress cell proliferation were selected for the following assays.

After treating with 5-, 10-, and 50-µM Zn²⁺ supplementation in the medium for 24 h, the labile intracellular zinc content was increased in both the SW620 and LoVo cells, with 50-µM Zn²⁺ having the most obvious promotion effect (Figure 1B). Following this, SW620 and LoVo cells were treated with 50-µM Zn²⁺ supplementation in the medium for 12, 24, and 48 h. Our findings indicate that 50-µM Zn²⁺ supplementation can increase the labile intracellular zinc content from 12 to 48 h, reaching a peak at 24 h (Figure 1C and 1D). These results all indicate that zinc supplementation can increase the labile intracellular zinc content in CRC cells. The specific mTOR inhibitor rapamycin cannot affect the labile intracellular zinc content with or without 50-µM Zn²⁺ supplementation in the medium (Figure 1E). However, the labile intracellular zinc content was increased in siCRIP1 transfected cells with or without 50-µM Zn²⁺ supplementation in medium (Figure 1F).

Zinc supplementation activated the GSK3/mTOR signaling pathway in the CRC cells

After the treatment with 5-, 10-, and 50-µM Zn²⁺ supplementation in medium for 24 h, the phosphorylation levels of GSK-3beta and mTOR were increased in both the SW620 and LoVo cells, with 50-µM Zn²⁺ having the most obvious promotion effect (Figure 2A). Following this, SW620 and LoVo cells were treated with 50-µM Zn²⁺ supplementation in medium for 12, 24, and 48 h. Our findings indicate that 50-µM Zn²⁺ supplementation can increase the phosphorylation levels of GSK-3beta and mTOR in a time-dependent manner (Figure 2B and 2C). These results indicate that zinc supplementation can activate the GSK3/mTOR signaling pathway in the CRC cells.

Zinc supplementation promotes migration, invasion, and EMT activation through the GSK3/mTOR signaling pathway

After being treated with 50-µM Zn²⁺ for 24 h, the migration and invasion capabilities of SW620 and LoVo cells were examined using the Transwell assays. The results showed that a
50-µM Zn²⁺ treatment could increase the number of migratory and invasive cells (Figure 3A). In addition, we found that a 50-µM Zn²⁺ supplementation significantly decreased the protein level of the epithelial marker, E-cadherin, and correspondingly increased the levels of mesenchymal marker, vimentin (Figure 3B). In contrast, the specific mTOR inhibitor rapamycin alleviated the effects of Zn²⁺ supplementation on the migration and invasion capabilities and EMT markers (Figure 3A and 3B). Compared to the cell group, rapamycin can suppress migration and invasion capabilities and the expression of EMT markers (Figure 3A and 3B). These results indicate that a 50-µM Zn²⁺ supplementation promotes migration, invasion, and EMT of SW620 and LoVo cells and that the GSK3/mTOR signaling pathway was involved in this process.

**CRIP1 silencing inhibits EMT and GSK3/mTOR signaling pathway**

Our previous study indicated that CRIP1 silencing could effectively inhibit migration and invasion of CRC cells [5]. To further determine whe-
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CRIP1 silencing suppressed invasion and CRC metastasis by inhibiting the EMT, we detected the expression levels of EMT markers and the phosphorylation levels of p-GSK-3beta and p-mTOR after transfection with siCRIP1 or siNC. Compared to the siNC transfection group, siCRIP1 transfection significantly increased the protein level of the epithelial marker, E-cadherin, and correspondingly decreased the levels of mesenchymal marker, vimentin (Figure 4A). In addition, the phosphorylation levels of GSK-3beta and mTOR decreased in both SW620 and LoVo cells after being transfected with siCRIP1 compared to those transfected with siNC (Figure 4B). These results all indicate that CRIP1 silencing inhibits EMT and GSK3/mTOR signaling pathway.

CRIIP1 silencing alleviated the effects of zinc supplementation on the migration, invasion, EMT, and GSK3/mTOR signaling pathway

To further investigate the relationship between CRIP1 and zinc supplementation in regulating migration, invasion, EMT, and GSK3/mTOR signaling pathway, SW620 and LoVo cells were treated with 50-µM Zn²⁺ after being transfected with siNC or siCRIP1 for 24 h. The results of Transwell assays showed that the number of migratory and invasive cells in the siCRIP1 transfection and zinc supplementation group (siCRIP1+Zn²⁺) was less than that in the siNC transfection and zinc supplementation group (siNC+Zn²⁺) (Figure 5A and 5B). The results of western blotting showed that the protein level
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*Figure 3.* Zinc supplementary in medium promoted migration, invasion, and EMT, and the specific mTOR inhibitor rapamycin alleviated the effects of $\text{Zn}^{2+}$ supplementation. A: The effects of 50-µM $\text{Zn}^{2+}$ supplementation on the migration and invasion capabilities of SW620 and LoVo cells. B: The effects of 50-µM $\text{Zn}^{2+}$ supplementation on the expression levels of EMT markers. The representative graphs are shown on the left. The statistical results are shown on the right. *P < 0.05.
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of the epithelial marker, E-cadherin, was increased in the siCRIP1+Zn²⁺ group compared to that in the siNC+Zn²⁺ group (Figure 5C). In addition, the level of mesenchymal marker vimentin and the phosphorylation levels of GSK-3beta and mTOR were decreased in the siCRIP1+Zn²⁺ group compared to those in the siNC+Zn²⁺ group (Figure 5C and 5D). These results indicated that CRIP1 silencing alleviated the effects of zinc supplementation on migration, invasion, EMT, and GSK3/mTOR signaling pathway.

Discussion

Zinc, an essential micronutrient, is required for the activity of > 300 enzymes, including key kinases, proteases, and phosphatases. The benefits of zinc supplementation have been widely accepted. In addition, its tumor-suppressing role and the association between zinc deficiency and cancer have been widely reported [19-22]. Our previous study indicated that CRIP1 is overexpressed in the CRC tissues and cell lines, and CRIP1 functioned as an oncogene in regulating migration and invasion of CRC cells [5]. CRIP1 has a role in zinc absorption and functions as an intracellular zinc transport protein [13-15]. These results suggested that an overactivity of intracellular zinc transport may also be involved in cancer development. CRC cells have a notable capability to migrate and invade; therefore, we focused on the function and underlying mechanism of excessive zinc supplementation on CRC cell migration and invasion capabilities.

First, 50-µM Zn²⁺ supplementation in medium increased the labile intracellular zinc content and enhanced the migratory and invasive capabilities of SW620 and HT29 cells, indicating that 50-µM Zn²⁺ supplementation elicited the effects on CRC cell migration and invasion. However, our present study showed that the survival rate of SW620 and LoVo cells was clearly inhibited after 100-, 125-, 150-, or 200-µM Zn²⁺ supplementation, indicating that excessive Zn²⁺ supplementation inhibits the effects on CRC cell growth. These inhibiting effects of Zn²⁺ supplementation have also been reported previously [23]. Although 50-µM Zn²⁺ can significantly inhibit the survival rate of SW620, the survival rate remains > 95% in our study. Thus, the effects of 50-µM Zn²⁺ supplementation on the cell survival rate are limited. In other words,
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50-µM Zn²⁺ supplementation cannot obviously suppress the cell growth. This result is consistent with the previous study [23]. In addition, the number of invasive or migrated cells was...
adjusted by dividing with the survival rate. Therefore, the conclusion that 50-µM Zn\textsuperscript{2+} supplementation promotes CRC cell migration and invasion cannot be mediated by its limited inhibiting effect on CRC cell growth. Zinc is an essential trace element with cofactor functions in a large number of proteins, such as zinc finger proteins [24]. However, some zinc finger proteins functioned as oncogene [25, 26], but some as tumor suppressor [27]. Therefore, we predicted that different concentrations of labile intracellular zinc may activate various proteins, resulting in different changes in the cells. These may explain the double-edged role in the development of CRC. Zinc concentration in the cell culture medium can simulate normal physiological concentrations of zinc. As such, 50-µM Zn\textsuperscript{2+} supplementation in a cell culture medium can stimulate a zinc excess condition, which suggests that an excess of zinc at some concentrations may play a cancer-promoting role in the development of CRC. This conclusion can be supported by the reports in Hep-2 tumor cells [28]. External zinc has also been reported to stimulate proliferation of tumor Hep-2 cells [28]. These results indicated that zinc does not always play a beneficial role. Clinicians and patients may need to reconsider the practice of using zinc supplements. However, further in vivo studies should be conducted to confirm this finding. In addition, only one concentration of Zn\textsuperscript{2+} was investigated in this study. The function of excessive zinc supplementation on CRC cell migration and invasion should be further confirmed by more concentration of Zn\textsuperscript{2+} in other CRC cells.

Second, we found that excessive zinc supplementation activated the GSK3/mTOR signaling pathway in SW620 and LoVo cells. This regulatory mechanism has also been reported in SH-SY5Y neuroblastoma cells [16]. GSK-3beta can bind to and phosphorylate Snail at two consensus motifs to dually regulate the function of Snail, which triggers the EMT of epithelial cells through repressing E-cadherin expression [17, 18]. Due to a 50-µM Zn\textsuperscript{2+} supplementation that enhanced the EMT of SW620 and LoVo cells, we predicted that zinc enhanced EMT of SW620 and LoVo cells through the activation of GSK3/mTOR signaling pathway, which was confirmed by our results. We found that the effects of zinc supplementation on the EMT were blocked by rapamycin, a specific mTOR inhibitor. Moreover, rapamycin alleviated the effects of a 50-µM Zn\textsuperscript{2+} supplementation on the migratory and invasive capabilities of SW620 and LoVo cells. Cancer-associated EMT is known to contribute to increased invasiveness and metastasis [29]. Based on these results, we predicted that excessive Zn\textsuperscript{2+} would promote the migratory and invasive capabilities of SW620 and LoVo cells through GSK3/mTOR signaling pathway-induced EMT. Our previous study demonstrated that CRIP1 silencing inhibited the migratory and invasive capabilities of SW620 and LoVo cells [5] and that CRIP1 silencing inhibited the EMT and GSK3/mTOR signaling pathway. In addition, CRIP1 silencing alleviated the effects of zinc supplementation on migration, invasion, EMT, and GSK3/mTOR signaling pathway. Furthermore, the labile intracellular zinc content was increased in siCRIP1 transfected cells with 50-µM Zn\textsuperscript{2+} supplementation in medium. As the intracellular zinc transport role of CRIP1, we predicted that CRIP1 affect GSK3/mTOR signaling pathway by increasing the zinc transport to the target protein. However, other zinc transporters also existed in the CRC cells, such as G protein-coupled receptor 39 [30]. Moreover, many enzymes or signaling pathways could also be regulated by zinc [20, 28, 31, 32]; thus, the underlying mechanism in which excessive Zn\textsuperscript{2+} promotes the migratory and invasive capabilities of SW620 and LoVo cells may be more complex than the results indicated in our report.

In conclusion, our results indicated that excessive zinc supplementation enhanced the migratory and invasive capabilities of SW620 and LoVo cells, and GSK3/mTOR signaling pathway-induced EMT was involved in this regulation. In addition, CRIP1 silencing alleviated the effects of excessive zinc supplementation on migration, invasion, EMT, and GSK3/mTOR signaling pathway. Based on these results, we predicted that CRIP1 promoted EMT through zinc-induced p-GSK-3beta in the CRC. Although further studies in vivo are needed to confirm the function and underlying mechanisms of zinc on CRC cell migratory and invasive capabilities, our data reveal a novel molecular basis for the understanding and clinical treatment of CRC.

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Disclosure of conflict of interest

None.

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