Norcantharidin induces ferroptosis via the suppression of NRF2/HO-1 signaling in ovarian cancer cells

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Abstract. Increasing evidence has indicated a crucial role of ferroptosis in ovarian cancer (OC). Norcantharidin (NCTD), a normethyl compound of cantharidin, is extensively used in clinical practice as an optional anticancer drug. However, whether NCTD leads to ferroptosis in OC has not been previously explored, at least to the best of our knowledge. In the present study, the effect of NCTD on SKOV3 and OVCAR-3 cells was evaluated. The experimental data of the present study revealed that NCTD significantly suppressed SKOV3 and OVCAR-3 cell viability in a concentration- and time-dependent manner. The results of Cell Counting Kit-8 assay revealed that NCTD treatment decreased SKOV3 and OVCAR-3 cell viability. In comparison, pre-incubation with ferrostatin-1 (Fer-1) significantly reversed the NCTD-induced reduction in SKOV3 and OVCAR-3 cell viability; however, no changes in cell viability were observed when the SKOV3 and OVCAR-3 cells were treated with NCTD, in combination with the apoptosis inhibitor, Z-VAD-FMK, the ferroptosis inhibitor, necrostatin-1, and the autophagy inhibitor, 3-methyladenine. Additionally, it was observed that NCTD markedly enhanced reactive oxygen species production and malondialdehyde and ferrous ion levels in the SKOV3 and OVCAR-3 cells; however, pre-incubation with Fer-1 abolished these effects. Flow cytometry also demonstrated a significant increase in cell death following treatment of the SKOV3 and OVCAR-3 cells with NCTD; however, pre-incubation with Fer-1 also reversed these effects. In vivo experiments demonstrated that NCTD significantly reduced tumor volume and weight. More importantly, it was revealed that nuclear factor erythroid 2-related factor 2 (NRF2), heme oxygenase 1 (HO-1), glutathione peroxidase 4 (GPX4) and solute carrier family 7 member 11 (xCT) expression levels were significantly decreased following NCTD treatment. Collectively, NCTD may represent a potent anticancer agent in OC cells, and NCTD-induced ferroptotic cell death may be achieved by inhibiting the NRF2/HO-1/GPX4/xCT axis.

Introduction

Ovarian cancer (OC) poses a serious threat to women's health and the incidence of ovarian cancer has increased to 11.7 per 100,000 women per year. Moreover, 30% of patients are diagnosed with advanced illness due to the recurrence of cancer and chemotheraphy resistance (1). Tumor metastasis is one of the main characteristics of OC and the most important cause of morality in patients bearing advanced tumors (2). Among patients with OC, ~90% do not survive due to metastasis-related complications (3). However, patients often suffer from cancer recurrence due to chemotheraphy resistance after the initial treatment (4). Therefore, elucidating the underlying mechanisms through which cancer metastasis is regulated is of utmost importance (5).

Ferroptosis is an iron-dependent, lipid peroxidation-driven cell death cascade that is critical for the progression of anticancer therapies (6,7). Prostaglandin-endoperoxide synthase 2 (PTGS2), also known as cyclooxygenase-2 (Cox-2), is the key enzyme that catalyzes prostaglandin biosynthesis (8). It mainly functions both as a peroxidase and a dioxygenase (8). During ferroptosis, the significant upregulation of PTGS2 has been identified (9). Glutathione-specific γ-glutamylcyclotransferase 1 (CHAC1) has been reported to decrease glutathione (GSH) levels and enhance cystine-starvation-induced ferroptosis (10). Both PTGS2 and CHAC1 are markers of ferroptosis (8).

The transcription factor, nuclear factor erythroid 2-related factor 2 (NRF2), has been suggested as a major regulator of intracellular oxidation homeostasis and lipid peroxidation (11,12). There is increasing evidence to indicate that NRF2 is strongly associated with the process of ferroptosis (11,12). The silencing of NRF2 may significantly reduce the expression of solute carrier family 7 member 11 (SLC7A11; xCT) and heme oxygenase 1 (HO-1) (12,13). xCT is a key gene that has been suggested to contribute to ‘iron overload-ferroptosis’ (12). The knockdown of xCT expression has been reported to result in reduced cystine-dependent GSH peroxidase activity and increased reactive oxygen species (ROS) and malonaldehyde (MDA) production, subsequently resulting in cellular ferroptosis (13). GSH is a necessary substrate for

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glutathione peroxidase 4 (GPX4), which is a key antioxidant enzyme (14). The depletion of GPX4 has been demonstrated to disrupt the balance of oxygen homeostasis and result in ferroptosis (14). In addition, NRF2 has also been revealed to directly interact with GPX4, which ultimately results in ferroptosis, by inducing intracellular antioxidant system damage (15). Therefore, targeting NRF2 may be useful for the induction of ferroptosis-related therapy in cancer patients.

Norcantharidin (NCTD), a normethyl compound of cantharidin, is extensively used in clinical practice as an optional anticancer drug in China, considering advantages of easy synthesis, potent activity as compared with cantharidin, and limited side-effects (16,17). In recent years, increasing evidence has demonstrated that NCTD significantly suppresses tumor cell proliferation and migration both in vitro and in vivo (16,18). However, whether NCTD suppresses OC via ferroptosis has not been previously reported, at least to the best of our knowledge.

In the present study, the specific function and underlying mechanisms of NCTD were first explored in SKOV3 and OVCAR-3 cells. The findings presented herein may prove to be of therapeutic value for patients with OC.

Materials and methods

Cells and cell culture. SKOV3 and OVCAR-3 human OC cell lines were purchased from Procell Life Science & Technology Co., Ltd. (cat. nos. CL-0215 and CL-0178). Short tandem repeat (STR) analysis was performed to confirm cell line authentication. SKOV3 or OVCAR-3 cells were cultured in McCoy's 5A or RPMI-1640 culture (Cytiva) supplemented with 10% fetal bovine serum (FBS; Invitrogen; Thermo Fisher Scientific, Inc.), streptomycin (100 mg/ml; Invitrogen; Thermo Fisher Scientific, Inc.) and penicillin (100 U/ml; Invitrogen; Thermo Fisher Scientific, Inc.) at 37°C in a humidified atmosphere, containing 5% CO2.

Cell Counting Kit-8 (CCK-8) assay. In brief, the SKOV3 and OVCAR-3 cells were seeded in 96-well plates at a density of 1x10^4 cells/well overnight. The SKOV3 and OVCAR-3 cells were then incubated with 0.25, 0.5, 1, 2, 4, 8, 16, 32 and 64 µg/ml NCTD (cat. no. C7632-25MG; Sigma-Aldrich; Merck KGaA) for 24 h at 37°C. Subsequently, 10 µl CCK-8 reagent (Beijing Solarbio Science & Technology Co., Ltd.) was added to each well and the cells were incubated for 4 h at 37°C. Cell viability was determined using a BD FACSCalibur system (BD Biosciences), 1 ml of DAPI (Beijing Solarbio Science & Technology Co., Ltd.) at 4˚C for 15 min. The intracellular MDA and Fe2+ contents were calculated according to a previous study (19): (Test group-NC)/(control group-NC) x100. The cell survival rate was calculated as the OD570nm of the necrosis inhibitor, necrostatin-1 (Nec-1; cat. no. HY-15760; MedChemExpress), 10 µM of the necrosis inhibitor, necrostatin-1 (Nec-1; cat. no. HY-15760; MedChemExpress), and 10 µM of the autophagy inhibitor, 3-methyladenine (3-MA; cat. no. HY-19312; MedChemExpress), for 1 h at 37°C. Subsequently, the SKOV3 and OVCAR-3 cells were treated with 10 and 20 µg/ml NCTD for an additional 24 h at 37°C. CCK-8 assay was carried out as described above.

2',7'-Dichlorofluorescin diacetate (DCFH-DA) staining. In brief, the SKOV3 and OVCAR-3 cells were seeded in 6-well plates at a density of 10^5 cells/well overnight. The SKOV3 and OVCAR-3 cells were then pre-incubated with or without 1 µM Fer-1 (cat. no. HY-100579; MedChemExpress) at 37°C for 1 h followed by treatment with 10 or 20 µg/ml NCTD for a further 24 h at 37°C. Furthermore, the SKOV3 and OVCAR-3 cells were incubated with 20 µM H2O2 for 24 h at 37°C. Subsequently, 1 ml DCFH-DA (cat. no. HY-D0940; MedChemExpress) was added to each well at a final concentration of 10 µM for 20 min at 37°C. After washing with 2 ml serum-free DMEM culture medium (HyClone; GE Healthcare Life Sciences), 1 ml of DAPI (Beijing Solarbio Science & Technology Co., Ltd.) was added to each well at a final concentration of 100 ng/ml for 20 min at 37°C. Subsequently, all cells were washed with 2 ml serum-free DMEM culture medium (HyClone; GE Healthcare Life Sciences) and the fluorescence was observed under a fluorescence microscope (x20; Olympus Corporation).

Flow cytometric assay. The SKOV3 and OVCAR-3 cells were seeded in 6-well plates at a density of 10^6 cells/well overnight. The SKOV3 and OVCAR-3 cells were then pre-incubated with or without 1 µM Fer-1 at 37°C for 1 h, followed by treatment with 10 or 20 µg/ml NCTD for an additional 24 h at 37°C. Subsequently, cell death was analyzed using an Annexin V/7-AAD double staining kit (cat. no. KGA1026; Nanjing KeyGen Biotech Co., Ltd.) according to the manufacturer's instructions. Briefly, the SKOV3 and OVCAR-3 cells were centrifuged at 1,000 x g for 3 min at 4°C and washed with PBS (Beijing Solarbio Science & Technology Co., Ltd.) thrice. The cells were then re-suspended using 500 µl 1X Annexin V Binding Buffer (cat. no. KGA1026; Nanjing KeyGen Biotech Co., Ltd.). Subsequently, 5 µl of the Annexin V/7-AAD reagent was added, and the cells were incubated for 10 min at room temperature. Following a 1-h incubation, cell death was determined using a BD FACSCalibur system (BD Biosciences), and data were analyzed using ModFit software version 4.1 (Verity Software House, Inc.). Annexin V/7-AAD+ cells were considered alive cells (Q4), Annexin V+/7-AAD- cells were considered early apoptotic cells (Q3), and Annexin V+/7-AAD+ cells were considered late apoptotic and necrotic cells (Q2).

Quantification of MDA, ROS, GSH and ferrous ion (Fe2+) contents. The SKOV3 and OVCAR-3 cells were seeded in 6-well plates at a density of 1x10^5 cells/well overnight. The SKOV3 and OVCAR-3 cells were then pre-incubated with or without 1 µM Fer-1 at 37°C for 1 h, followed by treatment with 10 or 20 µg/ml NCTD for a further 24 h at 37°C. Subsequently, 10^5 cells were treated in 100 µl RIPA buffer (Beijing Solarbio Science & Technology Co., Ltd.) at 4°C for 15 min. The intracellular MDA and Fe2+ contents were then
quantified using a Micro MDA assay kit (cat. no. BC0025; Beijing Solarbio Science & Technology Co., Ltd.) and Iron assay kit (cat. no. MAK025; Sigma-Aldrich; Merck KGaA), respectively, according to the manufacturer's instructions.

For the tumor tissues, MDA, ROS, GSH and Fe²⁺ levels were quantified using a Micro MDA assay kit (cat. no. BC0025; Beijing Solarbio Science & Technology Co., Ltd.), a Reactive Oxygen Species assay kit (cat. no. CA1401; Beijing Solarbio Science & Technology Co., Ltd.), a Micro Reduced Glutathione assay kit (cat. no. BC1175; Beijing Solarbio Science & Technology Co., Ltd.) and an Iron assay kit (cat. no. MAK025; Sigma-Aldrich; Merck KGaA), respectively, according to the manufacturer's instructions.

Reverse transcription-quantitative PCR (RT-qPCR). Total RNA was isolated from the SKOV3 and OVCAR-3 cells using RNAzol (Vigorous Biotechnology Beijing Co., Ltd.), according to the manufacturer's protocol. The concentration and purity of the RNA samples were determined by measuring the optical density ratio, OD260/OD280 using a Nanodrop One Ultra Micro Ultraviolet Spectrophotometer (ND-ONE-W(A30221); Thermo Fisher Scientific, Inc.). RT-qPCR was performed using a Takara PrimeScript™ One Step RT-PCR kit Ver 2.0 (cat. no. RR055A; Takara Bio, Inc.) according to the manufacturer's instructions. The following PCR reagents were used: 20 µl RNase Free ddH₂O, 25 µl 2X 1 step buffer, 2 µl PrimeScript 1 step enzyme mix, 1 µl forward primer, 1 µl reverse primer and 1 µl RNA template. The cycling conditions were as follows: 50˚C for 30 min, 94˚C for 2 min, 30 cycles of 94˚C for 30 sec, 55˚C for 30 sec, and 72˚C for 1 min, followed by 72˚C for 10 min. GAPDH was used as an internal control. Relative mRNA expression was normalized to GAPDH using the 2⁻ΔΔCt method (21). The primers used in the present study are listed in Table I.

Western blot analysis. Proteins were isolated from the SKOV3 and OVCAR-3 cells using a total protein extraction kit (Beijing Solarbio Science & Technology Co., Ltd.). A BCA protein assay kit (Pierce; Thermo Fisher Scientific, Inc.) was used to quantify the protein concentration. Protein samples (30 µg/lane) were then loaded onto 12% SDS-PAGE gels, and proteins were then transferred onto polyvinylidene difluoride (PVDF) membranes. Subsequently, protein was blocked with 8% skim milk (Pierce; Thermo Fisher Scientific, Inc.) in 0.1% Tris buffered saline with Tween-20 (TBST; OriGene Technologies, Inc.) for 2 h at room temperature. After washing with TBST three times, the membranes were incubated with primary antibodies against NRF2 (cat. no. 12282) and GAPDH (cat. no. 5174), all of which were purchased from Cell Signaling Technology, Inc., and CHAC1 (cat. no. MA5-2631; Thermo Fisher Scientific, Inc.), overnight at 4˚C. Primary antibodies were diluted with antibody diluent reagent solution (1:5,000 dilution; cat. no. ZB-2301; OriGene Technologies, Inc.) at room temperature, for 1 h. Immobilon Western Chemilum HRP (cat. no. WBKLS0500; MilliporeSigma) was used to visualize the antibody-antigen interactions. ImageJ software version 1.8.0 (National Institutes of Health) was also applied for densitometric analysis.

Human cell line xenografts. Athymic nu/nu female mice aged 6-8 weeks (n=9; mean weight, 20.2±1.54 g) were purchased from the specific pathogen SPF (Beijing) Lab Animals Technology Co. Ltd. Mice were housed in a temperature- and humidity-controlled environment (20-24˚C, 45-55% humidity), with free access to food and water and in groups of three. All procedures were reviewed and approved by the Institutional Animal Care and Use Committee (IACUC ID: 17-3256) at Nantong University and performed in accordance with the NIH Guide for the Care and Use of Laboratory Animals. Briefly, the mice were anesthetized with isoflurane (Sigma-Aldrich; Merck KGaA) inhalation at a concentration of 2.5% for anesthetic induction and then at 1.5% for anesthetic maintenance (22-24). SKOV3 cells (10⁶ cells per mouse in 100 µl PBS) were implanted subcutaneously into the right flanks of 6- to 8-week-old female nu/nu mice. Animal health and behavior were monitored each day. The mice were randomly divided three treatment groups (n=3 mice in each group, n=9 in total) as follows: i) The vehicle control [4% DMSO with 30% polyethylene glycol (PEG) 300 and double distilled H₂O]; ii) NCTD (100 mg/kg/day); and iii) NCTD (200 mg/kg/day) for 4 weeks. Tumor weight and tumor volume were determined at the end of a 4-week period. To ensure the mice were fully anesthetized, regular and even breathing was looked for. All mice were euthanized by decapitation under deep isoflurane anesthesia (5%) (25). The successful induction of anesthesia was confirmed by observation of the following parameters: respiration decreased in frequency and increased in depth, eyelid and cornea reflexes disappeared, muscle tension and the reflex response reduced, and no response to pain or other stimulation. Death was confirmed by a cessation of breathing.

Table I. Sequences of primers used in the present study.

| Primer name | Sequence (5'-3') |
|-------------|-----------------|
| CHAC1-Fw    | AGCAGATATGGTGTTGTC |
| CHAC1-Rv    | GGAATTCGAGTCTCATGGA |
| PTGS2-Fw    | GAGGGATCTGTGATGCTTCG |
| PTGS2-Rv    | AAACCACTGTTGCTACG |
| NRF2-Fw     | TGGCCCTGGAAAGTGTCAAC |
| NRF2-Rv     | CCCCCGGATGGACAAAGG |
| HO-1-Fw     | AGGGAATTCTCTTGCTGGC |
| HO-1-Rv     | GTGCTCACATTAGGTGCT |
| xCT-Fw      | TCTCCCTATGGCCAAACAGG |
| xCT-Rv      | TCCCCACTGCGTAAATGGAC |
| GPX4-Fw     | GTTTTCGCCCAAGGACATC |
| GPX4-Rv     | TGAGGAACTGTTGGAGAC |
| GAPDH-Fw    | TTCACAGGCAGACCCACACT |
| GAPDH-Rv    | CTGGTGGTCCAGGGTCTTA |

Fw, forward; Rw, reverse; PTGS2, prostaglandin-endoperoxide synthase 2; CHAC1, glutathione-specific γ-glutamylcyclotransferase 1; NRF2, nuclear factor erythroid 2-related 2; HO-1, heme oxygenase 1; xCT, solute carrier family 7 member 11; GPX4, glutathione peroxidase 4.
Tumor volumes were calculated using the following formula: \( a^2 \times b \times 0.4 \), where ‘a’ corresponds to the smallest diameter and ‘b’ to the diameter perpendicular to ‘a’. GSH, ROS, Fe\(^{2+}\) and MDA levels were also assessed in the tumor tissues with the aforementioned kits.

**Statistical analysis.** Statistical analysis was performed with SPSS version 13.0 (SPSS Inc.), and all data were quantitative. Statistical analysis was performed using an unpaired Student's t-test for comparisons between two groups and one-way analysis of variance (ANOVA) followed by Tukey's post hoc test was used for comparisons of more than two groups. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**NCTD decreases OC cell viability in a concentration- and time-dependent manner.** Firstly, the SKOV3 and OVCAR-3 cells were treated with NCTD at concentrations of 0.25, 0.5, 1, 2, 4, 8, 16, 32, and 64 \( \mu \)g/ml for 24 h. NCTD treatment decreased (A) SKOV3 and (B) OVCAR-3 cell viability in a concentration-dependent manner. SKOV3 and OVCAR-3 cells were treated with 10 and 20 \( \mu \)g/ml NCTD for 0, 12, 24, 48, and 72 h. The results of CCK-8 assay demonstrated that NCTD reduced (C) SKOV3 and (D) OVCAR-3 cell viability in a time-dependent manner. One-way ANOVA followed by Tukey's post hoc test was used for statistical analysis. *P<0.05 and **P<0.01 and, ***P<0.001 vs. control (no treatment). NCTD, norcantharidin.

![Figure 1. NCTD decreases ovarian cancer cell viability in a concentration- and time-dependent manner.](image)

Figure 1. NCTD decreases ovarian cancer cell viability in a concentration- and time-dependent manner. SKOV3 and OVCAR-3 cells were treated with NCTD at 0.25, 0.5, 1, 2, 4, 8, 16, 32 and 64 \( \mu \)g/ml for 24 h. NCTD treatment decreased (A) SKOV3 and (B) OVCAR-3 cell viability in a concentration-dependent manner. SKOV3 and OVCAR-3 cells were treated with 10 and 20 \( \mu \)g/ml NCTD for 0, 12, 24, 48 and 72 h. The results of CCK-8 assay demonstrated that NCTD reduced (C) SKOV3 and (D) OVCAR-3 cell viability in a time-dependent manner. One-way ANOVA followed by Tukey's post hoc test was used for statistical analysis. *P<0.05 and **P<0.01 and, ***P<0.001 vs. control (no treatment). NCTD, norcantharidin.

**NCTD induces ferroptosis in OC cells.** SKOV3 and OVCAR-3 cells were pre-incubated with different inhibitors, including Z-VAD-FMK (an apoptosis inhibitor), Fer-1 (a ferroptosis inhibitor), Nec-1 (a necrosis inhibitor) and 3-MA (an autophagy inhibitor), and then treated with NCTD. The results of CCK-8 assay revealed that treatment with NCTD significantly decreased SKOV3 and OVCAR-3 cell viability (Fig. 2A and B). In comparison, pre-incubation with Fer-1 significantly reversed the NCTD-induced reduction in SKOV3 and OVCAR-3 cell viability. However, no changes in cell viability were observed when the SKOV3 and OVCAR-3 cells were treated with NCTD in combination with Z-VAD-FMK, Nec-1 and 3-MA (Fig. 2A and B). Additionally, DCFH-DA staining indicated that H_2O_2 increased ROS accumulation in the SKOV3 and OVCAR-3 cells, with the pre-incubation with Fer-1 abolishing these effects (Fig. 2C and D). Flow cytometric assays also demonstrated a significant increase in cell death following treatment of the SKOV3 and OVCAR-3 cells with NCTD. However, cell death significantly decreased when the SKOV3 and OVCAR-3 cells were pre-incubated with Fer-1 (Fig. 2E and F). These observations indicated that NCTD mainly contributed to OC cell death by inducing ferroptosis.

**NCTD increases intracellular MDA and Fe\(^{2+}\) levels.** The intracellular MDA and Fe\(^{2+}\) contents were then quantified. The
results revealed that NCTD significantly increased the intracellular MDA and Fe$^{2+}$ levels in the SKOV3 and OVCAR-3 cells (Fig. 3A-D). However, following pre-incubation with Fer-1, the MDA and Fe$^{2+}$ levels were significantly decreased in the SKOV3 and OVCAR-3 cells (Fig. 3A-D). PTGS2 and CHAC1 ferroptosis marker mRNA levels were also quantified. The results of RT-qPCR analysis indicated that NCTD significantly increased the PTGS2 and CHAC1 mRNA levels in SKOV3 and OVCAR-3 cells. By contrast, pre-incubation with Fer-1 significantly reversed these effects (Fig. 3E-H).
Figure 3. NCTD increases the intracellular Fe$^{2+}$ content in SKOV3 and OVCAR-3 cells. SKOV3 and OVCAR-3 cells were pre-incubated with 1 µM Fer-1 for 1 h. The SKOV3 and OVCAR-3 cells were then treated with 10 and 20 µM NCTD for an additional 24-h time period. Pre-incubation with Fer-1 reduced the NCTD-induced upregulation of MDA contents in (A) SKOV3 and (B) OVCAR-3 cells. Fer-1 decreased the NCTD-induced elevation in the Fe$^{2+}$ contents in (C) SKOV3 and (D) OVCAR-3 cells. Reverse transcription-quantitative PCR analysis demonstrated that the PTGS2 and CHAC1 mRNA levels were decreased in the (E and G) SKOV3 and (F and H) OVCAR-3 cells, pre-incubated with Fer-1. Western blot analysis indicated that the expression of PTGS2 and CHAC1 was enhanced in (I) SKOV3 and (J) OVCAR-3 cells treated with NCTD, compared with that of the control. One-way ANOVA followed by Tukey’s post hoc test was used for statistical analysis. *P<0.05, **P<0.01, ***P<0.001 vs. control (DMSO); #P<0.05, ##P<0.01, ###P<0.001 vs. NCTD. NCTD, norcantharidin; Fer-1, ferrostatin-1; MDA, malonaldehyde; Fe$^{2+}$, ferrous ion; PTGS2, prostaglandin-endoperoxide synthase 2; CHAC1, glutathione-specific $\gamma$-glutamylcyclotransferase 1.
Western blot analysis also revealed that PTGS2 and CHAC1 expression levels were increased in the SKOV3 and OVCAR-3 cells treated with NCTD, compared with those of the untreated control, with the application of Fer-1 eliminating these effects (Fig. 3I and J).

**Figure 4.** NCTD suppresses NRF2/HO-1 signaling in ovarian cancer cells. SKOV3 and OVCAR-3 cells were treated with 10 and 20 µM NCTD for 24 h. Reverse transcription-quantitative PCR analysis indicated that the mRNA levels of NRF2, HO-1, xCT and GPX4 were significantly suppressed in the (A) SKOV3 and (B) OVCAR-3 cells treated with NCTD. (C) Western blot analysis demonstrated that the NRF2, HO-1, xCT and GPX4 protein levels were significantly suppressed in SKOV3 and OVCAR-3 cells incubated with NCTD. An unpaired Student's t-test was used for statistical analysis. *P<0.05, **P<0.01 and ***P<0.001 vs. control (DMSO). NCTD, norcantharidin; NRF2, nuclear factor erythroid 2-related 2; HO-1, heme oxygenase 1; xCT, solute carrier family 7 member 11; GPX4, glutathione peroxidase 4.

### Western blot analysis

Western blot analysis also revealed that PTGS2 and CHAC1 expression levels were increased in the SKOV3 and OVCAR-3 cells treated with NCTD, compared with those of the untreated control, with the application of Fer-1 eliminating these effects (Fig. 3I and J).

**NCTD suppresses NRF2/HO-1 signaling in OC cells.** Previous studies have indicated that NRF2 may be critical for ferroptosis-related cancer cell death (26,27). Hence, the effects of NCTD on NRF2 activation in SKOV3 and OVCAR-3 cells were examined in the present study. The results of RT-qPCR revealed that the NRF2, HO-1, xCT and GPX4 mRNA levels were significantly decreased in the SKOV3 and OVCAR-3 cells treated with NCTD (Fig. 4A and B). Moreover, western blot analysis demonstrated that the NRF2, HO-1, xCT and GPX4 protein levels were significantly suppressed in the SKOV3 and OVCAR-3 cells incubated with NCTD (Fig. 4C).

**NCTD suppresses tumor growth in vivo.** The *in vivo* experimental results demonstrated that treatment with 100 and 200 mg/kg NCTD for 4 weeks significantly suppressed tumor volume and weight (Fig. 5A-C). Treatment with 100 and 200 mg/kg NCTD also significantly decreased the GSH content in the tumors (Fig. 5D). In comparison, the MDA, ROS and Fe²⁺ contents were significantly increased in the tumor tissues of SKOV3-injected nu/nu female mice following treatment with 100 and 200 mg/kg NCTD treatment for 4 weeks (Fig. 5E-G). Western blot analysis also demonstrated that NCTD decreased the NRF2, HO-1, xCT and GPX4 protein levels, whereas it increased the PTGS2 and CHAC1 expression levels in SKOV3 tumor xenografts (Fig. 5H).

### Discussion

OC is the leading cause of gynecological malignancy-related mortality worldwide, and >75% of affected women are diagnosed at advanced stages of OC, presenting with indistinct and non-specific symptoms (28). Following diagnosis, the 5-year survival rate of patients with late-stage disease is limited to less than one-third (29). Metastatic disease following surgery and intensive platinum-taxane chemotherapy is suggested to be the major cause of mortality (29).
Figure 5. NCTD suppresses tumor growth in vivo. (A) Representative tumor images. Treatment with 100 and 200 mg/kg NCTD for 4 weeks significantly suppressed (B) tumor volume and (C) weight. (D) Treatment with 100 and 200 mg/kg NCTD significantly increased the GSH content in the tumors. The (E) MDA, (F) ROS and (G) Fe\(^{2+}\) contents were significantly decreased following treatment with 100 and 200 mg/kg NCTD for 4 weeks. (H) The results of western blot analysis demonstrated that NCTD reduced the NRF2, HO-1, xCT and GPX4 protein levels, elevating the expression of PTGS2 and CHAC1 in SKOV3-derived xenografts. One-way ANOVA followed by Tukey's post hoc test was used for statistical analysis. *P<0.05, **P<0.01 and ***P<0.001 vs. control.

Con, control; NCTD, norcantharidin; ROS, reactive oxygen species; MDA, malondialdehyde; Fe\(^{2+}\), ferrous ion; NRF2, nuclear factor erythroid 2-related 2; CHAC1, glutathione-specific γ-glutamylcyclotransferase 1; HO-1, heme oxygenase 1; xCT, solute carrier family 7 member 11; GPX4, glutathione peroxidase 4.
Previous studies have confirmed that ferroptosis-induced cell death plays a crucial role in the development of OC (30-32). NCTD prevents tumorigenesis by suppressing cell proliferation, and inducing apoptosis and cell cycle arrest in various tumors, including hepatocellular carcinoma, breast cancer and urinary bladder carcinoma (33-35). However, whether NCTD induces ferroptosis in OC has not been explored to date, at least to the best of our knowledge. In the present study, the results of CCK-8 assay indicated that NCTD suppressed OC cell viability in a time- and concentration-dependent manner. In the SKOV3 cells, a slight increase in cell viability was observed at 1 µg/ml. Hence, 0.25 and 0.5 µg/ml NCTD were used for a detailed exploration. However, in the OVCAR-3 cells, a stable downward trend of cell viability was noted following treatment with 1 µg/ml NCTD. In a previous study, a slight increase in cell viability following NCTD treatment was also observed in human MG63 osteosarcoma cells (36).

In line with this finding, it was revealed that at a concentration of 0.25 and 0.5 µg/ml, the SKOV3 cells exhibited enhanced proliferation, statistically significant as compared with the control group. It is suggested that this may be attributed to not having reached the dose-effect association in SKOV3 cells. Evidently, at low concentrations (0.25 and 0.5 µg/ml), NCTD did not inhibit the proliferation of SKOV3 cells. Apart from the dose-effect association, it was hypothesized that the SKOV3 cells may activate a stress reaction in response to low concentration of NCTD treatment, thereby promoting cell proliferation. However, this phenomenon was observed in the OVCAR-3 cells. These data indicated that OVCAR-3 cells were more sensitive to NCTD treatment.

Multiple mechanisms underlying ferroptotic cell death have been reported (37,38). In non-small cell lung cancer, NCTC has been demonstrated to trigger apoptotic cell death, by activating mitophagy-mediated autophagy signaling (39). In prostate cancer cells, NCTD has been revealed to induce endoplasmic reticulum stress-mediated apoptosis via suppressing SIRT1 (40). However, NCTD has not been previously reported to induce ferroptosis in other cell types. To explore which form of cell death was induced by NCTD, in the present study, NCTD was combined with different cell death inhibitors, including z-VAD, Nec-1, 3-MA and Fer-1. The results of the present study revealed that only Fer-1, a ferroptosis inhibitor, abolished NCTD-induced cell death in SKOV3 and OVCAR-3 cells, whereas the other inhibitors did not. Furthermore, NCTD significantly increased ROS, MDA and Fe²⁺ production. However, pre-incubation with Fer-1 abolished these effects. Flow cytometry also confirmed that NCTD increased SKOV3 and OVCAR-3 cell death. In contrast to other types of cell death, including apoptosis and necrosis, it is suggested that NCTD may have activated ferroptosis in OC cells due to the characteristics of iron dependence and accumulation of lipid ROS.

Ferroptosis is a new form of regulated cell death in the intracellular microenvironment that is induced by redox state disorder regulated via NRF2, a key regulator in iron chelators and lipophilic antioxidants (11,12). NRF2 contributes to ferroptosis by regulating its downstream target genes, including HO-1, xCT and GPX4 (12,13). One of the roles of GPX4 is to remove lipid ROS production, and GPX4 suppression results in the accumulation of lipid ROS, thereby inducing ferroptosis in HEK293T cells and rat intestinal epithelial IEC-6 cells (41,42). xCT is composed of a cystine/glutamate transporter, and it mainly acts to provide a substrate for GSH synthesis (43). xCT inhibition decreases the capacity of GPX4 to clear lipid ROS via inadequate glutathione synthesis and ultimately induces cell death (43,44). The present study demonstrated that NCTD suppressed the expression of NRF2, HO-1, GPX4 and xCT, suggesting that NCTD-induced ferroptotic cell death may be achieved by inhibiting the NRF2/HO-1/GPX4/xCT axis.

The role of NCTD in ovarian tumor growth was also explored in vivo. It was demonstrated that NCTD significantly suppressed tumor volume and weight. The GSH contents were significantly increased following NCTD treatment, whereas the ROS, MDA and Fe²⁺ levels were significantly reduced. Consistent with in vitro findings, the expression of NRF2, HO-1, GPX4 and xCT was significantly suppressed in nude mice treated with NCTD compared with that in control mice.

However, there are limitations to the present study. For instance, NRF2 overexpression or knockdown experiments are required to verify its involvement in NCTD-induced ferroptosis. Furthermore, concerning animal experiments, a control drug should have been used as a positive control. For instance, positive drugs, including erastin, should be used, which is a well-known ferroptosis activator and has been reported to inhibit OC cell growth via inducing ferroptosis in vitro and in vivo (31,45,46).

In conclusion, NCTD may represent a potent anticancer agent in OC cells and may induce OC cell death by blocking NRF2-related ferroptosis.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author upon reasonable request.

Authors' contributions

XZ performed the experiments, analyzed the data and wrote the manuscript. XC, LQ and JZ performed a part of the RT-qPCR experiments. JW designed the experiments, analyzed the data and gave final approval of the version to be published. All authors read and approved the final manuscript. XZ and JW confirm the authenticity of all the raw data.

Ethics approval and consent to participate

All procedures were reviewed and approved by the Institutional Animal Care and Use Committee (IACUC ID: 17-3256) at Nantong University and performed in accordance with the NIH Guide for the Care and Use of Laboratory Animals.
Competing interests
The authors declare that they have no competing interests.

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