

# Protective effect and mechanism of rat recombinant S100 calcium-binding protein A4 on oxidative stress injury of rat vascular endothelial cells

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**Abstract.** The aim of the present study was to examine the protective effects and mechanisms of S100 calcium-binding protein A4 (S100A4) on endothelial cell apoptosis induced by oxidative stress injury. Endothelial cells were cultured and divided into control and oxidative stress injury groups, with the latter state induced by H<sub>2</sub>O<sub>2</sub>. Endothelial cells in every group were incubated with or without 50 or 100 μM S100A4. The cell viability and amounts of malondialdehyde, nitric oxide and lactate dehydrogenase in the culture medium were measured. The apoptotic index was detected by TUNEL staining. Western blot and immunoprecipitation analyses were used to detect the expression levels and the association between S100A4 and P53. H<sub>2</sub>O<sub>2</sub> treatment led to oxidative stress injury in the cultured vascular endothelial cells, a decrease in the cell viability and an increase in the rate of apoptosis of vascular endothelial cells compared with the negative control group. Exogenous S100A4 serves a significant function against oxidative stress injury (P<0.05), increasing the viability and attenuating the apoptotic rate of endothelial cells. Western blotting results suggested that the protein levels of S100A4 and P53 increased subsequent to oxidative stress injury and that exogenous S100A4 increased the expression of P53 in the

cytoplasm and decreased the expression of P53 in nucleus. The immunoprecipitation assay results revealed a protein-protein interaction between S100A4 and P53. These results suggested that rat recombinant S100A4 serves an anti-apoptotic function in oxidative stress injury. This effect of S100A4 is mediated, at least in part, via the inhibition of the translocation of P53 to the nucleus.

## Introduction

In 2012 the World Health Organization (WHO) estimated that cardiovascular disease accounted for 30% of the total number of mortalities per year and is the leading cause of mortality worldwide (1). WHO predicts that cardiovascular mortality will rapidly increase up to 2030 (1,2). Vascular endothelial cells possess numerous biological functions, including the regulation of vascular tone, secretion of vasoactive substances and anti-platelet aggregation, serving an important role in maintaining the function of the cardiovascular system. A previous study demonstrated that the incidence of vascular disease in patients with impaired endothelial function was significantly increased compared with that in patients with normal endothelial function (3). Endothelial cell injury induced by oxidative stress is an important cause of vascular endothelial dysfunction and serves a crucial function in the occurrence of cardiovascular diseases (4). Vascular endothelial cell apoptosis is the primary form of oxidative stress injury and a prelude to the occurrence and evolution of multiple types of cardiovascular diseases (5). Wild type P53 is bound to four specific sites in the form of tetramer, which is associated with cell stress response, cell cycle and cell apoptosis (6).

The S100 protein, whose molecular weight is 11-12 kDa, belongs to the family of calcium binding proteins. It is highly homologous with calmodulin (7,8). The S100 protein family serves an important function in cell proliferation, differentiation, muscle contraction, gene expression and cell apoptosis (9,10). Dysfunction of S100 protein may cause

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different diseases. S100 calcium-binding protein A4 (S100A4) is a member of the S100 family and is associated with the occurrence, development and prognosis of tumors (11). Studies have indicated that the S100A4 protein not only is expressed in the cardiovascular endothelium, but also functions in promoting angiogenesis (12,13).

Previous studies have indicated that S100A4 binds the C terminal region of the P53 protein, which affects the subcellular localization and transcriptional activity of P53 (14). This combination affects apoptosis. The aim of the present study was to investigate whether the S100A4 protein exhibits a protective effect on oxidative stress injury of endothelial cells and to elucidate its mechanism. By establishing an oxidative stress damage model in vascular endothelial cells, the effect of S100A4 protein on oxidative stress injury of vascular endothelial cells was observed. The effect of S100A4 protein on cardiovascular endothelial cell apoptosis through the P53 pathway was also examined. Based on these experiments, the present study preliminarily clarified the regulatory mechanism of S100A4 protein on the apoptosis of vascular endothelial cells.

## Materials and methods

**Antibodies and reagents.** The following antibodies were used: P53 (cat. no. 10442-1-AP; ProteinTech Group, Inc. (Chicago, IL, USA), Caspase 3 (cat. no. ab13847; Abcam, Cambridge, MA, USA), p17-specific antibody (cat. no. 25546-1-AP; Proteintech Group, Inc.), S100A4 (ab41532; Abcam) and  $\beta$ -actin (cat. no. 60008-1-Ig; Proteintech Group, Inc.). S100A4 (cat. no. ab41532) was purchased from Abcam. The following reagents were used: Rat Lactic Dehydrogenase (LDH; cat. no. E-0672; Meilian Shengwu, Shanghai, China), nitrous oxide (NO; cat. no. KGE001; R&D Systems, Inc. Minneapolis, MN, USA), Malondialdehyde (MDA; cat. no. ab118970; Abcam), superoxide dismutase (SOD; cat. no. DYC3419-2; R&D Systems). Rat recombinant S100A4 protein was expressed and purified in the Performance Medicine Laboratory of Tianjin Institute of Health and Environmental Medicine (Tianjin, China). Cell Death Detection kit (cat. no. G3250, Promega) was from Promega Corporation (Madison, WI, USA). Hydrogels incubated with PBS alone were used as negative controls. All the experiments were repeated at least three times. The microscopy (OLMPUS CK40, Olympus corporation, Tokyo, Japan) was preserved in cell culture room and utilized to observe endothelial cells without fixing embedding and staining at 25°C. The magnification used was x100.

**Cell culture and treatment.** Rat aortic endothelial cells (RAECs) were isolated and cultured as described previously, with minor modifications (15). The thoracic aorta were excised from 3 male Wistar rats (150-180 g) after they were sacrificed humanely by injecting with 40 mg anesthetic (chloral hydrate) per 100 g of body weight and immediately placed in cold PBS containing 100 U/ml penicillin and 100 mg/ml streptomycin. The aorta was cut into 1 mm-wide rings subsequent to the removal of the periaortic fat. Following transfer to a T-25 cm<sup>2</sup> flasks (NalgeNunc International, Penfield, NY, USA), the rings were cultured in Medium 199 (cat. no. 11150059, Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA)

containing 20% fetal bovine serum (FBS; cat. no. 10099141, Gibco; Thermo Fisher Scientific, Inc.), 2.5 ng/ml basic fibroblast growth factors, 100 U/ml penicillin and 100 mg/ml streptomycin. The aorta rings were placed at 37°C in a humidified atmosphere with 5% CO<sub>2</sub> for 72-80 h without movement. All pieces of aorta rings were removed when cells migrated. In experiments, M199 medium supplemented with 1% bovine serum albumin (cat. no. B2064; Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) was used. All experiments were performed with RAECs up to 4 passages. Male Wistar rats were gained from the Experimental Animal Room of the Institute of Health and Environmental Medicine (Tianjin, China). The rats were maintained under standard conditions (ambient temperature 21-23°C; with a 12 h dark-light cycle) with ad libitum access to food and tap water. This study was approved by the committee of the Institute of Health and Environmental Medicine, Tianjin, China.

The cells were divided into the following groups: i) A normal control group incubated in Medium 199 (NC); ii) a H<sub>2</sub>O<sub>2</sub> injury group incubated in Medium 199 containing 100  $\mu$ M H<sub>2</sub>O<sub>2</sub> (HI); and two S100A4-treated groups incubated with iii) 50  $\mu$ M or iv) 100  $\mu$ M S100A4 in Medium 199 containing 100  $\mu$ M H<sub>2</sub>O<sub>2</sub> [(HI+S100A4 (50  $\mu$ M)/HI+S100A4 (100  $\mu$ M)]. In the MTT and ELISA analysis, two groups were added to detect whether S100A4 had damaged the cells. They were NC+S100A4 (50  $\mu$ M) and NC+S100A4 (100  $\mu$ M) groups.

**Cell proliferation assay.** Cell proliferation was determined using an MTT assay. A total of  $\sim 3 \times 10^3$  endothelial cells were plated into each well of a 96-well plate. Following overnight incubation at 37°C, the cells were treated with 100  $\mu$ M H<sub>2</sub>O<sub>2</sub>, H<sub>2</sub>O<sub>2</sub>+S100A4 (50/100  $\mu$ M) for 48 h. Then the medium was removed and MTT (20  $\mu$ l of 5 mg/ml) was added to each well and incubated at 37°C for 4 h. Plates were agitated at low speed (16.77 x g) 37°C for 10 min and the purple-colored precipitates of formazan were dissolved in 150  $\mu$ l dimethyl sulfoxide. Absorbance was measured at 490 nm using an ELISA plate reader. The reduction in viability of in H<sub>2</sub>O<sub>2</sub>-treated endothelial cells was expressed as a percentage compared with non-H<sub>2</sub>O<sub>2</sub> treated control cells. Control cells were considered 100% viable.

**ELISA assay.** The levels of LDH (Rat LDH ELISA kit; cat. no. E-0672; Meilian Shengwu), NO (Total Nitric Oxide and Nitrate/Nitrite Parameter assay kit; cat. no. KGE001; R&D Systems), MDA (MDA assay kit; cat. no. ab118970; Abcam), and SOD (Human/Mouse/Rat Total SOD2/Mn-SOD DuoSet IC ELISA kit; cat. no. DYC3419-2; R&D Systems) from endothelial cell medium were quantified by ELISA assay according to the manufacturer's protocol.

**Assessment of apoptosis by TUNEL staining.** Cells apoptosis was determined by double-labeling TUNEL immunofluorescence staining. A total of  $2 \times 10^4$  RAECs with Medium 199 (cat. no. 11150059; Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% FBS were seeded into each well of a 48-well plate. Following the aforementioned experimental treatments, cells were fixed in 4% paraformaldehyde for 30 min at room temperature. Subsequently, the cells were incubated with 3% H<sub>2</sub>O<sub>2</sub> in methanol for 10 min at room temperature to block endogenous peroxidase activity, and were then incubated

with 0.1% Triton X-100 in 0.1% sodium citrate for 2 min at 4°C. After washing in PBS, the cells were incubated with the TUNEL reaction mixture containing 5  $\mu$ l enzyme solution and 45  $\mu$ l fluorochrome-labeled solution at 37°C for 60 min in the dark. For counterstaining, the cells were incubated with DAPI for 15 min at room temperature. The cells were observed using a fluorescence microscope, in which 10 fields were randomly selected. The TUNEL+ and DAPI+ nuclei in the cells were counted manually. The percentage of apoptotic cells was calculated as the ratio of the number of TUNEL-positive cells to the total number of cells, which were counted using a fluorescence microscope.

**Western blot analysis.** The 4 groups of cells (described above) were harvested. The cells samples were homogenized in a Radio-Immunoprecipitation Assay (RIPA) lysis buffer [50 mM Tris (pH 7.4), 150 mM NaCl, 1% TritonX-100, 1% sodium deoxycholate, 0.1% SDS]. Protein concentrations were determined with a BCA assay kit (Pierce; Thermo Fisher Scientific, Inc.). The proteins (10  $\mu$ l in each line) were separated using a 10% gel with SDS-PAGE and transferred onto nitrocellulose membranes. Membranes were blocked with TBS-Tween-20 (0.05%) containing 5% non-fat milk for 2 h at room temperature and incubated with the following primary antibodies, anti- $\beta$ -actin (1:1,000, 60008-1-Ig, Proteintech Group, Inc.), anti-P53 (1:500, 10442-1-AP, Proteintech Group, Inc.), anti-cleaved-caspase-3 (1:1,000, 25546-1-AP, Proteintech Group, Inc.), anti-S100A4 (1:250, ab41532, Abcam) overnight at 4°C. The membranes were then incubated with secondary antibody, goat anti-rabbit antibodies (1:5,000, ZB-2301, ZSGB-Bio) conjugated with horseradish peroxidase for 1 h at room temperature (25°C). A SuperSignal™-enhanced chemiluminescent substrate (Pierce; Thermo Fisher Scientific, Inc.) was applied to the probed membrane for 3 min at room temperature. The blots were quantified via densitometry using ImageJ software (version 1.37; National Institutes of Health, Bethesda, MA, USA).

**Immunoprecipitation (IP).** Endothelial cells were seeded in 10-cm dishes, followed by stimulation (37°C) with or without 100  $\mu$ M H<sub>2</sub>O<sub>2</sub> for 48 h. Subsequently, the HepG2 cell (Tianjin Saierbio Biotechnology Co., Ltd., Tianjin, China) lysates were prepared with 1% Tris-Triton cell lysis buffer (Cell Signaling Technology, Inc., Danvers, MA, USA) pre-mixed with 1 mM phenylmethanesulfonyl fluoride on ice for 30 min and centrifuged (4°C) at 12,000 x g for 30 min. The supernatants were incubated (4°C) overnight with 30  $\mu$ l Dynabeads protein A or protein G (Thermo Fisher Scientific, Inc.) pre-coated with anti-S100A4 (1:250, Abcam) antibodies. The immunocomplexes were subjected to western blot analysis. The normal corresponding immunoglobulin G control was assayed simultaneously.

**Statistical analysis.** Data are presented as the mean  $\pm$  standard deviation. Statistical analysis was performed using a Student's t-test for paired samples or a single-factor analysis of variance with Student-Newman-Keuls post-hoc test as appropriate by SPSS software (version 20.0; IBM Corp., Armonk, NY, USA). P<0.05 was considered to indicate a statistically significant difference.

## Results

*S100A4 may inhibit the cell morphological changes induced by H<sub>2</sub>O<sub>2</sub>.* Visible under a phase contrast microscope, in the NC group the endothelial cells were polygonal and exhibited a clear structure, uniform size and single paving stone-like close spacing. The cell morphology of the NC+S100A4 groups was almost identical to that of the NC group. In the HI group, the endothelial cells exhibited shrinkage, roundness and clearance. There were vacuoles and nuclear chromatin condensation and aggregation in the cytoplasm of the cell. Cells demonstrated a disordered arrangement, losing the typical single layer paving stone-like arrangement. In the HI+S100A4 groups, cell morphology and structure were predominantly clear (Fig. 1A). The arrangement and morphology of the cells was not markedly changed compared with the NC group.

The MTT assay easily and effectively evaluates the activity of cells. The cell survival rate of the oxidative stress injury group was significantly decreased compared with that of the NC group (P<0.05), suggesting that H<sub>2</sub>O<sub>2</sub> may cause endothelial cell death. There were no significant differences in cell survival rate between the NC, NC+S100A4 (50  $\mu$ M) and NC+S100A4 (100  $\mu$ M) groups following administration of S100A4 protein, which indicated that exogenous S100A4 protein demonstrated no significant effect on endothelial cells. In addition, the cell survival rate of the 50 and 100  $\mu$ M HI+S100A4 groups was significantly increased compared with that in the HI group, but remained significantly decreased compared with that of the NC group (P<0.05) and the cell survival rate of HI+S100A4 (100  $\mu$ M) group was increased compared with that of the HI+S100A4 (50  $\mu$ M) group (Fig. 1B). This indicated that S100A4 protein may improve the survival rate of these cells.

*Effect of S100A4 protein on LDH, NO, MDA and SOD in cultured endothelial cells induced by oxidative stress.* The results indicated that the LDH activity and MDA content of HI group were significantly increased compared with those of the NC group (P<0.01), while NO content and SOD activity were significantly decreased compared with those of the NC group (P<0.01), suggesting that H<sub>2</sub>O<sub>2</sub> oxidatively damaged these cells. However, the LDH values of the 50 and 100  $\mu$ M HI+S100A4 groups were significantly decreased compared with that of the HI group, but remained significantly increased compared with that of the NC group (P<0.05). The LDH value of the HI+S100A4 (100  $\mu$ M) group was decreased compared with that of the HI+S100A4 (50  $\mu$ M) group (P<0.05; Fig. 2A). The NO content of 50 and 100  $\mu$ M HI+S100A4 groups was significantly increased compared with that of the HI group, but remained significantly decreased compared with that of the NC group (P<0.05; Fig. 2B). The content of NO in the HI+S100A4 (100  $\mu$ M) group was increased compared with that in the HI+S100A4 (50  $\mu$ M) group (P<0.05; Fig. 2B). The MDA content results demonstrated that different doses of S100A4 protein may significantly reverse the increase in MDA levels induced by H<sub>2</sub>O<sub>2</sub> injury (P<0.05; Fig. 2C). The SOD activities in the 50 and 100  $\mu$ M HI+S100A4 groups were significantly increased, compared with the HI group (P<0.05; Fig. 2D). The experimental results indicated that S100A4 protein prevented oxidative damage.

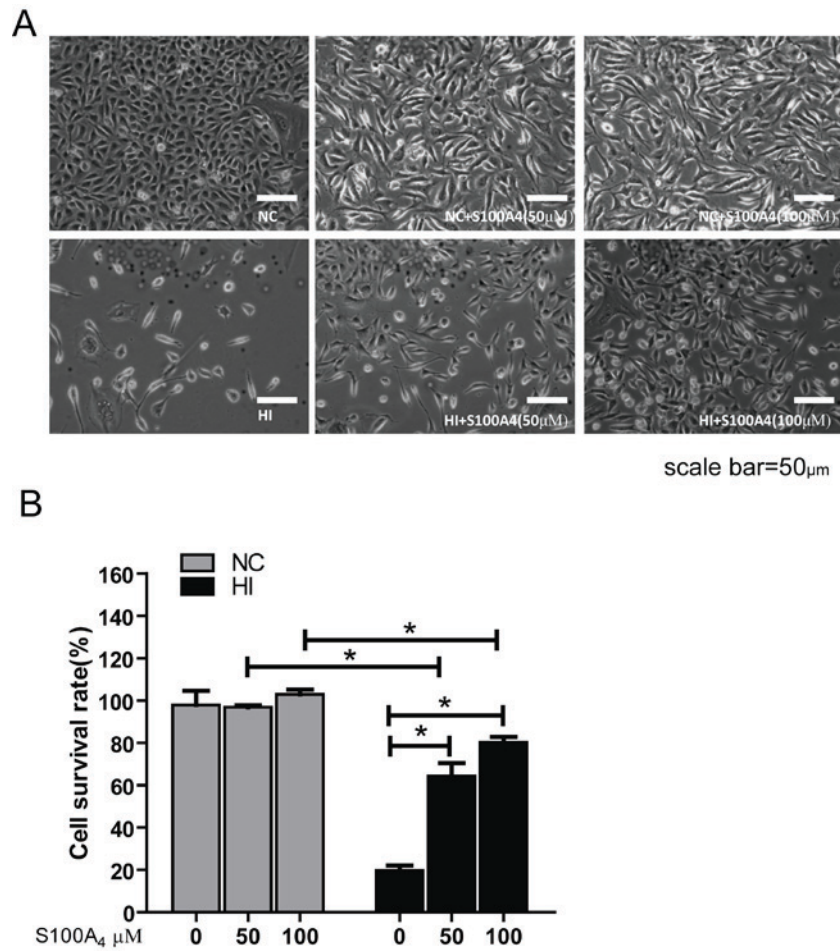


Figure 1. S100A4 suppresses apoptosis. Cultured rat vascular endothelial cells 12 h after H<sub>2</sub>O<sub>2</sub> injury with and without S100A4 treatment prior to fixing with 4% paraformaldehyde for the apoptosis assay. (A) Cell morphology was detected by light microscopy scale bar 50  $\mu$ m; (B) Proliferation of cells was determined by the MTT assay. \*P<0.05 vs. H<sub>2</sub>O<sub>2</sub> group. S100A<sub>4</sub>, S100 calcium-binding protein A4; NC, normal control; HI, hypoxic injury induced by H<sub>2</sub>O<sub>2</sub>.

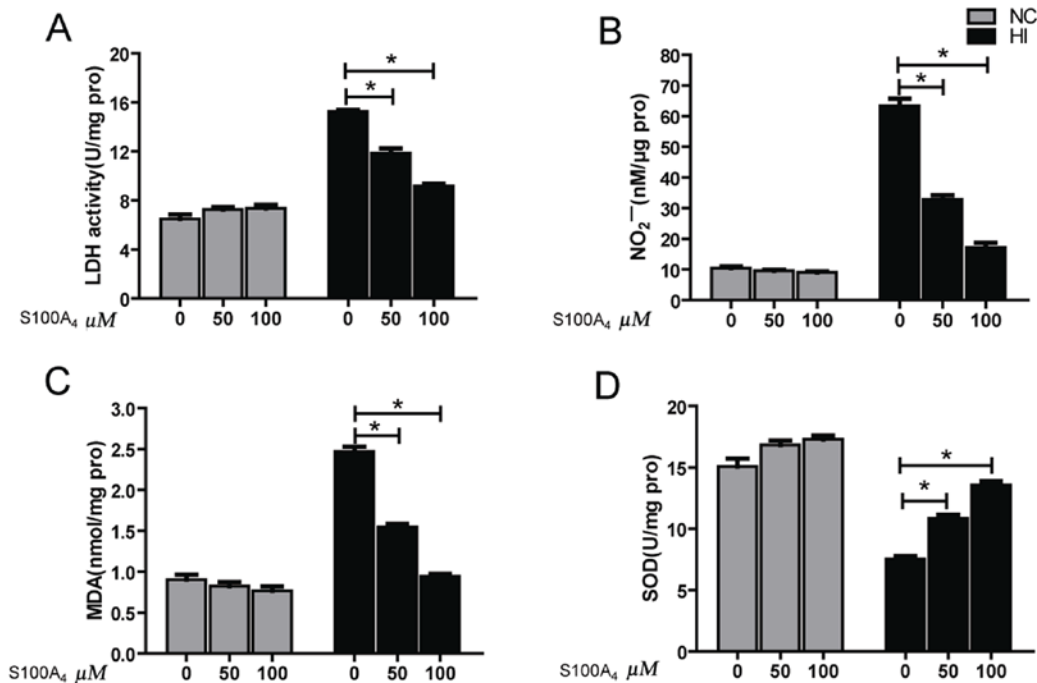


Figure 2. Effect of S100A4 protein on (A) LDH, (B) NO, (C) MDA and (D) SOD in cultured endothelial cells induced by oxidative stress. \*P<0.05 vs. the H<sub>2</sub>O<sub>2</sub> group. LDH, lactate dehydrogenase; NO, nitrous oxide; MDA, malondialdehyde; SOD, superoxide dismutase; S100A<sub>4</sub>, S100 calcium-binding protein A4; NC, normal control; HI, hypoxic injury induced by H<sub>2</sub>O<sub>2</sub>.

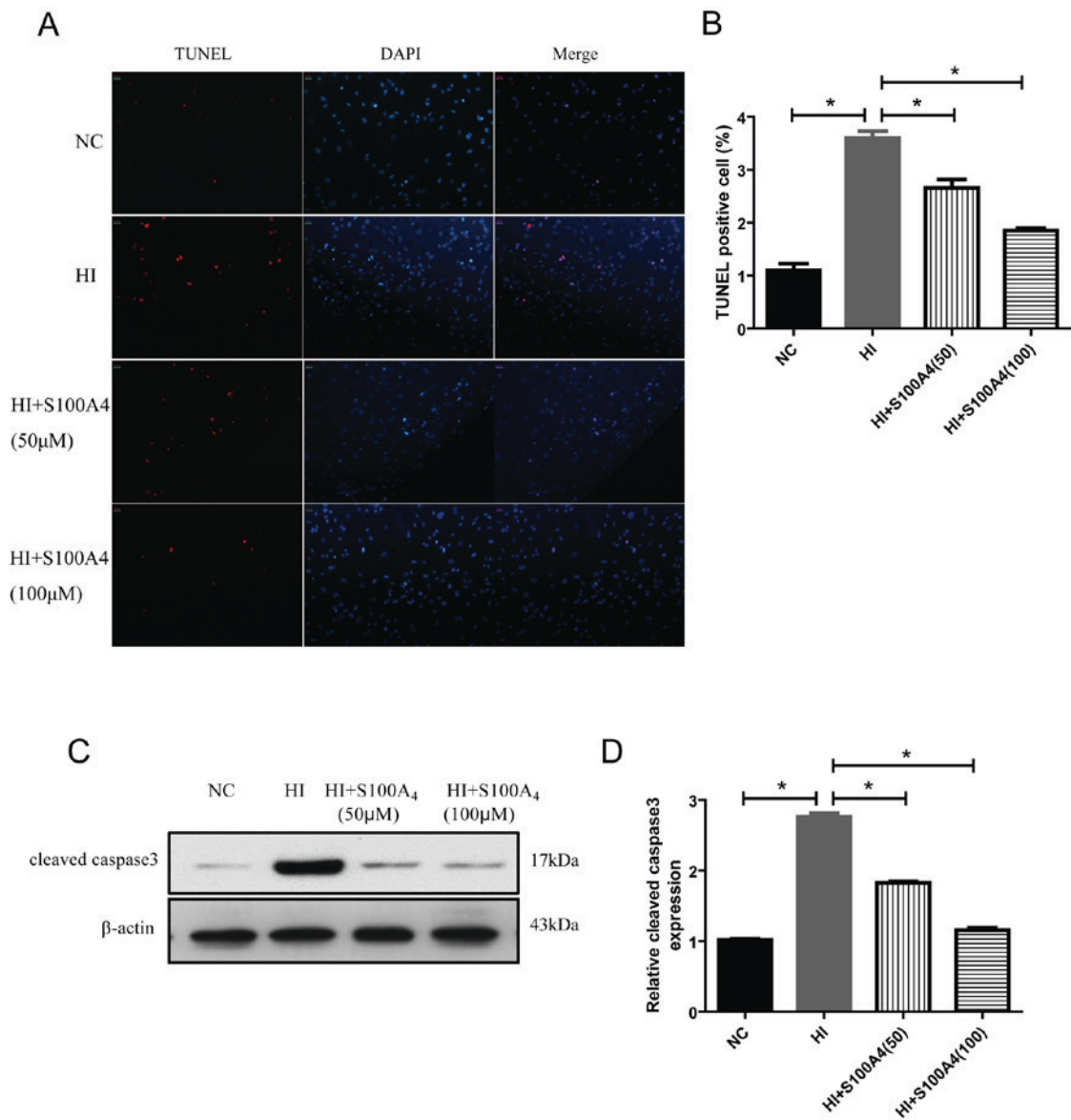


Figure 3. S100A4 causes a suppression of apoptosis induced by H<sub>2</sub>O<sub>2</sub>. (A) Images and (B) quantification of TUNEL apoptosis analysis. A total of ~4x10<sup>5</sup> rat vascular endothelial cells were cultured on the cover slips in 24-well plates and treated with 100 μM H<sub>2</sub>O<sub>2</sub>, with or without S100A4 concurrently. Following incubation, cells were washed with cold PBS (pH 7.4) and stained according to the protocol of the TUNEL assay kit. H<sub>2</sub>O<sub>2</sub> induced endothelial cell injury and cell apoptosis. The endothelial cell apoptosis rate in the HI+S100A4 (50 μM) and HI+S100A4 (100 μM) group was significantly decreased compared with that in the HI group. (C) Western blotting and (D) quantification of protein expression of caspase 3 in rat vascular endothelial cells. The expression of cleaved caspase-3 in ECs increased significantly after H<sub>2</sub>O<sub>2</sub> administration. S100A4 decreased caspase 3 expression in endothelial cells treated with 50 μM or 100 μM H<sub>2</sub>O<sub>2</sub> for 12 h. \*P<0.05 vs. H<sub>2</sub>O<sub>2</sub> group. S100A4, S100 calcium-binding protein A4; NC, normal control; HI, hypoxic injury induced by H<sub>2</sub>O<sub>2</sub>.

*Effect of S100A4 protein on the rate of endothelial cell apoptosis induced by oxidative stress.* The results demonstrated that the endothelial cell apoptosis rate of the oxidative stress injury group was significantly increased compared with that of the control group (P<0.01), suggesting that H<sub>2</sub>O<sub>2</sub> may induce endothelial cell injury and cell apoptosis. The endothelial cell apoptosis rate in the HI+S100A4 (50 μM) and HI+S100A4 (100 μM) group was significantly decreased compared with that in the HI group (P<0.05; Fig. 3A and B).

Since mitochondrial function is associated with caspase activity (16), the function of caspase 3 was assessed. The expression of cleaved (active) caspase 3 expression was demonstrated using western blot analysis. In Fig. 3, western blot analysis of cleaved caspase 3 (Fig. 3C and D) was performed following H<sub>2</sub>O<sub>2</sub> administration in ECs and revealed significant caspase

3 activity compared with NC (P<0.05). However, S100A4 significantly inhibited cleaved active caspase 3 activity. The results indicated that S100A4 protein exhibited prevented oxidative damage, inhibiting apoptosis.

*Determination of total protein, cytoplasmic protein and nuclear protein of endothelial cell protein by western blot analysis.* In Fig. 4, following S100A4 intervention, the level of intracellular S100A4 protein was significantly increased in the S100A4 groups (P<0.05). Although there was no significant change in the expression of P53 in the cell, the distribution of P53 protein in the cell was altered; the effect of S100A4 protein on P53 was dose-dependent. As the dose of S100A4 increased, the aggregation of P53 in the cytoplasm increased and the corresponding distribution in the nucleus decreased.

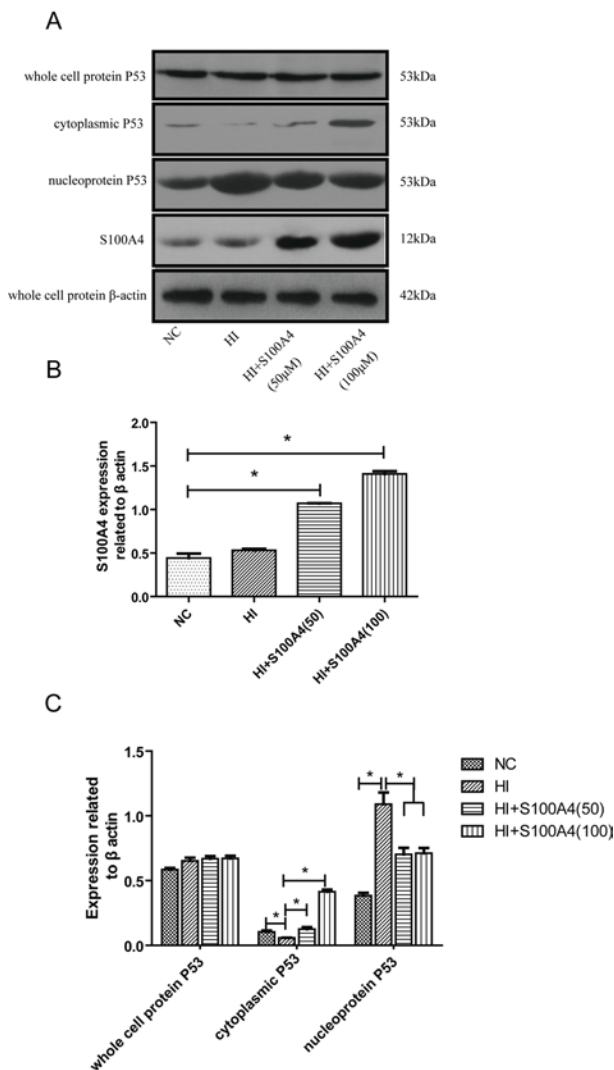


Figure 4. S100A4 functions in the cytoplasmic compartment. Western blot analysis of indicated proteins in nuclear, cytoplasmic and whole-cell extracts of endothelial cells treated with  $H_2O_2$  with or without S100A4 for 12 h. (A) Western blotting and (B) quantification of the expression of cytoplasmic S100A4 protein could not be effected by  $H_2O_2$  treatment, while it was significantly increased after exogenous S100A4 addition. (C) After  $H_2O_2$  treatment, the aggregation of P53 in the cytoplasm increased and the corresponding distribution in the nucleus increased comparison with the NC group. However, exogenous S100A4 reversed the changes mentioned above. \* $P < 0.05$  vs.  $H_2O_2$  group. NC, normal control; HI, hypoxic injury induced by  $H_2O_2$ ; S100A4, S100 calcium-binding protein A4.

This suggests that S100A4 protein may inhibit P53 movement from the cytoplasm into the nucleus. Therefore, we hypothesize that the function of S100A4 protein in endothelial cell apoptosis is inhibited in the oxidative stress damage model of endothelial cells and that this effect is mediated by P53.

**S100A4 interacts with P53 in the nucleus.** S100 family proteins have no known enzymatic activity; therefore, it may be hypothesized that S100 proteins function through interaction with other proteins to regulate the function of interacting proteins. Previous studies have focused on non-muscle myosin IIA and P53 as potential S100A4-interacting proteins. Therefore, the present study investigated the potential interaction between S100A4 and P53. IP of endogenous S100A4 in endothelial cells resulted in the co-precipitation of endogenous P53 in untreated

cells (Fig. 5). Following  $H_2O_2$  treatment, the total cell P53 did not change significantly. However, P53 in the nucleus was significantly increased. P53 binding to S100A4 was also significantly increased in the nucleus ( $P < 0.05$ ). Therefore, we hypothesized that S100A4 may combine with P53 in the nucleus to increase the degradation of P53, thereby inhibiting the apoptosis of cells.

## Discussion

The vascular endothelium is composed of a monolayer of vascular endothelial cells, which is a barrier between circulating blood and vascular smooth muscle. The vascular endothelium participates in the contraction and relaxation of blood vessels, the formation of blood vessels, thrombosis and the inflammatory response (17,18). Oxidative stress is the key factor that causes changes in endothelial function and cell damage (19,20). Oxygen free radicals, collectively known as reactive oxygen species (ROS), are one of the primary factors that cause oxidative damage of endothelial cells (18,19). Free radical reactions cause cell membrane lipid peroxidation, and then alterations to the cell transport and enzyme functions (19,20). Oxidative stress is associated with the pathological process of the endothelial cells, including increasing vascular endothelial cell permeability, affecting cell proliferation, increasing leukocyte infiltration and interfering with the signal transduction in the cell, which causes cell death and apoptosis (21). It serves a key function in the occurrence and development of vascular diseases (22). Therefore, it is important to prevent the occurrence of cardiovascular disease through the protection of vascular endothelial cells from oxidative damage.

The S100 protein family is one of the largest families of calcium-binding proteins. Including the S100A4 protein, at present, there are 21 members of the S100 family (23). Previous studies have demonstrated abnormal expression of S100 in numerous diseases, including psoriasis, Alzheimer's disease, cystic fibrosis, cardiomyopathy, muscle atrophy (spinal cord) lateral sclerosis and cancer (24,25). The S100A4 protein belongs to the S100 protein family and is composed of 101 amino acids with a molecular weight of 11.5 kD (23). Studies have confirmed that the S100A4 protein may promote the invasion and metastasis of tumors and regulate the function of the cell, including cell growth and cell signal transduction (26,27). Multiple studies demonstrate that the S100A4 protein is associated with vascular regeneration and extracellular matrix remodeling and may even be used as an independent vascular-stimulating factor (13,28). Schmidt-Hansen *et al* (29) identified that intracellular S100A4 stimulates endothelial cells to produce matrix metalloproteinases, which promotes the remodeling of the extracellular matrix and the degradation of matrix remodeling is a necessary step in angiogenesis. An additional study indicated that the release of S100A4 from the cornea of the implanted rat cornea may induce novel blood vessel formation (30). Furthermore, the interaction of S100A4 protein and membrane protein may promote the activation of plasminogen activator, which is induced by the activation of plasminogen activator (31).

To assess the function of S100A4 protein on endothelial cell damage induced by oxidative stress and the potential

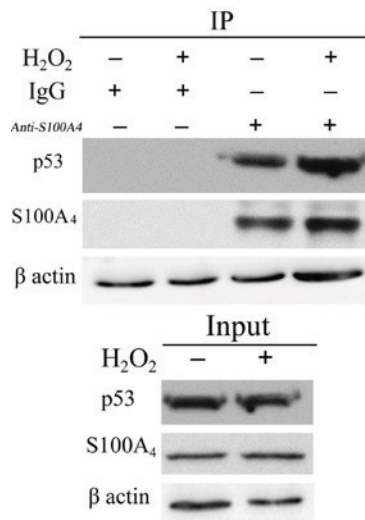


Figure 5. S100A4 interacts with P53, as indicated by IP. The whole-cell extracts of ECs were treated with H<sub>2</sub>O<sub>2</sub>. Immunoblot analysis of P53 and S100A4 protein in S100A4 IP experiments in untreated and H<sub>2</sub>O<sub>2</sub>-treated ECs. IP, immunoprecipitation; ECs, endothelial cells; S100A4, S100 calcium-binding protein A4; Ig, immunoglobulin.

mechanism, the present study selected 100  $\mu$ M H<sub>2</sub>O<sub>2</sub>-induced injury for 12 h to establish a model of endothelial cell oxidative stress injury (32). The results indicated that the oxidative stress caused the loss of the typical morphology of the endothelial cells. The cells exhibited shrinkage and roundness, the gap increased and vacuoles in the cytoplasm of cells were not arranged in order. The activity of LDH was increased, the content of NO was decreased and the content of MDA was increased, while the activity of SOD was decreased. The survival rate of endothelial cells was decreased and the apoptosis rate was increased. The experiment suggested that oxidative stress resulted in the injury of endothelial cells. S100A4 prevented oxidative stress to protect endothelial cells.

The important mechanism of oxygen free radical-induced cell and tissue damage is to induce peroxidation of the polyunsaturated fatty acids in the biological membrane, consequently producing lipid peroxide. Lipid peroxidation may be associated with the occurrence of a chain reaction of polyunsaturated fatty acids (28). As one of the unsaturated fatty acids, linoleic acid can induce both NF- $\kappa$ B and AP-1 transcriptional activation (33). Lipid free radicals and their degradation products (MDA) are formed, causing membrane fluidity, permeability and integrity damage and then the cell membrane is destroyed (34,35). ROS mainly includes oxygen ions and peroxides. As a superoxide scavenging enzyme, active SOD is able to neutralize O<sub>2</sub><sup>-</sup> and presents antioxidative effect (36). With the aid of SOD, superoxide radical anion or hyperoxide in biological tissues can be changed into hydrogen peroxide (HO) and singlet oxygen (<sup>1</sup>O<sub>2</sub>). HO may be changed into water by catalase or glutathione peroxidase (36). Therefore, the content of MDA may reflect the degree of lipid peroxidation and indirectly reflect the degree of cell damage. SOD activity indirectly reflects the ability of the body to clear oxygen free radicals. Oxidative stress may increase the secretion of endothelin-1 and diminish the bioavailability of nitric oxide (NO) (3,37). These vasoactive molecules promote the contraction of blood vessels and then initiate a series of post-injury reactions, resulting in the occurrence of cardiovascular disease (38,39).

Oxidative stress may lead to endothelial dysfunction (40,41), resulting in decreased NO levels and increased LDH release (42). The results indicated that H<sub>2</sub>O<sub>2</sub> may increase the content of MDA, the activity of LDH was increased and that the activity of SOD was decreased, suggesting that the endothelial cells were damaged by lipid peroxidation. S100A4 protein intervention may be effective against oxygen free radical damage.

Therefore, we hypothesized that the protective mechanism of S100A4 on the endothelial cells may be associated with the protection of cell mitochondria, increased cell activity.

Endothelial cell activity is key to the maintenance of endothelial function. If the apoptosis rate of endothelial cells exceeds the normal level, it will affect the integrity of the blood vessels and induce the injury of endothelial cells. The P53 gene is an important anti-cancer gene. A previous study demonstrated that wild type P53 may induce the apoptosis of leukocytes (43). Three members of the S100 protein family, namely S100B, S100A2 and S100A4, physiologically interact with P53 in a calcium dependent manner. Phosphorylation and acetylation of P53 may be affected by the interaction of S100 protein, which results in the regulation of the subcellular localization and transcriptional activity of P53 (44). Previously, studies have focused on the role of the P53 gene in cardiovascular diseases and have indicated that P53 serves an important function in the oxidative stress of blood vessels as an important type of transcription protein (45,46). The results of the present study suggested that oxidative stress causes an increased rate of endothelial cell apoptosis and P53 may be the target protein of S100A4. Subsequent to the cells being administered S100A4 protein, the total level of P53 protein was not increased but the distribution of P53 protein in the cells was changed, primarily with an increase in the aggregation of P53 in the cytoplasm and a decrease in the distribution of P53 in the nucleus. As the dose of S100A4 increased, the aggregation of P53 in the cytoplasm increased and the corresponding distribution in the nucleus decreased. This suggested that S100A4 protein may inhibit P53 movement from the cytoplasm into the nucleus.

To conclude, exogenous S100A4 protein may reverse oxidative stress damage, improve the survival rate of cells and inhibit the apoptosis of endothelial cells. The protective mechanism of S100A4 protein on the apoptosis of endothelial cells may be mediated by inhibiting P53. This provides a reference for the additional exploration of the application of S100A4 protein in endothelial cell damage induced by oxidative stress.

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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 on reasonable request.

### Authors' contributions

TW and BS developed the project. XM, XG and ZZ performed experiments and wrote the manuscript. XZ, LW and MY analyzed the data. KW and HR helped with computational analysis. All authors read and approved the final manuscript.

### Ethics approval and consent to participate

This study was approved by the committee of the Institute of Health and Environmental Medicine, Tianjin, China.

### Patient consent for publication

Not applicable.

### Competing interests

The authors declare that they have no competing interests.

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