Calcium Sensing Receptor Regulating Smooth Muscle Cells Proliferation Through Initiating Cystathionine-Gamma-Lyase/Hydrogen Sulfide Pathway in Diabetic Rat

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Key Words
Calcium sensing receptor • Smooth muscle cells • Proliferation • Hydrogen sulfide • Diabetic rat

Abstract
Aims: Hydrogen sulfide (H₂S) inhibits the proliferation of vascular smooth muscle cells (VSMCs). However, how cystathionine-gamma-lyase (CSE), a major enzyme that produces H₂S, is regulated remains unknown. Whether calcium-sensing receptor (CaSR) inhibits the proliferation of VSMCs by regulating the endogenous CSE/H₂S pathway in diabetic rat has not been previously investigated. Methods and Results: The morphological and ultrastructure alterations were tested by transmission electron microscopy, changes in the H₂S concentration and the relaxation of the mesenteric secondary artery loop of diabetic rats were determined by Multiskan spectrum microplate spectrophotometer and isometric force transducer. Additionally, the expression levels of CaSR, CSE and Cyclin D1 in the mesenteric arteries of rats were examined by western blotting. The intracellular calcium concentration, the expression of p-CaMK II (phospho-calmodulin kinases II), CSE activity, the concentration of endogenous H₂S and the proliferation of cultured VSMCs from rat thoracic aortas were measured by using confocal microscope, western blotting, microplate spectrophotometer, MTT and BrdU, respectively. The VSMC layer thickened, the H₂S concentration dropped, the relaxation of the mesenteric secondary artery rings weakened, and the expression of CaSR

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and CSE decreased whereas the expression of Cyclin D1 increased in diabetic rats compared with the control group. The [Ca$^{2+}$] of VSMCs increased upon treatment with CaSR agonists (10 μM Calindol and 2.5 mM CaCl$_2$), while it decreased upon administration of calhex231, U73122 and 2-APB. The expression of p-CaMK II and CSE increased upon treatment with CaSR agonists in VSMCs. CSE activity and the endogenous H$_2$S concentration decreased in response to high glucose, while it increased with treatment of CaSR agonists. The proliferation rate increased in response to high glucose, and CaSR agonists or NaHS significantly reversed the proliferation of VSMCs caused by high glucose. **Conclusions:** Our results demonstrated that CaSR regulated the endogenous CSE/H$_2$S pathway to inhibit the proliferation of VSMCs in both diabetic and high glucose models.

**Introduction**

Diabetes increases the risk of cardiovascular morbidity and mortality by promoting cardiovascular injury, which is a major cause of death and disability in patients [1]. During vascular injury, vascular smooth muscle cell proliferation is a critical factor and a noted characteristic. Some studies have suggested that oxidative stress [2] and growth factors [3, 4] are involved in the proliferation of VSMCs in diabetes models. Recent studies have suggested that vasodilators play a major role in modulating the proliferation of VSMCs in diabetic models [5].

Hydrogen sulfide (H$_2$S) is the third member of the gaseous transmitter family. It is a strong reducing agent [6, 7] and vasodilator [8] and is endogenously synthesized from L-cysteine. In mammalian cardiovascular tissues, the biosynthesis of H$_2$S is mainly catalyzed by cystathionine-gamma-lyase (CSE). Recent studies have shown that H$_2$S is involved in vasorelaxation [8], cardioprotection [9] and the inhibition of proliferation of VSMCs [10]. A deficiency in the CSE/H$_2$S pathway plays an important role in the development of certain cardiovascular diseases, such as spontaneous hypertension, atherosclerosis and heart failure [11-13]. However, whether H$_2$S protects the periphery vasculature against diabetes-induced injury remains unclear.

The calcium-sensing receptor (CaSR) belongs to the transmembrane G-protein coupled receptor family. Extracellular calcium binds CaSR to cause phosphatidylinositol biphosphate (PIP$_2$) to be cleaved into the secondary messengers diacylglycerol (DAG) and inositol 1,4,5-trisphosphate (IP$_3$) through the PLC (phospholipase C) pathway. IP$_3$ binds the IP$_3$ receptor on the sarcoplasmic reticulum, which causes calcium to be released and thus increases the calcium concentration in the cytoplasm [14, 15]. Some evidence suggests that a functional CaSR is expressed in VSMCs [16]. Some studies also confirmed that a positive modulator of CaSR induces the relaxation of rat isolated small mesenteric arteries [17, 18]. A recent study found that increasing the [Ca$^{2+}$]$_i$ increases endogenous H$_2$S production in smooth muscle cells [19].

The role of H$_2$S in inhibiting the proliferation of VSMCs has been investigated previously [10]. However, the effects of the CSE/H$_2$S pathway on SMC proliferation have not been characterized, and identifying an interaction between CaSR and the CSE/H$_2$S pathway may uncover the mechanism of endogenous H$_2$S production.

In the present study, we determined that the reduction of H$_2$S production in a STZ-induced diabetic rat model stimulated SMC proliferation. CaSR could influence CSE activity and H$_2$S production and arrest the progression of diabetic vascular injury.

**Materials and Methods**

**Animals**

Male Wistar rats (200-250 g) were obtained from the Experimental Animal Center of Harbin Medical University (Harbin, People’s Republic of China). All animal experimental protocols complied with the 'Guide
for the Care and Use of Laboratory Animals’ published by the United States National Institutes of Health. The study was approved by the Institutional Animal Research Committee of Harbin Medical University. All animals were housed at the animal care facility of Harbin Medical University at 25 °C with 12/12-h light/dark cycles and allowed free access to normal rat chow and water throughout the study period. Rats were randomly assigned to different treatment groups.

**Experimental groups**

Diabetes was induced by streptozotocin (STZ) through intraperitoneal as previously reported [20]. Diabetes was induced by tail vein injection of STZ (streptozotocin, 50 mg/kg) dissolved in 0.1 M citrate buffer (pH 4.5), whereas the control group were injected citrate buffer alone by the same route. Glucose (10 %) was added into the drinking water of STZ-treated rats during the first 24 h to reduce hypoglycemic phase following STZ injection. Blood glucose levels of the rats were measured using blood glucose test strips after 72 h of STZ injection, and blood glucose levels over 16.7 mM were accepted as diabetes. Rats were divided into 4, 8, and 12 weeks diabetes groups and diabetes + NaHS treatment groups, the daily administration of 80 μM NaHS intraperitoneal injection in the NaHS treatment group rats. Measurement of blood glucose was repeated after 2 weeks of treatment in each group.

**Morphological changes in the mesenteric secondary arteries of rats**

Ultrastructural alterations of mesenteric secondary arteries of rats were detected by transmission electron microscopy (TEM). Samples were taken at the mesenteric secondary arteries in five rats from each group. Samples for TEM were cut into pieces less than 1 mm³ and fixed in 2.5 % glutaraldehyde in 0.1 M sodium cacodylate buffer (pH 7.4) for 4 h. Tissues were post-fixed in 1 % osmium tetroxide in 1 % KFe(CN)₄, buffered with 0.1 M sodium cacodylate, dehydrated through graded concentrations of ethanol and propylene oxide and subsequently embedded in Epon 812. Ultrathin sections were cut from blocks and mounted on copper grids. Then, the grids were counterstained with lead citrate and uranyl acetate. Photomicrographs were obtained using Zeiss Axiophot microscopes.

**Histological Examination**

Some sections of the mesenteric secondary artery tissue were fixed in 4% paraformaldehyde in PBS (pH 7.4). The fixed mesenteric secondary artery tissues were embedded in paraffin and cut into 4 μm-thick sections. The sections were then stained with Masson trichrome stain to reveal histological changes and areas of interstitial fibrosis [21].

**The changes on the relaxation rate of rat mesenteric secondary artery rings**

The rat mesenteric secondary arteries were isolated, and rings of approximately 2–3 mm in length were prepared. Subsequently, the rings were mounted under a resting tension of 1.0 g in organ baths filled with Krebs–Henseleit solution. Tissues were equilibrated for 1.5 h and washed by Krebs–Henseleit solution every 15 min before each experimental procedure. Isometric changes in tension were recorded with an isometric force transducer (MAY95-transducer data acquisition system). The composition of the Krebs–Henseleit solution (in mM) was NaCl 113; KCl, 4.7; MgSO₄, 1.2; CaCl₂, 2.5; KH₂PO₄, 1.2; NaHCO₃, 25.0; and glucose, 11.6 and this was gassed with 95 % O₂/5 % CO₂ at 37 °C and pH 7.4.

Reagents aqueous solution was introduced directly into the organ bath by an automated pipette. At the beginning of each experiment, KCl (60 mM)-induced contractions were elicited in mesenteric arteries isolated from different groups of rats. After a 30 min washout period, the arteries were precontracted by 1 nM phenylephrine and then the relaxation response to reagents (Calindol, 5 μM or NaHS, 60 μM) were evaluated by cumulative addition to the organ bath. The functionality of the endothelium was also determined after obtaining the reagents induced relaxation response. For this purpose, after an hour of washout period, the arteries were again precontracted by phenylephrine (1 nM), and then the concentration-dependent relaxation response (<70 %) to acetylcholine (0.1 mM) was elicited by cumulative addition to the organ bath.

**Vascular smooth muscle cells of rat thoracic aorta (A7r5) culture**

Vascular smooth muscle cells of rat thoracic aorta (A7r5) were maintained in DMEM containing 10 % fetal bovine serum (FBS) (Gibco-BRL, Life Technologies, Gaithersburg, MD), penicillin (100 IU/ml), and
strectomycin (100 μg/ml) at 37 °C in a humidified chamber containing 5 % CO₂, incubator. The experiments were performed when the cells reached 80–90 % confluence. In all studies, cells were incubated in the low glucose (5.56 mM) medium. In certain selective experiments, cells were subsequently incubated in the high glucose (25 mM) medium for 72 h.

Measurement of [Ca²⁺], with Fluo-4/AM
The VSMCs were placed onto coverslips, which were covered in 12-well culture plates. After 72 h at 37 °C, the VSMCs were washed with PBS and incubated with 5 μM Fluo-4/AM for 30 min at 37 °C in the dark. The cells were rinsed three times with Tyrode’s solution to remove the remaining dye. During experiment, FI (fluorescence intensity) of fluo-4/AM in VSMCs was recorded using a laser-scanning confocal microscope (Olympus, Japan) at an excitation wavelength of 488 nm and an emission wavelength of 530 nm.

In positive and negative group, cells were exposed to CaSR agonists (Calindol, 2 μM, or CaCl₂ 2.5 mM) and BAPTA (5 μM, calcium chelator) recorded for 330 s at 3s intervals. In Calindol experiments, the VSMCs preincubated with specific inhibitor, U73122 (10 μM, PLC-specific inhibitor), Calhex231 (3 μM, CaSR-specific inhibitor), 2-APB (75 μM, IP₃ receptor inhibitor) and TG (10 μM, Ca²⁺-ATP inhibitor) for 20 min before the experiment. In CaCl₂ experiments, preincubated with NiCl₂ (10 mM, Na⁺-Ca²⁺ channel inhibitor) and CdCl₂ (0.2 mM, L-type Ca²⁺ channel inhibitor) 30 min before the procedure. Image analysis was performed offline using Fluoview-FV300 (Olympus, Japan) to select cell regions from which FI was extracted, and further analysis was conducted with Excel (Microsoft) and Origin Version 7.5 software (OriginLab Corporation). [Ca²⁺] changes were expressed as fluorescence intensity representing FI and normalized to initial fluorescence intensity.

Measurement of H₂S production
H₂S production rate was measured as described previously [22]. Briefly, after different treatments, the cells were collected and homogenized in 50 mM ice-cold potassium phosphate buffer (pH 6.8). The flasks containing reaction mixture (100 mM potassium phosphate buffer, 10 mM L-cysteine, 2 mM pyridoxal 5'-phosphate and 10 % (w/v) cell homogenates) and center wells containing 0.5 ml 1 % zinc acetate and a piece of filter paper were flushed with N₂, and incubated at 37 °C for 90 min. The reaction was stopped by adding 0.5 ml of 50 % trichloroacetic acid, and the flasks were incubated at 37 °C for another 60 min. The contents of the center wells were transferred to test tubes each containing 3.5 ml of water, 0.5 ml of 20 mM N,N-dimethyl-p-phenylenediamine sulfate in 7.2 M HCl and 0.5 ml 30 mM FeCl₃ in 1.2 M HCl were added. The absorbance of the resulting solution at 670 nm was measured 20 min later with a Multiskan spectrum microplate spectrophotometer.

Cell viability and proliferation assay
Vascular smooth muscle cells of rat thoracic aorta (A7r5) were cultured in 96-well tissue culture plates (1×10⁶ cells/well) with 10 % FBS for 24 h. Then the serum-free medium was used and cells were exposed to different reagents for another 24 h. Cell viability and proliferation were measured respectively by the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) [23] and 5-bromo-2'-deoxyuridine (BrdU) [10] incorporation assays.

Immunofluorescence study
The VSMCs were placed onto coverslips, which were covered in 24-well culture plates with polylysine. After cultured for 72 h at 37°C, the VSMCs were washed with PBS, fixed with 4 % formaldehyde in PBS for 10 min, blocked in 1 % BSA for 30 min. The cells were incubated with antibody against PCNA (1:100) or the P21 (Santa Cruz, CA) overnight at 4°C. Then, the cells were incubated with secondary IgG (Santa Cruz, CA) (1:1000) conjugated with fluorescein isothiocyanate (FITC), for 1 h at 37 °C and washed in PBS and 0.1 % Tween 20. Fluorescence images were collected with a fluorescence microscope (Leica, Germany).

Total protein extraction from rat mesenteric secondary arteries and cultured vascular smooth muscle cells of rat thoracic aorta (A7r5) and protein analyzed by western blotting
Rat mesenteric secondary arteries and VSMCs were homogenized in 0.5 mL of RIPA buffer prior to being transferred into small tubes and rotated 1 h at 4 °C. Solubilized proteins were collected after centrifugation at 3,000 × g for 30 min. The supernatant was collected and stored at -80 °C. The protein concentration of
each sample was quantified using the enhanced BCA Protein Assay kit. To detect protein levels, protein lysates from each group of cells and tissues were separated by SDS-PAGE and electrotransferred onto a PVDF membrane (Millipore). Polyacrylamide gels (10 %) were used for CSE (42 kDa), CaSR (90-150 kDa), Cyclin D1 (36 kDa), α-smooth muscle actin (42 kDa), calponin (34 kDa), MMP9 (67-92 kDa) and p-CaMK II (54 kDa) protein testing. The nonspecific proteins on membranes were blocked with 5 % nonfat dried skimmed milk powder prepared in TBS + 0.2 % Tween 20 for 2 h at room temperature. Immunoblotting was then performed using 2 μg/mL rabbit anti-rat CSE, calponin, MMP9, p-CaMK II and CaSR antibodies, rabbit anti-human Cyclin D1 and α-smooth muscle actinpolyclonal antibody, respectively. Membrane blots were washed and incubated with anti-rabbit or mouse IgG antibodies at a 1:1,000 dilution. Immunoreactive proteins were then visualized using AP colorimetric analysis. The volume of the protein bands was quantified using a Bio-Rad Chemi EQ densitometer and Bio-Rad Quantity One software (Bio-Rad laboratories, Hercules, USA).

Statistical analysis
Data are expressed as mean ± SEM. All data involving multiple groups were analyzed by one-way ANOVA. Differences between individual groups were analyzed using the Student’s t test. A difference with P<0.05 was considered statistically significant.

Results

General characteristics and blood glucose in diabetic rats
To investigate the in vivo effect of H2S on diabetic vessels, we first established a diabetic rat model by injecting rats with STZ. Compared with the control group, food and water intake and the concentration of blood glucose were significantly increased in the 4, 8 and 12 week diabetic groups. These results demonstrated that our rat diabetic model was established successfully (Table 1).

|                      | Food intake(g/day) | Water intake(ml/day) | Glucose (mM) |
|----------------------|--------------------|----------------------|--------------|
| Control              | 13.2±1.5           | 27±3.1               | 5.2±0.5      |
| Dia 4w               | 39.4±2.3*          | 183±4.9*             | 29.5±5.4*    |
| Dia 8w               | 46.8±2.5*          | 228±5.6*             | 25.6±4.2*    |
| Dia 12w              | 48.6±4.6*          | 231±4.2*             | 24.3±4.5*    |

Exogenous H2S decreased the thickness of smooth muscle in the mesenteric secondary arteries of diabetic rats
To explore the in vivo effect of H2S on the proliferation of smooth muscle in the mesenteric secondary arteries of diabetic rats, the thickness of the mesenteric artery was evaluated by TEM. Compared with the control group, the thickness of the smooth muscle significantly increased and the H2S concentration decreased in the mesenteric arteries of the 4, 8 and 12 week diabetic groups. The thickness of the smooth muscle and the H2S concentration in the groups treated with NaHS obviously improved compared with the diabetic groups (Fig. 1).

Protein expression in the mesenteric arteries of diabetic rats
Our results showed that compared with the control group, the expression of CaSR and CSE decreased while Cyclin D1 expression increased in the 4, 8 and 12 week diabetic groups, while the expression of these proteins reversed in the NaHS treatment groups (Fig. 2).

CaSR or H2S increased the relaxation rate of the mesenteric artery rings of diabetic rats
Functional changes in vessels are critical indicators of vascular injury. To investigate the effects of CaSR and H2S on the functional changes of mesenteric artery rings, we
examined the relaxation of mesenteric artery rings subjected to CaSR or NaHS treatment. Compared with the control group, the relaxation rates of the mesenteric artery rings of the 4, 8 and 12 week diabetic groups decreased dramatically. In contrast, the relaxation rates significantly increased after treatment with NaHS compared with the diabetic groups after treatment with Calindol (5 \( \mu \)M, a CaSR specific agonist) or NaHS (60 \( \mu \)M, \( H_2 S \) donor) (Fig. 3).

**CaSR activation induced an increase in \([Ca^{2+}]\), in VSMCs via the PLC-IP3 signal transduction pathway and modulated the expression of CSE via p-CaMKII**

To investigate the relationship between CSE and CaSR activation, we examined the signal transduction pathways among CaSR, \([Ca^{2+}]\), and CSE. We found that the FI of \([Ca^{2+}]\) was significantly increased in the CaSR agonist (CaCl\(_2\), and Calindol) group compared with the control group. The FI of \([Ca^{2+}]\), was decreased in the Calindol (Calhex231/2-APB/U73122/ TG pretreated) and CaCl\(_2\), (NiCl\(_2\), + CdCl\(_2\), + Calhex231/2-APB/U73122/ TG pretreated) experimental groups compared with the CaSR agonist group. Furthermore, as a negative control, BAPTA had an obvious effect on the FI of \([Ca^{2+}]\). Western blot analysis showed that, compared with the control group, the expression of p-CaMKII and CSE increased in the CaSR agonist group; in contrast, the expression of CSE decreased in the KN93-pretreated (10 \( \mu \)M, calmodulin antagonist) group compared with the CaSR agonist group (Fig. 5).
CaSR activation mediated the CSE/H₂S pathway in VSMCs

To further explore the modulation of CSE/H₂S by CaSR activation, we also examined the effect of CaSR agonists that mediate the CSE/H₂S pathway of VSMCs both in low and high glucose. CaSR agonists (CaCl₂ or Calindol) increased the expression of CaSR compared with the control (5.56 mM glucose, LG) group of VSMCs from the rat thoracic aorta. CaCl₂
Fig. 3. The changes of relaxation rate in mesenteric artery rings after the treatment with Calindol or NaHS (A: The changes of relaxation rate of 4-12w Dia were decreased while 4-12w Dia + NaHS were increased after treated with Calindol. 4-12w Dia vs. Control, *p<0.01; 8w and 12w Dia vs. 4w Dia, *p<0.01; 4w Dia + NaHS vs. 4w Dia, *p<0.05; 8w and 12w Dia + NaHS vs. 8w and 12w Dia, #p<0.01. B: The changes of relaxation rate of 4-12w Dia were decreased while that of 4-12w Dia + NaHS were increased after treated with NaHS. 4-12w Dia vs. Control, #p<0.01; 12w Dia vs. 4w and 8w Dia, *p<0.05; 4w and 8w Dia + NaHS vs. 4w and 8w Dia, *p<0.05; 12w Dia + NaHS vs. 12w Dia, #p<0.01. C: 4-12w Diabetes vs. Control, *p<0.01; Diabetes + NaHS vs. Diabetes, *p<0.05, #p<0.01; and 12w Diabetes vs. 4w and 8w Diabetes, *#p<0.05. n=6-8. Dia means Diabetes). The curves (A and B) show the changes of the relaxation rate at different time, and the bar graph (C) shows the relaxation rate at the 5 min point.

also increased the expression of CSE compared with the control group. In contrast, the expression of CSE was decreased in the pretreated BAPTA groups compared with the CaCl₂ group. Both CaCl₂ and Calindol increased H₂S production in the VSMCs compared with the control group, while H₂S production was decreased in the EGTA (calcium chelator), BAPTA and 2-APB groups compared with the CaSR agonist group (Fig. 4).
control group. These changes obviously increased in the Calindol, CaCl₂, Ca²⁺ production vs. Ca²⁺ vs. Ca²⁺, 2-APB + Ca²⁺ vs. Ca²⁺, *p<0.05; and D: Calindol vs. control, BAPTA + Calindol vs. Calindol, 2-APB + Calindol vs. Calindol, *p<0.05, n=4-5).

In the high glucose (HG, 25 mM glucose) group of VSMCs from the rat thoracic aorta, the expression levels of CaSR and CSE and CSE activity significantly decreased compared with the control group. These changes obviously increased in the Calindol, CaCl₂, NaHS and A23187 (Ca²⁺ transporter) groups compared with the HG group. We also measured H₂S production in the different groups; H₂S concentration was decreased in the HG group compared with the control group, whereas the CaSR agonist increased the H₂S level compared with the HG group (Fig. 6).

*H₂S inhibited the alteration of the VSMC phenotype, as observed with Masson’s trichrome staining and western blot analysis*

A previous study showed that VSMCs proliferate after the phenotypic modulation from a contractile state to a proliferative state [24]. To test the inhibition of H₂S on the proliferation of VSMCs in diabetic rats, we examined the interstitial fibrotic areas of the mesenteric arteries and the expression of related proteins in VSMCs. The results of Masson’s trichrome staining showed that, compared with the control group, the interstitial fibrotic areas of the smooth muscle layer in the diabetic group were significantly increased, while the interstitial fibrotic areas reduced in the diabetes + NaHS group. Western blot analysis showed that the expression of α-smooth muscle actin and calponin decreased while MMP-9 increased in the
CaSR activation mediated the CSE/H₂S pathway, which inhibited VSMC proliferation

Treatment with 25 mM HG significantly increased cell viability and proliferation of A7r5 compared with the control group. However, cell viability and proliferation decreased in the Calindol, CaCl₂, NaHS and A23187 groups compared with the HG group based on the cell viability data (MTT) and cell proliferation analysis (BrdU). Immunofluorescence staining showed that the expression of proliferating cell nuclear antigen (PCNA) increased while P21 decreased in HG group compared with the control group. In contrast, the expression of PCNA and P21 reversed in the Calindol, NaHS and A23187 groups compared with the HG group (Fig. 8).
The proliferation of VSMCs plays a key role in the maintenance of vascular structure and functions. However, its phenotypic alteration leads to vascular remodeling and various proliferative vascular diseases [25]. High glucose induces VSMC proliferation in vitro, mimicking a key process involved in subendothelial arterial wall thickening and subsequent progression to atherosclerosis in diabetic patients [26]. Recently, some studies suggested that vasodilators such as nitric oxide (NO), play a major role in modulating the alteration...
Fig. 7. Masson’s trichrome staining of rat mesenteric arteries and protein expression levels in VSMCs (A: the interstitial fibrotic areas of the smooth muscle layer in the diabetic group were increased while those reduced in the diabetes + NaHS group. a1-a3, 4w-12w Diabetes; b1-b3, 4w-12w Diabetes + NaHS, (40×). B: The α-smooth muscle actin (a), Calponin (b) and MMP-9 (c) expression determined by Western blot. HG vs. LG, *p<0.05 and HG + NaHS vs. HG, #p<0.05. n=4-6).

of phenotype and proliferation of VSMCs in diabetic models [5]. Similar to NO, H₂S has also been suggested to regulate VSMCs proliferation [27]. Growing evidence supports H₂S as an important endogenous modulator of cell proliferation and apoptosis. Overexpression of CSE in human aortic SMCs inhibited cell growth and induced cell apoptosis [28]. The absence of
endogenous H$_2$S in VSMCs in CSE-deficient mice (KO mice) leads to a significant increase in cell growth rate [8]. Clearly, endogenous CSE/H$_2$S limits the proliferation and growth of VSMCs. However, previous to this study, how the CSE/H$_2$S pathway is regulated to affect the proliferation of VSMCs has been unknown. Here, we provide evidence that CaSR regulates the CSE/H$_2$S pathway, which leads to an inhibition of VSMCs proliferation in diabetic models. Our present study showed that the activation of CaSR increased the intracellular calcium concentration in VSMCs, which modulated phosphorylation of CaMK II to elevate CSE activity, and increased the endogenous H$_2$S level to inhibit the proliferation of VSMCs in the diabetic models.
As a G-protein coupled receptor, CaSR initiates intracellular signals through the modulation of a series of intracellular signaling proteins, including G proteins and PLC, and promotes [Ca\textsuperscript{2+}] release from the endoplasmic/sarcoplasmic reticulum [29,30]. Our previous study proved that the activation of CaSR induced the release of Ca\textsuperscript{2+} from endoplasmic/sarcoplasmic reticulum and increased the intracellular calcium concentration [30].

Previous studies suggested that endogenous H\textsubscript{2}S, which is a strong vasodilator, can be produced by cystathionine-gamma-lyase (CSE) or cystathionine-beta-synthase (CBS) [31]. In mammalian cardiovascular tissues, the biosynthesis of H\textsubscript{2}S is mainly catalyzed by CSE [32]. A recent study found that the elevation of [Ca\textsuperscript{2+}], increased CSE activity and the vasodilator production in smooth muscle cells [8, 19]. However, whether CaSR plays a role in the modulation of CSE/H\textsubscript{2}S and, in turn, influences the vasorelaxation and proliferation of VSMCs has been poorly explored. Our data prove that, in response to low glucose, CaSR agonists increased the [Ca\textsuperscript{2+}] of VSMCs in a time-dependent manner. Further, pretreatment with the CaSR inhibitor calhex231, PLC-IP\textsubscript{3} receptor pathway inhibitor (U73122 or 2-APB) or chelating the intracellular free calcium with BAPTA significantly inhibited the increase of [Ca\textsuperscript{2+}], induced by Calindol (Fig. 5A and C) or CaCl\textsubscript{2} (Fig 5B). The results are in accordance with previous study [30, 33] and suggest that CaSR is involved in the increase of [Ca\textsuperscript{2+}], and that the activation of CaSR increased the [Ca\textsuperscript{2+}] through the PLC-IP\textsubscript{3} receptor pathway. Moreover, CaSR and CSE expression and the H\textsubscript{2}S concentration in VSMCs were increased in response to treatment with CaSR agonists (Fig 4A, B). In contrast, chelating extracellular calcium with EGTA or intracellular free calcium with BAPTA inhibits the effect of CaSR activation on H\textsubscript{2}S concentration (Fig 4C, D).

Some studies [8] have demonstrated that H\textsubscript{2}S generation by CSE is physiologically regulated by calcium-calmodulin. In this study, CaSR activation increased the expression of p-CaMK II and CSE (Fig 5D, E), while the calmodulin antagonist KN93 inhibited the expression of CSE; these results indicate that CaSR modulates the expression of CSE through the calcium-calmodulin pathway of VSMCs. The above results also demonstrated that the rates of [Ca\textsuperscript{2+}] increase induced by Calindol are much slower and the elevated [Ca\textsuperscript{2+}] levels persisted longer. Thus, we reasoned that the CaSR-induced upregulation of CSE expression may be related to the kinetics and the sustained increase levels of [Ca\textsuperscript{2+}].

VSMCs proliferation is a critical factor and is characteristic of diabetes or hyperglycemia. In the present study, we generated the diabetic rat model (Table 1) and confirmed that the SMCs proliferated in the mesenteric arteries (Fig 1A-B). In addition, changes in the relaxation rate of the diabetic rats' mesenteric arteries were evaluated to assess the function of vessels. It has been demonstrated that the H\textsubscript{2}S and Calindol are potent arterial relaxants [18, 34]. The study also suggested that vasodilators play a major role in modulating the proliferation of VSMCs in diabetic models [5]. Some studies have shown that H\textsubscript{2}S dilates and hyperpolarizes animal arteries through activation of the BK\textsubscript{Ca} or K\textsubscript{ATP} channels [35, 36], while the mechanisms of Calindol have remained poorly understood [37]. In this study, we verified that Calindol (5 μM) or NaHS (60 μM) elicited a relaxation response in isolated rat mesenteric arteries (Fig 3A-C), which proved the vasodilator's function and vascular protection of Calindol and exogenous H\textsubscript{2}S. Our data also confirmed that the activation of CaSR and NaHS inhibited the viability and proliferation of VSMCs (Fig 8A, D), and the immunofluorescence of PCNA decreased (Fig 8B) while P21 increased (Fig 8C). These results demonstrated that the activation of CaSR and NaHS inhibited VSMCs proliferation. Furthermore, the expression of CaSR and CSE and the H\textsubscript{2}S concentration decreased in the diabetic and hyperglycemic conditions (Fig 1C and Fig 6A-B, D), while the CaSR agonist increased the expression and activity of CSE and the H\textsubscript{2}S concentration in response to hyperglycemia (Fig 6C-E). Furthermore, pretreatment with 2-APB antagonized the effect of CaSR, and the results of VSMCs were in accordance with the protein expression patterns of the diabetic rat mesenteric arteries (Fig 2A-C) and low glucose condition (Fig 4). However, the expression of CaSR was not statistic significance in mesenteric arteries of diabetic rats at 4 weeks, this maybe involved in the experimental cases insufficient or short intervention time because we showed the reduced trend in the experiment (Fig 2A). The expression of CaSR was decreased in 8-12 weeks (Fig 2B-C) and...
in VSMCs with the treatment of CaSR activator or HG (Fig 4 and Fig 6) in vitro. These data suggest that activation of CaSR modulates the CSE/H$_2$S pathway via the IP$_3$ receptor pathway in response to hyperglycemia. The mechanisms underlying the antiproliferative effect of H$_2$S are multifaceted. A recent study suggested that one of the focal point of these studies is the involvement of the mitogen-activated protein kinase (MAPK) superfamily [31]. Whether the modulation of CaSR on CSE/H$_2$S pathway, which inhibits VSMC proliferation, is related to the MAPK superfamily needs further study.

A previous study [38] suggested that in adult blood vessels, VSMCs proliferate at an extremely low rate; however, in response to vascular injury, proliferation increases dramatically, and the cells become dedifferentiated owing to their inherent plasticity. Depending on the pathological state, the contractile phenotype of VSMCs can be altered from a quiescent contractile to a proliferative synthetic phenotype [39]. In the present study, we have proved that NaHS inhibited the alteration of VSMC phenotype by Masson's trichrome staining (Fig 7A) and western blot analysis (Fig 7B). These results are in accordance with the ultrastructure alteration of rat mesenteric arteries (Fig 1A-B) and elucidated the biological mechanism through which H$_2$S inhibits the proliferation of VSMCs.

CaSR-mediated H$_2$S generation in VSMCs via Ca$^{2+}$ signaling may serve as a common pathway in regulating the proliferation of VSMCs. If so, activation of Ca$^{2+}$ release from the ER through CaSR may attenuate VSMCs proliferation in the hyperglycemic condition. The contribution of CaSR-mediated Ca$^{2+}$ signaling to VSMCs proliferation shown in this study is the first demonstration of an important role for this receptor in diabetes.

**Conclusions**

The current study is the first to demonstrate an important role for CaSR in Ca$^{2+}$ signaling, which is involved in VSMCs proliferation, and this role is mediated by the PLC-IP3 receptor pathway and initiated by CSE/H$_2$S pathway. Our study opens a new avenue to explore the potential roles of Ca$^{2+}$ signaling in regulating VSMCs proliferation and could lead to the development of more effective approaches for treating VSMCs proliferation in diabetic vessels.

**Disclosure Statement**

The authors declare that they have no conflict of interest.

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