Hydrogen Sulfide Ameliorates Blood-Spinal Cord Barrier Disruption and Improves Functional Recovery by Inhibiting Endoplasmic Reticulum Stress-Dependent Autophagy

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Spinal cord injury (SCI) induces the disruption of blood-spinal cord barrier (BSCB), which elicits neurological deficits by triggering secondary injuries. Hydrogen sulfide (H2S) is a gaseous mediator that has been reported to have neuroprotective effect in the central nervous system. However, the relationship between H2S and BSCB disruption during SCI remains unknown. Therefore, it is interesting to evaluate whether the administration of NaHS, a H2S donor, can protect BSCB integrity against SCI and investigate the potential mechanisms underlying it. In present study, we found that SCI markedly activated endoplasmic reticulum (ER) stress and autophagy in a rat model of complete crushing injury to the spinal cord at T9 level. NaHS treatment prevented the loss of tight junction (TJ) and adherens junction (AJ) proteins both in vivo and in vitro. However, the protective effect of NaHS on BSCB restoration was significantly reduced by an ER stress activator (tunicamycin, TM) and an autophagy activator (rapamycin, Rapa). Moreover, SCI-induced autophagy was remarkably blocked by the ER stress inhibitor (4-phenylbutyric acid, 4-PBA). But the autophagy inhibitor (3-Methyladenine, 3-MA) only inhibited autophagy without obvious effects on ER stress. Finally, we had revealed that NaHS significantly alleviated BSCB permeability and improved functional recovery after SCI, and these effects were markedly reversed by TM and Rapa. In conclusion, our present study has demonstrated that NaHS treatment is beneficial for SCI recovery, indicating that H2S treatment is a potential therapeutic strategy for promoting SCI recovery.

Keywords: hydrogen sulfide (H2S), spinal cord injury (SCI), blood-spinal cord barrier (BSCB), ER stress, autophagy

INTRODUCTION

The blood-spinal cord barrier (BSCB) is primarily composed of a specialized system of endothelial cells (ECs) and accessory structures, including the basement membrane, astrocytic end feet processes and pericytes. BSCB forms a tight structure owing to the well-developed tight junctions (TJs) proteins and adherens junctions (AJs) proteins that block the entry of molecules into the spinal cord (Bartanusz et al., 2011; Zheng et al., 2017). The degradation of TJs and AJs...
proteins causes BSCB destruction and consequently increases BSCB permeability after spinal cord injury (SCI) (Lee J.Y. et al., 2012). BSCB disruption allows blood cells to infiltrate the injured parenchyma and aggravates secondary injuries, such as local edema, ischemia, focal hemorrhage, and inflammation (Fang et al., 2015; Li et al., 2017; Lee et al., 2018). Thus, targeting the BSCB disruption is considered as a primary therapeutic approach for SCI.

Hydrogen sulfide (H\textsubscript{2}S) is a gaseous molecule with a rotten egg smell that has been recognized as having a toxic effect. Recent studies have also demonstrated that H\textsubscript{2}S plays a crucial role in physiological and biological effects (Abe and Kimura, 1996; Fiorucci et al., 2006; Martelli et al., 2012). As an almost ubiquitous novel gasotransmitter, H\textsubscript{2}S plays an important role in oxidative stress (Xie et al., 2017b), inflammation (Magierowski et al., 2017) and cardiovascular protection (Calvert et al., 2009). Moreover, it has been reported that H\textsubscript{2}S has neuroprotective effect against oxygen-glucose deprivation (OGD)-induced neuron injury (Yu et al., 2017). Recently, H\textsubscript{2}S has been demonstrated to ameliorate homocysteine-induced brain blood barrier disruption in mice (Kamat et al., 2016). However, it is unclear whether H\textsubscript{2}S plays a role in protecting BSCB integrity after SCI.

Under physiological conditions, the aggregation of misfolded proteins in the endoplasmic reticulum (ER) disrupt ER function, and this process is known as ER stress (Lai et al., 2007). Recent studies have reported that ER stress plays a vital role in a range of neurological diseases, including cerebral ischemia, neurodegeneration disorders and SCI (Gong et al., 2017; Mercado et al., 2018; Wu et al., 2018). Previous studies showed that various cells, such as neurons, oligodendrocytes and astrocytes, could trigger ER stress after SCI (Kuang et al., 2016; Mecha et al., 2012; Wu et al., 2016). Inhibiting ER stress markedly protects neurons against SCI (Acioğlu et al., 2016), but the role of ER stress in BSCB disruption after SCI is not completely defined. Increasing evidences have shown that H\textsubscript{2}S is involved in regulating ER homeostasis, and one study indicated that H\textsubscript{2}S can reduce ER stress and apoptosis in cigarette smoke-treated bronchial epithelial cells (Lin et al., 2017). Here, we try to explore whether reduction of ER stress is involved in the protective effect of H\textsubscript{2}S on BSCB restoration after SCI.

Autophagy, a lysosome-dependent self-degradation process, is a vital pathway for the maintenance of cellular homeostasis under both pathological and physiological conditions in CNS (Yang and Klionsky, 2010; Mizushima and Komatsu, 2011). A previous study suggests that autophagy are activated to remove dysfunctional proteins caused by ER stress, indicating that ER stress may be a trigger for the activation of autophagy (Ding and Yin, 2008; Rzymski et al., 2010). In addition, a large number of studies have shown that autophagy plays an important role in acute injuries, such as SCI and traumatic brain injury (He et al., 2016; Bai et al., 2017; Xie et al., 2017a). Recently, some studies have indicated that H\textsubscript{2}S is associated with autophagy. It has showed that H\textsubscript{2}S inhibits autophagy by upregulating the PI3K/AKT pathway during therapy for myocardial fibrosis (Xiao et al., 2016). Additionally, H\textsubscript{2}S is involved in restoring cardioprotection from post-conditioning by increasing autophagy (Xiao et al., 2015; Chen et al., 2016). However, evidences concerning the role of autophagy during H\textsubscript{2}S-mediated protection against BSCB disruption after SCI have not been reported, and the relationship between autophagy and ER stress during H\textsubscript{2}S treatment for SCI remains unclear.

In the present study, we had investigated the effect of H\textsubscript{2}S on SCI recovery in vivo and in vitro, and discovered that H\textsubscript{2}S alleviated the BSCB disruption and permeability, consequently improved functional recovery after SCI. Mechanistic studies had also demonstrated that H\textsubscript{2}S prevent the loss of TJ and AJ proteins, including P120, β-catenin and Occludin, in vivo and in OGD-treated ECs. Additionally, we had further explored the role of ER stress and autophagy during H\textsubscript{2}S treatment for SCI.

**MATERIALS AND METHODS**

**Animals and Drug Administration**

Eight-week-old adult female Sprague-Dawley rats (200–220 g, n = 80) were acquired from the Animal Center of the Chinese Academy of Sciences (Shanghai, China). The animals were housed under a 12 h light/dark cycle at 21–23°C and given ad libitum access to food and water. The protocol for the care and use of animals conformed to the guidelines from the National Institutes of Health. All experiments were approved by the Laboratory Animal Ethics Committee of Wenzhou Medical University. For drug administration, NaHS (Sigma-Aldrich, St. Louis, MO, United States) was dissolved in phosphate-buffered saline (PBS). The rats were intraperitoneally (i.p.) injected with NaHS (5.6 mg/kg) 30 min before SCI and were injected with the same dose of NaHS daily for 7d. The ER stress activator tunicamycin (TM) (10 μg/kg), the ER stress inhibitor 4-phenylbutyric acid (4-PBA) (100 mg/kg), the autophagy activator rapamycin (Rapa) (0.5 mg/kg) and the autophagy inhibitor 3-methyladenine (3-MA) (2.5 mg/kg) were i.p. injected immediately after SCI. TM, 4-PBA, Rapa, and 3-MA were purchased from Sigma (Sigma-Aldrich, St. Louis, MO, United States). The control group received the equivalent volume of saline for the same duration. Every effort was made to minimize the pain and discomfort of the animals.

**Induction of Spinal Cord Injury Rat Model**

To induce a SCI model, the rats were anesthetized with 10% (w/v) choral hydrate (3.6 mL/kg, i.p.). The skin and muscles adjacent to the spinous processes were incised to expose the vertebral column, and a laminectomy was then performed at the T9 level. The exposed spinal cord was subjected to a moderate crushing injury using a vascular clip (30 g force, Oscar, China) for 1 min (Zheng et al., 2016). The sham group underwent identical procedures but sustained no impact injury. Postoperative monitoring included manual emptying of the bladder twice a day. All rats showed no remarkable side effects resulting from the drug treatment.

**Evans Blue Dye Assays**

The integrity of BSCB was evaluated using Evans blue (EB) dye. The rats were injected with 2% EB dye (2 mL/kg) 1d after SCI by intravenous tail injection. Two hour after the injection, the rats

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were anesthetized and sacrificed via intra-cardiac perfusion with 0.9% saline. Then Spinal cord tissues were stripped and captured.

Locomotion Recovery Assessment
The Basso Beaattie Bresnahan (BBB) scale and footprint analysis were used to analyze locomotion recovery (Li et al., 2018). The scores ranged from 0 (complete paralysis) to 21 (normal locomotion). The locomotor activity of the rats was evaluated in an open experimental field for 5 min. The animals were evaluated 0, 1, 3, 5, 7, and 14d after surgery. The footprint analysis was performed by dipping the rat's posterior limb in a red dye and the fore limb in a blue dye. Outcome measures were obtained by five independent investigators who were blinded to the experimental conditions.

Cell Culture and Oxygen-Glucose Deprivation
HUVECs were expanded and maintained in endothelial cell medium (ECM, ScienCell, Carlsbad, CA, United States) supplemented with 1% ECGS (ScienCell, Carlsbad, CA, United States), 5% FBS (ScienCell, Carlsbad, CA, United States), and antibiotics (100 µg/mL streptomycin and 100 U/mL penicillin, ScienCell, Carlsbad, CA, United States) in a humidified atmosphere of 5% CO₂ and 95% air at 37°C. The forth generation of cells was used in our study. For the OGD treatment, the cells were refreshed with glucose-free DMEM containing TM (3 mM) or Rapa (100 nM) and immediately placed in a sealed chamber loaded with a mixed gas of 5% CO₂ and 95% N₂ for 6 h. The cells were pre-treated with NaHS (100 µM) for 2 h before OGD stimulation. All experiments were performed in triplicate.

Western Blotting Analysis
Spinal cord tissue samples were extracted 1d after surgery and immediately stored at −80°C for western blotting. Briefly, the tissues were lysed using RIPA buffer (0.5% sodium deoxycholate, 1% Triton X-100, 1 mM EDTA, 1 mM PMSF, 10µg/mL leupeptin, 20 mM Tris-HCl, pH 7.5, and 150 mM NaCl). In vitro, the cells were washed twice with PBS and lysed in lysis buffer (0.1% SDS, 1% sodium deoxycholate, 1% Nonidet P-40, 25 mM Tris-HCl, pH 7.6, and 150 mM NaCl). Tissue and cell lysates were centrifuged at 12,000 rpm for 10 min at 4°C, and the supernatant was obtained for a protein assay. Protein concentrations were quantified with BCA reagents (Thermo). An 80µg (in vivo) or 40 µg (in vitro) aliquot of protein was separated by SDS-PAGE and transferred onto a PVDF membrane (Bio-Rad). The membrane was blocked with 5% (w/v) non-fat milk (Bio-Rad) in TBST (Tris-buffered saline with 0.1% Tween-20) for 2 h at room temperature, and the membranes were then incubated with the following primary antibodies: GAPDH (1:10000 Bio-world), β-catenin (1:1000, Abcam), P120 (1:1000, Abcam), Occludin (1:1000, Abcam), PDI (1:1000, Abcam), ATF6 (1:1000 Abcam), CHOP (1:1000 CST), Cle-caspase 12 (1:1000 Abcam), phosphor-mTOR(1:1000 Abcam), mTOR (1:1000 Abcam), ATG7(1:1000 Abcam), ATG5(1:1000 Novus), Beclin1 (1:1000, Abcam), and LC3B I/II (1:1000, Novus) at 4°C overnight. The membranes were washed with TBST three times and incubated with the secondary antibodies for 1 h at room temperature. Signals were visualized by a Chemi DocXRS+ Imaging System (Bio-Rad). We analyzed the bands by using the Quantity-One software.

Immunofluorescence Staining
The rat tissues were fixed in 4% paraformaldehyde (PFA) in PBS for 24 h, embedded in paraffin, and then sectioned into 5 µm slices. The sections were deparaffinized, rehydrated and washed three times for 15 min in PBS. For immunofluorescence staining in vitro, the cells were incubated on cover glasses and fixed for 15 min using 4% PFA. Next, the cells were washed in PBS three times for 2 min each. The sections and cells were incubated with 5% BSA in PBS containing 0.1% Triton X-100 at 37°C for 30 min. Then, the sections were incubated overnight at 4°C with the following antibodies: P120 (1:250 Abcam), β-catenin (1:300, Abcam), Occludin (1:200, Abcam), PDI (1:200, Abcam), and LC3B (1:500, Novus). The sections were washed four times with PBS and incubated with AlexaFluo 488 donkey anti-rabbit/mouse secondary antibodies (1:1000) for 1 h at 37°C. The sections were rinsed four times, incubated with DAPI for 10 min, washed with PBS and sealed with a coverslip. All images were captured on a confocal fluorescence microscope (Nikon, A1 PLUS, Tokyo, Japan).

TUNEL Assay
DNA fragmentation was detected using an In Situ Cell Death Detection Kit (Roche, South Francisco, CA, United States), and TUNEL staining was performed 1d after SCI. The sections were deparaffinized and rehydrated. Then, these sections were treated in a 20 µg/ml proteinase K working solution for 20 min at 37°C. The sections were washed three times in PBS and incubated with TUNEL reaction mixture in a dark humidified chamber for 1 h at 37°C. Afterward, the sections were rinsed with PBS and treated with DAPI for 10 min at room temperature. Positive control was obtained with 10U/mL DNase I buffer for 10 min at room temperature before incubation with the TUNEL reaction mixture. Negative control was incubated with the TUNEL reagent without the TdT enzyme. Positive cells were observed under a confocal fluorescence microscope (Nikon, A1 PLUS, Tokyo, Japan) and analyzed by using Image J software.

Statistical Analysis
All data were expressed as the mean ± standard error of the mean (SEM) from at least three independent experiments. Statistically significant differences were evaluated using one-way analysis of variance (ANOVA) with Dunnett’s post hoc test. Differences were considered to be statistical significance when P value < 0.05.

RESULTS
SCI Activates ER Stress and Autophagy
To detect whether ER stress and autophagy were involved in SCI, we had detected the expression levels of ER stress makers (GRP78 and PDI) and autophagy makers (LC3 and Beclin-1) in
a rat model of SCI. It was found that GRP78 and PDI levels were significantly up-regulated after SCI and peaked at 1d and 12 h post-surgery, respectively, (Figures 1A–C). Additionally, the LC3-II/I ratio and Beclin-1 levels were markedly increased 1d after SCI (Figures 1D–F). Based on the above findings, we concluded that SCI activated ER stress and autophagy, and 1 dpi rat model was used to perform the subsequent experiments.

**NaHS Administration Significantly Attenuates SCI-Induced ER Stress and Autophagy**

Here, we try to evaluate whether NaHS administration could attenuate SCI-induced ER stress and autophagy. The rats were randomly divided into three groups: (i) sham group; (ii) SCI group; and (iii) SCI with NaHS treatment (SCI + NaHS) group. It was observed that the levels of GRP78, PDI, p-JNK, ATF6, CHOP, and Cle-Caspase 12 were significantly increased after SCI, and those increases were markedly inhibited by NaHS treatment (Figures 2A, B). Then, we had further detected the expression levels of p-mTOR, ATG5, ATG7, LC3, and Beclin-1. As shown in Figures 2C, D, the expression of ATG5, ATG7, LC3-II/I ratio, and Beclin-1 levels were increased after SCI, and NaHS administration dramatically attenuated the SCI-induced increases of ATG5, ATG7, LC3-II/I ratio, and Beclin-1 levels (Figures 2C, D). Moreover, NaHS treatment significantly blocked SCI-induced decrease of p-mTOR (Figures 2C, D). Additionally, using TUNEL staining, it was found that SCI dramatically triggered cell apoptosis (45.4 ± 2.81 versus 4.15 ± 1.18, **P < 0.001, N = 3) and NaHS treatment ameliorated it (45.4 ± 2.81 versus 24.78 ± 2.33, **P < 0.01, N = 3) (Figures 3A, B). Consistent with TUNEL, NaHS treatment significantly blocked SCI-induced decreases of Bcl-2 expression (0.79 ± 0.07 versus 0.41 ± 0.06, **P < 0.01, N = 3) and increases of Bax expression (1.31 ± 0.09 versus 2.2 ± 0.2, **P < 0.01, N = 3) (Figures 3C–E). Taken together, the above results indicate that NaHS treatment significantly attenuated the levels of ER stress and autophagy after SCI.

**SCI Induces the Loss of TJ and AJ Proteins**

It is well known that the TJs and AJs in the ECs of blood vessels is essential for maintenance of BSCB integrity (Bartanusz et al., 2011; Zheng et al., 2017). Thus, we had evaluated the expression levels of Occludin, P120 and β-catenin to determine whether levels of TJ (Occludin) and AJ (P120 and β-catenin) proteins were altered in response to SCI. Compared with sham group, Occludin, P120 and β-catenin were significantly decreased 6 h, 12 h, 1d, and 3d after SCI (Figures 4A–D). In conclusion, consistency with previous studies (Lee J.Y. et al., 2012), these data suggests that SCI induced the loss of TJ and AJ proteins.

**NaHS Administration Prevents the Loss of TJ and AJ Proteins by Inhibiting ER Stress and Autophagy After SCI**

To determine whether NaHS prevents the loss of TJ and AJ proteins after SCI, we examined the expression levels of TJ and AJ proteins after NaHS treatment for SCI. Western blotting results had showed that the levels of Occludin, P120 and β-catenin in spinal cord were decreased after SCI, which is remarkably blocked by NaHS treatment (Figures 5A–D). To further evaluate whether NaHS treatment prevents the loss of TJs and AJs after SCI by inhibiting ER stress and autophagy, tunicamycin (TM, a specific ER stress activator) and rapamycin (Rapa, a specific autophagy activator) were used to activate ER stress and autophagy, respectively. As shown in Figures 5A–D, both TM and Rapa treatment significantly reversed the protective effect of NaHS against SCI-induced loss of TJ and AJ proteins. Immunofluorescence staining results had further confirmed the effect of NaHS (Figures 5E, F). The above results demonstrates that NaHS treatment prevented the loss of TJs and AJs by inhibiting ER stress and autophagy.

**ER Stress Triggers Autophagy After SCI**

Prior reports have indicated that ER stress can mediate autophagy (Kouroku et al., 2007; Lee H. et al., 2012). To further confirm the connection between ER stress and autophagy during SCI, 4-PBA (a classical ER stress inhibitor) and 3-MA (a classical autophagy inhibitor) were applied to treatment for SCI. It was observed that 4-PBA not only suppressed ER stress, as indicated by the reduction of GRP78 and PDI, but also inhibited autophagy, as evidenced by the reduction of Beclin-1 and the LC3-II/I ratio (Figures 6A–C). However, 3-MA only inhibited the expression of Beclin-1 and the LC3-II/I ratio, without obvious effects on GRP78 and PDI (Figures 6A, D, E). The above results have suggested that SCI-induced ER stress significantly triggered autophagy.

**NaHS Administration Decreases BSCB Permeability and Improves Functional Recovery by Inhibiting ER Stress and Autophagy After SCI**

To determine whether NaHS treatment mitigates SCI-induced increases of permeability after SCI, EB dye was used to examine the effect of NaHS on BSCB permeability at 1d post-SCI. As shown in Figures 7A, B, compared with the sham group, EB dye extravasation was significantly increased in SCI group, suggesting that the integrity of BSCB was disrupted after SCI. After administration of NaHS, the content of EB was markedly decreased compared with that in SCI group. However, both TM and Rapa significantly reduced the effect of NaHS on BSCB integrity.1.31 ± 0.15 versus 0.53 ± 0.09, **P < 0.01; 121 ± 0.15 versus 0.53 ± 0.09, \(*P < 0.05\) (Figures 7A, B). We also assessed the BBB rating scale and footprint analysis to evaluate the locomotor recovery in rats after SCI. There were no significant differences in BBB scores among the SCI, SCI + NaHS, SCI + NaHS + TM, and SCI + NaHS + Rapa groups 1, 3, and 5d after injury. However, 7 and 14d after injury, it was observed that the BBB score of SCI + NaHS group was higher than the SCI + NaHS + TM group (5.83 ± 0.65 versus 1.31 ± 0.15, **P < 0.01; SCI + NaHS + TM versus SCI + NaHS, 5.83 ± 0.65 versus 3.67 ± 0.56, \(*P < 0.05, N = 10\), SCI + NaHS + Rapa groups (5.83 ± 0.65 versus 3.33 ± 0.42, **P < 0.01, N = 10) (Figures 7C–E). In footprint analysis, NaHS-treated rats exhibited coordinated....
NaHS Administration Inhibits ER Stress and Autophagy in OGD-Treated ECs

Here, we had further determined the effect of NaHS treatment on ER stress and autophagy in vitro using OGD treating HUVECs model. Compared with the control group, OGD treatment had up-regulated the expression levels of GRP78 and PDI, and these increases was significantly inhibited by NaHS administration (Figures 8A–C). In addition, consistence with the western blotting results, the OGD-treated group exhibited an increased fluorescence intensity of PDI. However, NaHS treatment markedly weakened the fluorescence intensity of PDI (Figure 8G). Furthermore, we had also detected the expression levels of the markers of autophagy. Compared with control group, the levels of Beclin-1 and LC3-II/I ratio were significantly increased in the OGD-treated group, which was dramatically blocked by NaHS administration (Figures 8D–F). Moreover, the immunofluorescence staining findings had also showed that the OGD group exhibited an enhanced fluorescence intensity of LC3 when compared with that of the control group, which was significantly decreased after treatment with NaHS (Figure 8H). Taken Together, the above findings indicate that...
NaHS treatment significantly inhibited ER stress and autophagy in vitro.

**NaHS Administration Attenuates the Loss of TJ and AJ Proteins by Inhibiting ER Stress and Autophagy in vitro**

To further determine whether the protective effect of NaHS on BSCB is related to the inhibition of ER stress and autophagy, NaHS, TM, and Rapa were administered to HUVECs. Consistent with the findings obtained in vivo, the expression levels of P120, β-catenin and Occludin were decreased in the OGD group and significantly up-regulated in the OGD + NaHS group. However, both TM and Rapa markedly weakened the protective effect of NaHS (Figures 9A–D). In addition, immunofluorescence staining results had also showed that the intensity of β-catenin and Occludin in the OGD group was decreased compared to that of the control group. However, NaHS treatment significantly
FIGURE 3 | NaHS treatment blocked SCI-triggered apoptosis at 1d after SCI. (A,B) TUNEL staining in the sham, SCI model and SC+NaHS-treated groups, Scale bar = 50 µm. (C–E) Representative western blots of Bax and Bcl2 in the sham, SCI model and SC+NaHS-treated groups. All data were presented as the mean ± SEM, n = 3. **P < 0.01 and ***P < 0.001 versus the indicated group.

reversed the destructive effect of OGD, as evidenced by the increased intensity of β-catenin and Occludin (Figures 9E,F). Both TM and Rapa abolished such effect of NaHS. These data indicates that NaHS effectively attenuated the loss of TJs and AJs by inhibiting ER stress and autophagy in OGD-treated ECs.

DISCUSSION

Spinal cord injury is a devastating neurological disease that affects thousands of patients every year. A series of secondary injuries, including vascular changes, oxidative stress, and apoptosis, are considered as the major causes of disability (Fan et al., 2013; Colon et al., 2016; Izmailov et al., 2017). Thus, it is important to explore the effective therapeutic treatments for SCI and reduce SCI-induced tissue damage and neurological disorders. In present study, we had demonstrated that NaHS administration effectively decreases the degradation of TJ and AJ proteins, prevents BSCB permeability, and ultimately protects the spinal cord against secondary injury to improve functional recovery. The inhibitions of ER stress and autophagy were the potential molecular mechanisms underlying NaHS treatment for SCI.

As a H₂S donor, NaHS has been widely used to assess the therapeutic potential of exogenous H₂S delivery. Previous study employed aqueous NaHS solutions to assess the rat aortic ring response in vitro (Zhao et al., 2001). NaHS mitigated the degree of acute lung injury by attenuating IL-6 and IL-8 levels, while increasing IL-10 levels in the lung tissue and plasma (Li et al., 2008). Moreover, NaHS exhibited efficacy
FIGURE 4 | SCI induces the loss of TJ and AJ proteins. (A) Representative western blotting results for TJ proteins (Occludin) and AJ proteins (P120 and β-catenin) in the sham group, 6 h, 12 h, 1d, 3d, and 7d after SCI group. (B–D) Quantification of the expression of Occludin, P120, and β-catenin. The relative band density value was normalized to that of GAPDH. All data are presented as the mean ± SEM, n = 5. *P < 0.05, **P < 0.01, and ***P < 0.001 versus the sham group.

in preventing cell damage caused by ROS in brain cells (Whiteman et al., 2005). Recent studies had also indicated that NaHS has a protective effect in peripheral nerve injury and spinal cord ischemia reperfusion (Park et al., 2015; Xie et al., 2017b). Our findings had indicated that NaHS could improve functional recovery after SCI, suggesting that NaHS probably represents a potential therapeutic approach for traumatic SCI.

It is well known that the maintenance of BSCB integrity effectively promotes central nervous system (CNS) recovery after SCI, and TJ and AJ proteins are the crucial components of BSCB (Lee J.Y. et al., 2012). The disruption of BSCB allows neutrophils and other immune cells to infiltrate the area of injury, resulting in serious secondary injuries (Penas et al., 2007). Therefore, regulation of TJ and AJ proteins may be the potential mechanism of NaHS-induced neuroprotective role during SCI. In addition, it has been demonstrated that the administration of NaHS protects the blood-brain barrier (BBB) integrity after cerebral ischemia and traumatic brain injury (Wang et al., 2014; Xu et al., 2018). The BSCB is analogous to the BBB in that it is selectively permeable and blocks the entry of blood cells and other molecules into the CNS (Bartanusz et al., 2011). Thus, we evaluated the effect of NaHS on the BSCB after SCI. We discovered that NaHS treatment reduced the leakage of EB dye. In addition, the decreases of TJ (P120, β-catenin) and AJ (Occludin) proteins were significantly inhibited by NaHS treatment when comparing with the permeability of the BSCB during SCI. Furthermore, NaHS treatment inhibited the degradation of TJ and AJ proteins under OGD conditions *in vitro*. Taken together, these findings indicated that NaHS treatment promotes to maintain BSCB integrity following SCI.

ER stress occurs when misfolded proteins accumulate in the ER lumen and cause ER dysfunction, which is known as unfolded protein response (UPR) (Schroder and Kaufman, 2005). ER stress is one of the crucial molecular mechanisms underlies the pathogenesis of SCI. Recent studies indicated that the inhibition of ER stress prevents the disruption of the BSCB (Zheng et al., 2017), and NaHS treatment could suppress the activation of ER stress (Xu et al., 2017). Consistent with the prior study, we had evaluated the role of ER stress in the effect of NaHS on BSCB restoration after SCI or OGD, and found that NaHS treatment inhibited the activation of ER stress. Furthermore, the activation of ER stress with TM significantly reversed the protective effect of NaHS on BSCB restoration and consequently functional motor recovery after SCI. These data collectively suggest that NaHS treatment rescue the BSCB disruption from SCI via inhibiting ER stress.
**FIGURE 5** NaHS prevents the loss of TJ and AJ proteins by inhibiting ER stress and autophagy at 1 d post-SCI. (A) TM and Rapa were applied to specifically activate ER stress and autophagy, respectively. The rats were randomly divided into five groups: (i) the Sham group; (ii) the SCI model group; (iii) the SCI model with NaHS treatment (SCI + NaHS) group; (iv) the SCI model with NaHS and TM treatment (SCI + NaHS + TM) group; and (v) the SCI model with NaHS and Rapa treatment (SCI + NaHS + Rapa) group. Representative western blots of P120, β-catenin, and Occludin from each group. (B–D) Quantification of the expression of P120, β-catenin, and Occludin. The relative band density value was normalized to that of GAPDH. All data are presented as the mean ± SEM, n = 5. **P < 0.001** versus the sham group. *P < 0.05 and **P < 0.01 versus the indicated group. (E,F) Representative fluorescence images of β-catenin and P120 (Green) in sections from the spinal cord tissue in each group. Scale bar = 50 µm, n = 5.
Autophagy, a catabolic cellular organelle process, removes unwanted cellular components through double-membrane autophagosomes fused with lysosomes, and, thus, is essential for survival, differentiation, development and homeostasis (Noda and Inagaki, 2015). Previous studies had demonstrated that autophagy exerts a destructive role during SCI and inhibiting autophagy can promote locomotor functional recovery of mice (Wu et al., 2018). Consistent with prior study, our findings had revealed that autophagy is markedly activated during the acute phase of SCI. NaHS administration downregulated the expression of Beclin1 and the LC3-II/I ratio in vivo and in vitro, and prevented the loss of TJs and AJs protein. However, the role of autophagy in SCI and the reciprocal regulation between H\textsubscript{2}S and autophagy remained controversial. Recent study had reported that H\textsubscript{2}S promotes autophagy via the PI3K/Akt/mTOR signaling pathway in hepatocellular carcinoma cells (Wang et al., 2017). Moreover, autophagy stimulation or enhancement can improve neuron protection and functional recovery during SCI (Bai et al., 2017; Hu et al., 2017). It is well known that the pathological status of spinal cord is changing after SCI. We speculate that the different periods of SCI maybe the caused factor for the different role of autophagy and H\textsubscript{2}S on SCI, especially in acute phase and chronic phase.

Our mechanism studies have shown that ER stress and autophagy activation are both involved in NaHS treatment for BSCB disruption during SCI. However, whether ER stress has a cross-talking with autophagy during NaHS treatment for BSCB is still unclear. Prior and our current study has demonstrated that there are mutual cross-talking between ER stress and autophagy (Hoyer-Hansen and Jaattela, 2007). Additionally, some studies have indicated that activated ER stress can trigger autophagy (Chandrika et al., 2015; Feng et al., 2017). Furthermore, ER stress can induce the initiation of autophagosome formation via the IRE1/JNK signaling pathway (Ogata et al., 2006) and PERK/eIF2\alpha signaling pathway (Kouroku et al., 2007; B’Chir et al., 2013). In our present study, 4-PBA markedly blocked SCI-induced autophagy, however, the autophagy inhibitor (3-MA) can only inhibit autophagy activity without remarkable effects on ER stress. These results suggests that NaHS treatment blocked SCI-induced ER stress and ER stress-associated autophagy, subsequently ameliorated SCI.

An interesting finding in present study was verified that inhibition of autophagy and ER stress contributes to the H\textsubscript{2}S treatment for SCI. It has been demonstrated that H\textsubscript{2}S regulates oxidative stress (Kimura et al., 2010; Xie et al., 2017b). But the role of H\textsubscript{2}S on oxidative stress remained controversial. Some studies...
FIGURE 7 | NaHS prevents the disruption of BSCB and improves the functional recovery by inhibiting ER stress and autophagy after SCI. (A,B) Representative whole spinal cord tissues and quantification of BSCB permeability for each group, indicating that EB dye permeabilized into the spinal cord. All data are presented as the mean ± SEM, n = 5. *P < 0.05 and **P < 0.01 versus the SCI + NaHS group. (C) The BBB locomotion scores of the different groups 1, 3, 5, 7, and 14d after SCI, n = 10. (D,E) Quantification of the BBB locomotion scores at 7 and 14d from (C). All data are presented as the mean ± SEM, n = 10. *P < 0.05, **P < 0.01 and ***P < 0.001 versus the SCI + NaHS group. (F) Footprint analysis results from the different groups.
FIGURE 8 | NaHS treatment inhibits the activation of ER stress and autophagy in OGD-treated HUVECs. (A) Representative western blots of GRP78 and PDI in HUVECs subjected to OGD and treated with NaHS. (B,C) Quantification of the expression of GRP78 and PDI. The relative band density value was normalized to that of GAPDH. All data are presented as the mean ± SEM. *P < 0.05 and **P < 0.01 versus the OGD group. (D) Representative western blots of Beclin1 and LC3B II for each group after OGD. (E,F) Quantification of the expression of Beclin1 and LC3B II. All data are presented as the mean ± SEM. *P < 0.05 and **P < 0.01 versus the OGD group. (G,H) Representative fluorescence images of PDI and LC3 (Green) in HUVECs. The nucleus is labeled with DAPI (blue). Scale bar = 50 μm (PDI), and scale bar = 25 μm (LC3), n = 3.

demonstrate that H₂S directly suppress oxidative stress (Kimura et al., 2010), whereas the others studies have demonstrate H₂S has a deleterious effect on oxidative stress via increasing the formation of reactive oxygen species and leading to glutathione (GSH) depletion (Truong, 2006). The present study did not further provide the evidence of the relationship between H₂S and oxidative stress in the disruption of BSCB after SCI. Some studies has reported that concentrations of H₂S is the conclusive factor for its significant physiological and toxicological roles. It has demonstrated that the concentrations (~15 mM) of H₂S quickly
FIGURE 9 | NaHS protects OGD-treated HUVECs by inhibiting ER stress and autophagy in vitro. (A) Representative western blots of P120, β-catenin and Occludin for each group. (B–D) Quantification of the expression of P120, β-catenin and Occludin. The relative band density value was normalized to that of GAPDH. All data are presented as the mean ± SEM. **P < 0.001 versus the Con group. #P < 0.05 and ##P < 0.01 versus the indicated group. (E,F) Representative fluorescence images of β-catenin and Occludin (Green) in HUVECs for each group. The nucleus is labeled with DAPI (blue). Scale bar = 50 μm, n = 3.
In our current study, NaHS (5.6 mg/kg) and NaHS (100 μM) treatment was used in vivo and in vitro, which is less than 0.1 mM. Therefore, it is reasonable to speculate that the NaHS treatment can inhibit oxidative stress and ameliorate the disruption of BSCB after SCI in our study.

In summary, our study has demonstrated that NaHS administration significantly ameliorates the disruption of BSCB and subsequently improves functional motor recovery after SCI. Inhibition of ER stress and ER stress-associated autophagy were involved in NaHS treatment for SCI. Our finding suggested that rational H₂S treatment promotes the recovery of SCI, which is the potential therapeutic strategies for SCI.

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AUTHOR CONTRIBUTIONS
JX and HX conceived and designed the experiments. HW and WH performed the experiments. HW and YW performed statistical analysis and wrote the paper. JL, KX, ZL, QW, YL, LX, JW, and HHI provided assistance with experiments. All authors discussed the results and approved the final manuscript.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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