Oxidative Stress Inhibits Vascular $K_{ATP}$ Channels by S-Glutathionylation

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The $K_{ATP}$ channel is an important player in vascular tone regulation. Its opening and closure lead to vasodilation and vasoconstriction, respectively. Such functions may be disrupted in oxidative stress seen in a variety of cardiovascular diseases, while the underlying mechanism remains unclear. Here, we demonstrate that S-glutathionylation was a modulation mechanism underlying oxidant-mediated vascular $K_{ATP}$ channel regulation. An exposure of isolated mesenteric rings to hydrogen peroxide $(H_2O_2)$ impaired the $K_{ATP}$ channel-mediated vascular dilation. In whole-cell recordings and inside-out patches, $H_2O_2$ or diamide caused a strong inhibition of the vascular $K_{ATP}$ channel (Kir6.1/SUR2B) in the presence, but not in the absence, of glutathione (GSH). Similar channel inhibition was seen with oxidized glutathione (GSSG) and thiol-modulating reagents. The oxidant-mediated channel inhibition was reversed by the reducing agent dithiothreitol (DTT) and the specific deglutathionylation reagent glutaredoxin-1 (Grx1). Consistent with S-glutathionylation, streptavidin pull-down assays with biotinylated glutathione ethyl ester (BioGEE) showed incorporation of GSH to the Kir6.1 subunit in the presence of $H_2O_2$. These results suggest that S-glutathionylation is an important mechanism for the vascular $K_{ATP}$ channel modulation in oxidative stress.

ATP-sensitive K$^+~(K_{ATP})$ channels are regulated by intracellular ATP/ADP and couple intermediary metabolic states to membrane excitability. The activity of these channels is low under physiological conditions and drastically rises during metabolic stress (1, 2). The $K_{ATP}$ channels are expressed in almost all tissues. In vascular smooth muscle cells (SMCs), activation of $K_{ATP}$ channels by several vasodilators reduces SMC membrane excitability, leading to vasorelaxation (3, 4). Activity of the channels is inhibited by vasoconstrictors (5), resulting in depolarization of the SMCs and vasoconstriction. Such a common vascular regulator is thus targeted by a variety of cellular events in physiological and pathological conditions (1, 6).

One important cellular event is oxidative stress that is known to play an important role in the development and maintenance of several cardiovascular diseases, such as hypertension, atherosclerosis, and diabetic vascular complications (7, 8). During oxidative stress, excessive reactive oxygen species (ROS), such as superoxide (O$^-$), hydroxyl radical (OH$^-$), and hydrogen peroxide $(H_2O_2)$, are overly produced causing vascular dysfunction and structural damages (9, 10). Previous studies indeed have shown that O$^-$ suppresses $K_{ATP}$ channels and blunts the pial arterial dilation responses (11). In diabetic patients, in whom oxidative stress is evident, $K_{ATP}$ channel function is disrupted, leading to impaired vasodilation responses (12). In insulin-resistant rats, the $K_{ATP}$ channel-dependent vasodilation is also impaired, which is likely to be mediated by ROS (13). Although the dysfunction of $K_{ATP}$ channels in oxidative stress has been documented, the mechanism underlying channel modulation remains unknown (9, 14).

ROS can modulate proteins by intra- and intermolecular thiol oxidation (15, 16), which may be the underlying cause for the modulation of vascular $K_{ATP}$ channels in oxidative stress. To test this hypothesis, we performed studies using a combined molecular biology, electrophysiology, and biochemistry approach. Our results showed that the Kir6.1/SUR2B channel, the major isoform of vascular $K_{ATP}$ channels, was inhibited by micromolar concentrations of $H_2O_2$ as well as several other oxidants via S-glutathionylation.

MATERIALS AND METHODS

Chemicals and Reagents—Unless stated, all reagents and chemicals used in this study were purchased from Sigma. $H_2O_2$ and glutathione (GSH) were freshly made and used within 4 h. Other reagents were prepared as concentrated stocks in double-distilled water or dimethyl sulfoxide (DMSO). In cases where DMSO was used, the final concentration of DMSO in solution was <0.1% (v/v). At this concentration, DMSO did not have any detectable effect on the channel activity.

Mesenteric Artery Preparation and Tension Measurements—All animal experiments were performed in compliance with an approved protocol by the Institutional Animal Care and Use Committees (IACUC) at Georgia State University. Male Sprague-Dawley rats (200–250 g body weight) were deeply anesthetized and sacrificed. Mesenteric arteries were dissected, and connective tissues were removed in physiological saline solution (PSS) containing the following (concentrations in mM): NaCl, 140; KCl, 4.6; CaCl$_2$, 1.5; MgCl$_2$, 1; glucose, 10; and HEPES, 5 (pH 7.3). The arteries were cut into small rings (2 mm
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in length) and transferred to ice-cold Krebs solution containing the following (concentrations in mM): NaCl, 118.0; NaHCO<sub>3</sub>, 25.0; KCl, 3.6; MgSO<sub>4</sub>, 1.2; KH<sub>2</sub>PO<sub>4</sub>, 1.2; glucose, 11.0; and CaCl<sub>2</sub>, 2.5. The arterial ring was mounted on a force-electricity transducer (Model FT-302; iWorx/CBSciences Inc. Dover, NH) for measurements of isometric force contraction in a 5-ml tissue bath containing air-bubbled Krebs solution. Tissue vitality was confirmed with the contraction response to phenylephrine (PE). When endothelium-free rings were needed, the rings were rubbed with sanded polyethylene tubing. The endothelium-free rings were then tested for PE contraction followed by acetylcholine (ACh, 1 μM) for relaxation. The rings were considered to be endothelium-free, if >90% relaxation was eliminated. PE and ACh were then washed out, and the rings were allowed to equilibrate in the Krebs solution for another 30–60 min before the experiments.

Expression of K<sub>ATP</sub> Channel in HEK293 Cells—K<sub>ATP</sub> channels were expressed in HEK293 cells because these cells have low numbers of endogenous channels. The HEK293 cells were cultured in Dulbecco’s modified Eagle medium (DMEM)/F12 medium with 10% fetal bovine serum and penicillin/streptomycin at 37 °C with 5% CO<sub>2</sub>. A eukaryotic expression vector pcDNA3.1 was used to express rat Kir6.1 (GenBank<sup>TM</sup> No. D42145) in cells with SUR2B (GenBank<sup>TM</sup> No. D86038; mRNA isoform, NM_011511). A 35-mm Petri dish of cells was transfected with 1 μg of Kir6.1 or Kir6.2 and 3 μg of SUR2B using Lipofectamine<sup>2000</sup> (Invitrogen Inc., Carlsbad, CA). To facilitate the identification of positively transfected cells, 0.4 g of green fluorescent protein (GFP) cDNA (pEGFP-N2; Clontech, Palo Alto, CA) was included in the cDNA mixture. One day after transfection, cells were dissociated with 0.25% trypsin, split, and transferred to coverslips for further growth. Experiments were performed on the cells in coverslips during the following 12–48 h.

Electrophysiology—Patch-clamp experiments were carried out at room temperature as described previously (3–5, 17, 18). In brief, fire-polished patch pipettes with 2–5 MΩ resistance were made of 1.2-mm borosilicate glass capillaries. Whole-cell currents were recorded in single-cell voltage clamp with a holding potential of 0 mV and step to −80 mV. The bath solution contained the following (concentration in mM): KCl, 10; potassium gluconate, 135; EGTA, 5; glucose, 5; and HEPES, 10 (pH 7.4). The pipette was filled with a solution containing the following (concentration in mM): KCl, 10; potassium gluconate, 133; EGTA, 5; glucose, 5; K<sub>2</sub>ATP, 1; NaADP, 0.5; MgCl<sub>2</sub>, 1; and HEPES, 10 (pH = 7.4). To avoid nucleotide degradation, all intracellular solutions were freshly made and used within 4 h. All the recordings were made with the Axopatch 200B amplifier (Axon Instruments Inc., Foster City, CA). The data were low-pass filtered (2 kHz, Bessel 4-pole filter, −3 dB) and digitized (10 kHz, 16-bit resolution) with Clampex 9 (Axon Instruments Inc.). Macroscopic currents were recorded from giant inside-out patches, and single-channel currents were recorded from small inside-out patches with a constant single voltage of −80 or −60 mV. Symmetric high K<sup>+</sup> (145 mM in total) was used in both bath and pipette solutions and K<sub>2</sub>ATP (1 mM), and NaADP (0.5 mM), were included in the bath solution to maintain the channel activity. Higher sampling rate (20 kHz) was used to digitize the currents recorded from inside-out patch. Data were analyzed using Clampfit 9 (Axon Instruments Inc.).

Immunohistochemistry—Immunohistochemistry was performed on the HEK293 cells with and without Kir6.1/SUR2B transfection. Two hours before the experiments, the culture medium was replaced with fresh medium. Biotinylated glutathione ethyl ester (BioGEE; 250 μM; Invitrogen) was added to the medium and incubated for 1 h followed by an H<sub>2</sub>O<sub>2</sub> (750 μM) challenge for 15 min. The medium was then discarded, and the cells were washed three times with phosphate-buffered saline (PBS) (containing 0.3% Triton X-100) to remove the excessive free BioGEE that had not conjugated with the proteins. Cells were then fixed with 4% paraformaldehyde for 30 min followed by three washes with PBS. Dylight-488-conjugated streptavidin was diluted 1:1000 in PBS and added to the cells for 1 h of incubation at room temperature. After three PBS washes, the cells were examined under the LSM 510 confocal microscope (Zeiss). For double staining, the cells were further incubated with rabbit primary antibody against Kir6.1 for 2 h followed by three washes. Dylight-594-conjugated goat anti-rabbit secondary antibody (1:1000; Jackson ImmunoResearch) was used to visualize the Kir6.1 staining. Experiments were repeated three times.

Streptavidin Pull-down Assay and Western Blot—HEK293 cells expressing Kir6.1/SUR2B channels and the A10 smooth muscle cell line were used for this experiment. A10 cells were cultured in DMEM with 10% fetal bovine serum (FBS) at 37 °C in humidified atmosphere with 5% CO<sub>2</sub>. BioGEE and H<sub>2</sub>O<sub>2</sub> treatments were performed as described above in Immunohistochemistry. The cells were then washed once and lysed using RIPA buffer (Sigma). Samples were run on 10% SDS-polyacrylamide non-reducing gel and then transferred to a nitrocellulose membrane (Bio-Rad). Rabbit primary antibodies against Kir6.1 (1:500; Sigma) and secondary antibodies conjugated with alkaline phosphatase were used in the Western blot (1:10,000; Jackson ImmunoResearch). Signals were visualized by SuperSignal West Pico Chemiluminescent Substrate (Thermo Scientific).

Biotin-GSH-conjugated proteins were pulled down using streptavidin-dynabeads according to the instructions provided by Invitrogen. Briefly, the beads were washed three times before the immobilization. Samples were then mixed with beads and incubated at room temperature with gentle rotation for 30 min. A magnet was used to separate the biotinylated molecules-bead complex from other unlabelled proteins. Supernatant containing unlabelled proteins was discarded, and the pellet was resuspended followed by three washes. The biotinylated molecules-beads complex was then resuspended in the loading buffer with 0.1% SDS and boiled so that glutathionylated proteins were released into the solvent for further analysis. Experiments on HEK293 cells transfected with the Kir6.1/SUR2B channel were repeated four times, and experiments on A10 cells were repeated five times.

Data Analysis—Data were presented as means ± S.E. Differences were evaluated using Student’s t-tests or ANOVA, and statistical significance was accepted when p < 0.05.
The results of whole-cell recordings may be affected by washout or inadequate controls of cytosolic soluble factors that can be potentially involved in the channel modulation, such as endogenous GSH and GSSG. Therefore, further studies were performed in excised patches. In giant inside-out patches, millimolar concentrations of H$_2$O$_2$ were required to inhibit the Kir6.1/SUR2B channel in the absence of cytosolic soluble components (Fig. 2, B, E, and F). Strikingly, the administration of small amount of glutathione (GSH) drastically enhanced the channel sensitivity to H$_2$O$_2$. In the presence of 100 μM GSH, H$_2$O$_2$ as low as 10 μM began to inhibit channel activity, and clear concentration dependence was seen with an IC$_{50}$ of 25 μM (Fig. 2, C, E, and F). The effect of H$_2$O$_2$ was subsequently studied in the presence of 2 mM GSH and 40 μM GSSG, a ratio that is close to the physiological concentrations of GSH/GSSG in the cytosol (21). Under this condition, H$_2$O$_2$ also potently inhibited the channel with an IC$_{50}$ of 20 μM (Fig. 2, D–F).

Consistent with these observations in inside-out patches, a supplement of GSH/GSSG (2 mM and 40 μM, respectively) to the pipette solution significantly enhanced the channel sensitivity to H$_2$O$_2$ in whole-cell recordings. With GSH/GSSG in the pipette solution, 100 μM H$_2$O$_2$ inhibited the whole-cell currents by 36.2 ± 8.8% (n = 5) compared with 8.0 ± 5.1% when GSH/GSSG were absent (n = 4, p < 0.05; Fig. 2G). In contrast, the same concentration of H$_2$O$_2$ (100 μM) inhibited the channel by >80% in inside-out patches in the presence of GSH or GSH/GSSG (Fig. 2G). The difference in the H$_2$O$_2$ sensitivity between inside-out patches and whole-cell recordings may be due to the diffusion kinetics across plasma membranes. Using the pseudo-first-order reaction analysis described by Tang et al. (22) in their study on H$_2$O$_2$-mediated BK channel inhibition, we calculated the pseudo first order constant to be 724 M$^{-1}$ min$^{-1}$ based on the average Kir6.1/SUR2B channel inhibition (36.2%) by 100 μM H$_2$O$_2$ (with GSH/GSSG) during a period of <5 min. With this constant, our further calculation showed that a 50% inhibition of the channel was achieved by ~23 μM H$_2$O$_2$ in ~30 min, which implies a protective role of the membrane barrier against a burst of H$_2$O$_2$ (see discussion for details). The requirement of GSH for H$_2$O$_2$ to produce its channel inhibition effect indicates that GSH-mediated protein modifications of the Kir6.1/SUR2B channel, such as S-glutathionylation, are likely to occur when H$_2$O$_2$ is produced as an intermediary metabolite or a product of oxidative stress.

To understand the biophysical mechanisms underlying Kir6.1/SUR2B modulation, we analyzed single channel proper-
ties recorded in regular inside-out patches in the presence of 2 mM GSH and 40 μM GSSG. We found that the channel open-state probability was progressively suppressed with increased H$_2$O$_2$ concentrations while the unitary conductance remained unchanged (Fig. 2).

**GSH-dependent Modulation by Other Oxidants**—Diamide, an oxidant that produces intra- and intermolecular disulfide bonds, is known to cause S-glutathionylation in the presence of GSH (23, 24). In giant inside-out patches, we found that the joint application of diamide and GSH drastically inhibited channel activity within 2–8 min, regardless of the order of application, i.e. 78.4 ± 5.1% inhibition with GSH first and 87.9 ± 8.7% inhibition with diamide first (Fig. 3, A, B, and E). No statistical difference was found between the experiments (*p < 0.05; ***,p < 0.001) between these conditions.

GSSG is another S-glutathionylation inducer (23, 24). In giant inside-out patches, we found that the application of GSSG inhibited the Kir6.1/SUR2B channel in a concentration-dependent manner (Fig. 4, A and B). Evident channel inhibition was seen with 0.5 mM GSSG, and stronger inhibition was produced by higher concentrations (Fig. 4, A and D). Together, these data strongly suggest that S-glutathionylation appears to occur in

![FIGURE 2. Responses of vascular K$_{ATP}$ channel to H$_2$O$_2$. A, whole-cell currents were recorded from a cell expressing Kir6.1/SUR2B channels. The bath solutions contained 145 mM K$^+$, making the reversal potential of the K$^+$ currents close to 0 mV. Command pulses of −80 mV were given every 3 s. The recording pipette was filled with the same solution (bath solution) with the addition of 1 mM ATP, 0.5 mM ADP, and 1 mM free Mg$^{2+}$. Application of Pin (10 μM) markedly augmented the whole-cell currents that were subsequently inhibited by 0.3 mM and 3 mM H$_2$O$_2$ progressively. The K$_{ATP}$ channel-specific blocker, Glib (10 μM) further reduced channel activity. The lower panel shows individual current traces produced by single command pulses. B, currents were recorded from giant inside-out patches obtained from HEK293 cells expressing the Kir6.1/SUR2B channel with a holding potential of −60 mV. Currents were activated by Pin, followed by increasing doses of H$_2$O$_2$. The H$_2$O$_2$ exposures did not lead to significant changes in channel activity. C, in the presence of GSH (100 μM), the H$_2$O$_2$ application strongly inhibited the Pin-activated currents, and the channels were almost completely inhibited by 100 μM H$_2$O$_2$ in giant inside-out patches. D, similar channel inhibition by H$_2$O$_2$ occurred in the presence of 2 mM GSH and 40 μM GSSG. Single channel activity from different treatments was displayed in expanded records from the top trace. Five active channels were seen, which were dose-dependently inhibited by H$_2$O$_2$. Note that the channel retained the same conductance with H$_2$O$_2$ exposure. Labels on the left: C, closure and 1, 2…5, the first, second…fifth opening. The locations from where these traces were expanded were marked with i–iv. E, dose-response relationships obtained in the whole-cell or inside-out patch configurations with or without GSH/GSSG. WC, whole-cell configuration; IO GSH, inside-out patch with 100 μM GSH; IO–, inside-out patch without GSH; IO GSH/GSSG, inside-out patch with a mixture of 2 mM GSH and 40 μM GSSG; F, the half-maximal inhibitory concentration (IC$_{50}$) of H$_2$O$_2$ was compared under whole-cell (WC) and inside-out patch (IO) configurations in the absence or presence of GSH or GSH/GSSG; G, the effect of 100 μM H$_2$O$_2$ on the channel activity showed significant differences (*, p < 0.05; ***,p < 0.001) between these conditions.
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Reversal of the \( K_{\text{ATP}} \) Channel Inhibition by Deglutathionylation Reagents—After pinacidil-activated currents were strongly inhibited by the oxidants, washout with addition of pinacidil (10 \( \mu \)M) had only a modest effect on channel activity (10.2 ± 2.6% of the original pinacidil-induced currents; \( n = 4 \); Fig. 2C). Increasing the pinacidil concentration to 100 \( \mu \)M had no additional effect (3.9 ± 0.6% of the original pinacidil-induced current; \( n = 3 \); Fig. 3A). However, application of the reducing agent dithiothreitol (DTT; 5 \( \mu \)M) reversed the GSSG-mediated (5 \( \mu \)M) current inhibition by 36.1 ± 4.7% (\( p < 0.001; n = 4 \); Fig. 4, C and E). Similarly, the 2-PDS mediated channel inhibition was partially reversed by DTT (34.9 ± 10.3% inhibition, \( n = 5 \); Fig. 5, B and C).

An exposure of the cell to the specific deglutathionylation reagent glutaredoxin-1 (Grx1; 4 \( \mu \)M) immediately reversed current inhibition by 27.8 ± 10.5% (\( p < 0.05; n = 4 \); Fig. 4D). After washout of both GSSG and Grx1, the pinacidil-induced currents were reduced to 72.0 ± 7.1% of its original level (\( p < 0.001; n = 4 \); Fig. 4, D and F). Collectively, these data suggest that S-glutathionylation-mediated Kir6.1/SUR2B channel inhibition can be reversed substantially by reducing or deglutathionylation reagents.

Biochemical Evidence for the Kir6.1 S-Glutathionylation—Does S-glutathionylation occur on the channel protein or on another protein that regulates the channel activity? To address this question, we examined S-glutathionylation using membrane-permeable BioGEE. HEK293 cells transfected with Kir6.1/SUR2B or the expression vector alone were loaded with BioGEE (250 \( \mu \)M) for 1 h followed by an \( H_2O_2 \) (750 \( \mu \)M) challenge for 15 min. Clear labeling was observed in the Kir6.1/SUR2B-transfected cells (Fig. 6, A–C), whereas a rather weak stain was seen with the mock transfection (data not shown). When the cells were double-stained with BioGEE (green) and Kir6.1 antibodies (red), co-localization was observed (Fig. 6, D–F).

The Kir6.1 S-glutathionylation was further tested with the streptavidin pull-down assay (26). The A10 vascular smooth muscle cell line, in which the Kir6.1/SUR2B channel was endogenously expressed (17, 27), was loaded with BioGEE (250 \( \mu \)M) for 1 h followed by an \( H_2O_2 \) (750 \( \mu \)M) challenge for 15 min. A strong Kir6.1-reactive band (~32 kDa) was detected in the whole-cell lysate (Fig. 6G, lower panel). After pull-down with streptavidin, the cell lysate pretreated with a combination of BioGEE and \( H_2O_2 \) showed a clear band of Kir6.1 immunoreactivity (Fig. 6G, upper panel), whereas no band was observed in cell lysates treated with either BioGEE or \( H_2O_2 \) alone (Fig. 6G).

Similar results were obtained using HEK293 cells transfected with Kir6.1/SUR2B (supplemental Fig. S1).

Oxidant-mediated Blockade of the \( K_{\text{ATP}} \) Channel Activation by Natural Activators—Kir6.1/SUR2B channel activity is low under basal condition but increases significantly in the presence of several vasodilating hormones and neurotransmitters that are coupled to the adenyl cyclase-cAMP-PKA pathway (3, 18). Therefore, we examined the effect of S-glutathionylation on the Kir6.1/SUR2B currents activated by the vasoactive intestinal polypeptide (VIP; 100 nM) and \( \beta \)-adrenoreceptor agonist isoproterenol (Isop; 100 nM), both of which activate Kir6.1/SUR2B currents through the PKA pathway. In the whole-cell configuration, DTNP (50 \( \mu \)M) or 2-DTP (50 \( \mu \)M)
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In inside-out patches, basal currents as well as the currents (currents were also potently inhibited by these agents (Fig. 7). 2-DTP inhibited the VIP-activated currents by 89.7%. In a giant inside-out patch, the Pin-induced currents were strongly inhibited by 5 mM GSSG. The current partially recovered with the application of the reducing agent DTT (5 mM). Pin-activated currents were potently inhibited by GSSG and partially reversed by 4 µM Grx1. Additional pinacidil perfusion (Pin WS) further augmented the channel activity. E and F, quantification of the data from C and D, respectively (n = 4). *, p < 0.05; ***, p < 0.001.

**FIGURE 4. The effect of GSSG on the channel activity and its reversibility.** A, in a cell expressing the Kir6.1/SUR2B channel, the Pin-activated currents were dose-dependently inhibited by GSSG treatment. B, summary of different dosages of GSSG on the channel activity (n = 4–6). C, in a giant inside-out patch, the Pin-induced currents were strongly inhibited by 5 mM GSSG. The current partially recovered with the application of the reducing agent, DTT (5 mM). D, Pin-induced currents were inhibited by GSSG and partially reversed by 4 µM Grx1. Additional pinacidil perfusion (Pin WS) further augmented the channel activity. E and F, quantification of the data from C and D, respectively (n = 4). *, p < 0.05; ***, p < 0.001.

**FIGURE 5. Thiol modification by reactive disulfide reagents.** A, in whole-cell configuration, the Pin-activated currents were potently inhibited by DTNP (50 µM). B, another reactive disulfide, 2-DTP (50 µM) also had a strong inhibitory effect on the Pin-sensitive currents of the Kir6.1/SUR2B channel, and this inhibitory effect was reversed by the reducing agent DTT (5 mM). C, the effects of 2-DTP (50 µM), DTNP (50 µM), and DTT (5 mM) on the channel activity were summarized. ***, p < 0.001.

**DISCUSSION**

Inflammatory oxidative stress is a common pathogenesis of cardiovascular diseases, including hypertension, atherosclerosis, and diabetic vascular complications, which mostly result from the overproduction of ROS overwhelming the capacity of cellular antioxidant defense systems (7, 8). When excessively produced, ROS can cause damages to lipids, proteins, and nucleotides, leading to cell dysfunction, structural injuries, and death (7). As the universal antioxidant treatment did not yield promising results in clinic trials (28), the identification of specific molecules and the understanding of the molecular mechanisms underlying the oxidant-mediated protein modulation becomes crucial for the development of novel therapeutic strategies. Our studies indicate that the Kir6.1/SUR2B channel is inhibited by micromolar concentrations of H$_2$O$_2$ through S-glutathionylation. Such channel inhibition is of pathophysiological relevance and is likely to occur in vasculatures as the production of micromolar concentrations of H$_2$O$_2$ has been shown during oxidative stress (10).

H$_2$O$_2$, O$_2^•$, OH, and peroxyl (RO$_2^•$) are the major oxidants produced endogenously in biological systems. Their concentrations in the cytoplasm are tightly controlled by several antioxidant systems (29). Early studies have shown that ROS including H$_2$O$_2$ have modulatory effects on membrane proteins (22, 30, 31). Most of the studies, however, use millimolar concentrations of H$_2$O$_2$ (32, 33). In the present study, we have observed a clear inhibition of the Kir6.1/SUR2B channel by H$_2$O$_2$ with IC$_{50}$ of ~1.5 mM in whole-cell recording. In contrast, the inhibition of $K_{ATP}$ currents can be clearly seen with micromolar concentrations of H$_2$O$_2$ in vascular rings. The major reason for the discrepancy is likely to be the effect of washout or inadequate controls of the cytosolic soluble factors that play a role in the channel modulation in oxidative stress. Indeed, some of the cytosolic substances are identified to be GSH and GSSG in the present study.

Searching for the missing cytosolic factors to better characterize the effect of H$_2$O$_2$, we conducted experiments in giant inside-out patches. We have found that supplying a small amount of exogenous GSH dramatically reduces the concentration of H$_2$O$_2$ needed for channel inhibition. Using 2 mM GSH and 40 µM GSSG, which mimic the intracellular GSH/GSSG ratio (21, 34), we have also observed clear channel inhibition by micromolar concentration of H$_2$O$_2$. The drastic effect of GSH/GSSG on the channel sensitivity to H$_2$O$_2$ is not only seen in inside-out patches, but also takes place in the whole-cell recordings. A supplement of GSH/GSSG to the pipette solution together, these results indicate that the Kir6.1/SUR2B channels are inhibited by S-glutathionylation, regardless of how the channels are activated.
The channel is inhibited by H\textsubscript{2}O\textsubscript{2} or diamide potently only when there is GSH. (ii) The S-glutathionylation inducer GSSG causes the channel inhibition. (iii) Several 2-PDSs reactive with thiol groups to form adapters at cysteine residues inhibit the Kir\textsubscript{6.1}/SUR\textsubscript{2B} channel in micromolar concentrations. (iv) The oxidant-mediated channel inhibition can be reversed by the specific deglutathionylation reagent Grx1 and the general reducing reagent DTT. (v) Streptavidin pull-down assays reveal the incorporation of the GSH moiety to the Kir6.1 subunit in the presence but not in the absence of H\textsubscript{2}O\textsubscript{2}. This \( K_{\text{ATP}} \) channel modulation mechanism is not limited to the pinacidil-induced currents. The \( K_{\text{ATP}} \) currents activated by the natural vasodilators VIP and Isop are similarly inhibited by these oxidants.

The modulation of protein activity by S-glutathionylation is a newly recognized post-translational regulatory mechanism (35, 36). This process, facilitated by oxidative stress and also seen in unstressed cells, can result in major changes to protein conformations and functions (16). Such modulation has been demonstrated in a large number of proteins using microarray (37) or proteomic analysis (38), while the physiological relevance of this modulation remains to be understood (36). Several recent studies indicate that the following membrane proteins are modulated by S-glutathionylation: the cystic fibrosis transmembrane conductance regulator (24), the ryanodine receptor (39), the sarco/endoplasmic reticulum Ca\textsuperscript{2+} ATPase (40), and the Na\textsuperscript{+}/K\textsuperscript{+} pump (41). The present study has shown for the first time that an important vascular tone regulator, Kir\textsubscript{6.1}/SUR\textsubscript{2B} channel, is subject to S-glutathionylation produced by H\textsubscript{2}O\textsubscript{2} at physiological or pathophysiological concentrations. This channel modulation by H\textsubscript{2}O\textsubscript{2} can lead to impairment of the vasodilation responses in a variety of vascular complications. Therefore, the demonstration of S-glutathionylation as a regulatory mechanism has important clinical implications. With the information, new therapeutic strategies may be formulated by preventing the oxidative modulation of the Kir6.1/SUR2B channel.

Another similar post-translational regulation mechanism is S-nitrosylation through which a nitric oxide (NO) moiety is incorporated into the thiol group of a cysteine residue (25, 42). However, our initial attempts using NO donors yielded inconsistent results (data not shown).

In conclusion, our studies indicate that the Kir6.1/SUR2B channel is inhibited by H\textsubscript{2}O\textsubscript{2} at micromolar concentrations...
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![Diagram](image)

**FIGURE 7. Redox-mediated blockade of the $K_{\text{ATP}}$ channel activation by natural activators.** A, in whole-cell configuration, Isop (100 nM) activated Kir6.1/SUR2B currents were subsequently inhibited by DTNP (50 μM). B, in a giant inside-out patch, the channel was activated by the catalytic subunit of PKA (cPKA, 50 units/ml), and inhibited by 5 mM GSSG. C, summary of the effects of GSSG, DTNP, and 2-PDS on the Kir6.1/SUR2B channel induced by a variety of channel activators ($n = 3–5$). Abbreviations: FSK, forskolin; VIP, vasoactive intestinal polypeptide. ***, $p < 0.001$.

owing to $S$-glutathionylation. The demonstration of the mechanism underlying the impairment of the vascular $K_{\text{ATP}}$ channel should have impacts on the treatment and prevention of several vascular diseases caused by oxidative stress.

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