Histone Deacetylase Inhibitors Relax Mouse Aorta Partly through Their Inhibitory Action on L-Type Ca^{2+} Channels[§]

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ABSTRACT

Histone deacetylase (HDAC) inhibitors modulate acetylation/deacetylation of histone and nonhistone proteins. They have been widely used for cancer treatment. However, there have been only a few studies investigating the effect of HDAC inhibitors on vascular tone regulation, most of which employed chronic treatment with HDAC inhibitors. In the present study, we found that two hydroxamate-based pan-HDAC inhibitors, suberoylanilide hydroxamic acid (SAHA) and trichostatin A (TSA), could partially but acutely relax high extracellular K^{+} -contracted mouse aortas. SAHA and TSA also attenuated the high extracellular K^{+} -induced cytosolic Ca^{2+} rise and inhibited L-type Ca^{2+} channel current in whole-cell patch-clamp. These data

demonstrate that SAHA could inhibit L-type Ca^{2+} channels to cause vascular relaxation. In addition, SAHA and TSA dose dependently relaxed the arteries precontracted with phenylephrine. The relaxant effect of SAHA and TSA was greater in phenylephrine-precontracted arteries than in high K^{+} -contracted arteries. Although part of the relaxant effect of SAHA and TSA on phenylephrine-precontracted arteries was related to L-type Ca^{2+} channels, both agents could also induce relaxation via a mechanism independent of L-type Ca^{2+} channels. Taken together, HDAC inhibitors SAHA and TSA can acutely relax blood vessels via their inhibitory action on L-type Ca^{2+} channels and via another L-type Ca^{2+} channel-independent mechanism.

Introduction

Histone deacetylases (HDACs) are a family of enzymes that regulate protein activity through removal of acetyl groups (Richardson et al., 2017). A total of 18 histone deacetylases can be grouped into four classes (I, II, III, and IV). The most well known molecular targets of HDACs are the histone proteins that compact DNA in chromatin. Reversible acetylation and deacetylation of histone proteins in chromatin constitutes a major epigenetic mechanism for the control of gene expression. In addition, up to 1750 nonhistone target proteins of HDACs have been identified (Choudhary et al., 2009). It has been suggested that acetylation and deacetylation of these nonhistone proteins may regulate such crucial cellular processes as the cell cycle, cytoskeleton reorganization, endocytosis, and vesicular trafficking (Choudhary et al., 2009). Owing to the

crucial importance of HDAC activity in physiologic and pathologic processes, HDAC inhibitors have been developed as therapeutic options for a variety of diseases, including cancer, mood disorders, and HIV (Dokmanovic et al., 2007). In an experimental model, HDAC inhibitors have also been shown to be beneficial in arrhythmia, myocardial infarction, hypertension, and atherosclerosis (Eom and Kook, 2014). HDAC inhibitors can be divided into several structural classes, including hydroxamates, cyclic peptides, and others (Dokmanovic et al., 2007). Suberoylanilide hydroxamic acid (SAHA) and trichostatin A (TSA) are hydroxamate-based pan-HDAC inhibitors that inhibit class I and class II HDAC proteins (Dokmanovic et al., 2007). However, their inhibitory potency to different classes of HDAC proteins varies greatly (Ko et al., 2013). Of the two, only SAHA has been approved by the FDA for clinical use, for treatment of cutaneous T-cell lymphoma and multiple myeloma (Dokmanovic et al., 2007).

Only a few studies have investigated the action of HDAC inhibitors in vascular tone control. Most examined the effect of chronic treatment with HDAC inhibitors on vascular tone regulation, but the results are controversial. Usui et al. (2012) showed that chronic treatment of rat mesenteric arteries with TSA for 3 weeks enhanced the endothelium-dependent

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ABBREVIATIONS: BAPTA, 1,2-bis(o-aminophenoxy)ethane-*N,N,N',N'*-tetraacetic acid; DMSO, dimethyl sulfoxide; HDAC, histone deacetylase; Orai1, calcium release-activated calcium channel protein 1; p66shc, Src homology 2 domain-containing transforming protein C1, MW 66 isoform; RO2959, 2,6-difluoro-*N*-{5-[4-methyl-1-(5-methyl-thiazol-2-yl)-1,2,5,6-tetrahydro-pyridin-3-yl]-pyrazin-2-yl}-benzamide hydrochloride; SAHA, suberoylanilide hydroxamic acid; SOCE, store-operated Ca^{2+} entry; TSA, trichostatin A; U46619, (Z)-7-[(1S,4R,5R,6S)-5-[(E,3S)-3-hydroxyoct-1-enyl]-3-oxabicyclo[2.2.1]heptan-6-yl]hept-5-enoic acid.

Fig. 1. SAHA and TSA reversed the vascular contraction elicited by high extracellular K^+ in mouse aortas. (A) Representative traces (left) and summarized data (right) showing verapamil-induced relaxation in the arteries precontracted by 60 mM KCl. The vehicle control (0.1% DMSO) had no effect. (B and C) The vascular relaxant effect of SAHA (B) and TSA (C) on 60 mM KCl-precontracted aortas with endothelium (+Endo) or without endothelium (-Endo). Representative traces (left) and data summary (right) of relaxing effect are shown. The data are expressed as mean $\pm$ S.E.M. ($n = 5$). *** $P < 0.001$ vs. vehicle (DMSO) control.

relaxation and reduced the angiotensin II (Ang II)-induced contraction in spontaneously hypertensive rats. In contrast, several other studies reported that chronic treatment with TSA reduced the endothelium-dependent and endothelium-independent vascular relaxation (Colussi et al., 2012; Mathews et al., 2014; Pandey et al., 2014). These authors found that, whereas the endothelium-dependent component could be attributed to TSA actions on the expressional level of arginase 2 and nitric oxide synthase (Mathews et al., 2014), the endothelium-independent components might be related to acetylation/deacetylation of such smooth muscle contractile proteins as myosin light chain and smooth muscle actin (Colussi et al., 2012). Interestingly, a recent study also demonstrated an acute effect of HDAC inhibitors on vascular tone. In that study, acute treatment of rat aortic rings with 10 μ M TSA for 15 minutes substantially reduced Ang II-induced contraction, probably via an inhibitory action of TSA on p66shc phosphorylation in vascular smooth muscle cells (Kang et al., 2015).

In the present study, we aimed to investigate acute effects of HDAC inhibitors on vascular tone. The results show that application of two commonly used HDAC inhibitors, SAHA and TSA, could induce an acute relaxation in high extracellular K^+ - and phenylephrine-precontracted rat aortic rings. This relaxation was endothelium-independent. Ca^{2+} influx measurement and whole-cell patch-clamp studies demonstrated that SAHA and TSA inhibit the activity of L-type Ca^{2+} channels in rat vascular smooth muscle cells.

Material and Methods

Chemicals. Verapamil, methylene blue, and phenylephrine were from Sigma-Aldrich (St. Louis, MO). SAHA (PubChem CID:5311), TSA (PubChem CID:444732), LBH-589 (PubChem CID:Panobinostat) (6918837), FK228 (romidepsin) (PubChem CID:5352062), and PDX-101 (belinostat) (PubChem CID:6918638) were from Selleckchem (Houston, TX). U46619, SAHA, TSA, LBH-589, FK228, and PDX-101 were dissolved in dimethyl sulfoxide (DMSO), and other drugs were prepared in distilled water.

Fig. 2. SAHA and TSA attenuated the high extracellular K^{+} -elicited cytosolic Ca^{2+} rise and inhibited L-type Ca^{2+} channel current. (A) Effect of SAHA and TSA on high K^{+} -induced Ca^{2+} rise. Acutely isolated smooth muscle cells were treated with SAHA/TSA in normal Krebs solution bath, followed by a change to 60 mM KCl Krebs solution bath, which induced a Ca^{2+} rise. SAHA/TSA at 300/100 μM was applied about 5 minutes before high K^{+} treatment. Representative traces (left) and data summary (right) of Ca^{2+} responses are shown. (B) Effect of SAHA and TSA on ionomycin-induced Ca^{2+} rise. Acutely isolated smooth muscle cells were treated with TSA/SAHA in normal Krebs solution bath, followed by Krebs solution bath with 1 μM ionomycin, which induced a Ca^{2+} rise. SAHA/TSA at 300/100 μM was applied about 5 minutes before ionomycin treatment. Representative traces (left) and data summary (right) of Ca^{2+} responses are shown. (C and D) Whole-cell current recording in A7r5 cells. Shown are current-voltage (I - V) relationships (left), voltage protocol and representative whole-cell current traces (right) of A7r5 cells with or without SAHA (C) and TSA (D). SAHA/TSA at 300/100 μM was applied about 5 minutes before current recording. The data are expressed as the mean $\pm$ S.E.M. [$n = 5$ in (A and B), $n = 6$ in (C and D)]. * $P < 0.05$; *** $P < 0.001$ vs. vehicle (DMSO) control.

Measurement of Vascular Tension. All animals were obtained from the Laboratory of Animal Services Center, the Chinese University of Hong Kong, Hong Kong, China. The animal experiments were conducted in accordance with the Guide for the Care and Use of Laboratory Animals published by the U.S. National Institutes of Health. Isometric tension of aortic rings were recorded by Multi Wire Myograph

System (Danish Myo Technology A/S, Aarhus, Denmark) as described (Tian et al., 2012). Briefly, male C57 adult mice were killed by carbon dioxide suffocation. After euthanasia, the mouse thoracic aortas were quickly dissected free and cut into 2-mm long rings in ice-cold and oxygenated Krebs buffer, which contained in mM: 119 NaCl, 4.7 KCl, 2.5 CaCl_2 , 1 MgCl_2 , 25 NaHCO_3 , 1.2 KH_2PO_4 , and 11 D-glucose.

Fig. 3. Comparison of inhibitory potencies of SAHA and TSA on high extracellular K^+ -induced Ca^{2+} entry with their vasodilatory potencies. (A) Comparison of dose-response curves and IC_{50} values of SAHA on 60 mM KCl-induced cytosolic Ca^{2+} rise in smooth muscle cells (left panel) with vasodilatory action of SAHA in K^+ -contracted mouse aortas (right panel). (B) Comparison of dose-response curves and IC_{50} values of TSA on 60 mM KCl-induced cytosolic Ca^{2+} rise in smooth muscle cells (left panel) with vasodilatory action of TSA in K^+ -contracted mouse aortas (right panel). The data are expressed as the mean $\pm$ S.E.M. ($n = 3-6$).

The vascular rings were mounted onto two thin stainless-steel holders in a myograph filled with Krebs solution bubbled with 95% O_2 and 5% CO_2 at 37°C. The optimum baseline tension was set at 3 mN. After an equilibration period and 60 mM KCl solution pretreatment (which contained in mM: 63.7 NaCl, 60 KCl, 2.5 $CaCl_2$, 1 $MgCl_2$, 25 $NaHCO_3$, 1.2 KH_2PO_4 , and 11 D-glucose), the rings were exposed to 1 μ M phenylephrine to test their contractile responses and subsequently challenged with acetylcholine to certify the integrity of endothelium.

Preparation of Acutely Isolated Smooth Muscle Cells. The smooth muscle cells were freshly dispersed as described previously (Eto et al., 2001). Briefly, male C57 adult mice were anesthetized with ether. The aorta was dissected. The connective tissue and the endothelium were carefully removed under microscope and the smooth muscle was cut into 1- to 2-mm sections. The tissues were preincubated with oxygenated Ca^{2+} -free phosphate-buffered saline (PBS) with collagenase II (2% w/v in 1 $\times$ PBS) and trypsin (0.25% w/v) for 40 minutes at 18–22°C, and the smooth muscle was cut into one to two arterial branches and dissected. The tissues were washed three times with oxygenated Ca^{2+} -free PBS and then gently agitated with a blunt-tipped glass pipette until the solution became cloudy. The debris was then removed with a fine 200- μ m nylon mesh. The cell suspension was stored at 4°C and used within 5 hours of cell harvesting.

Preparation of Smooth Muscle Cell Line. An aortic smooth muscle cell line from rat (A7r5) was obtained from the American Type Culture Collection (ATCC). The A7r5 cell line was cultured in 90% Dulbecco's modified Eagle's medium (Gibco/Thermo Fisher Scientific, Gaithersburg, MD), and 10% fetal bovine serum (Gibco/Thermo Fisher Scientific) at 37°C and 5% CO_2 in air.

Cytosolic Ca^{2+} Measurement. The cytosolic Ca^{2+} was measured as described elsewhere (Qi et al., 2015). Briefly, smooth muscle cells were loaded with Fluo-4/AM (Invitrogen/Thermo Fisher Scientific, Waltham, MA) in normal Tyrode's solution that contained in mM: 140 NaCl, 5.4 KCl, 1 $MgCl_2$, 2 $CaCl_2$, 5.5 glucose, and 5 HEPES, pH 7.4. To deplete intracellular Ca^{2+} stores, the cells were treated with 4 μ M thapsigargin in Ca^{2+} -free Tyrode's solution, which contained in mM: 140 NaCl, 5.4 KCl, 1 $MgCl_2$, 5.5 glucose, 0.2 ethylene glycol tetraacetic acid (EGTA), and 5 HEPES, pH 7.4. Store-operated Ca^{2+} entry (SOCE) was initiated by the addition of 2 mM Ca^{2+} to the bath. The real-time fluorescent images were captured every 10 seconds and analyzed by MetaFluor imaging software (Molecular Devices, Sunnyvale, CA).

Patch-Clamp Recording. For whole-cell patch-clamp recordings, single A7r5 smooth muscle cells were seeded on glass coverslips precoated for 30 minutes with Matrigel containing

50 μ g/ml laminin (Corning, Corning, NY) in 0.1% gelatin (Sigma-Aldrich). The glass coverslips containing the cells were placed into a room-temperature-recording chamber and perfused continuously with Tyrode's solution, which contained in mM: 140 NaCl, 1 $MgCl_2$, 1.8 $CaCl_2$, 5.4 KCl, 10 glucose, and 5 HEPES, pH adjusted to 7.4 by NaOH. The whole-cell patch-clamp recordings (voltage- and current-clamp configurations) were performed with an Axopatch 200B amplifier (Molecular Devices) and pCLAMP 10.4 software (Molecular Devices). Signals were digitized at 10 kHz and filtered at 2 kHz. Microelectrodes (1B150F-4; World Precision Instruments, Sarasota, FL) with resistances of 3–7 M Ω were pulled from P-97 puller (Sutter Instrument, Novato, CA). L-type Ca^{2+} current was recorded from single cells with ruptured whole-cell patch-clamp techniques. After establishing patch-clamp, the Tyrode's solution was replaced with the Ba^{2+} -containing solution, which contained in mM: 108 NaCl, 20 triethylamine (TEA)-Cl, 5.4 CsCl, 10 $BaCl_2$, 1 $MgCl_2$, 5.5 glucose, and 10 HEPES, pH adjusted to 7.4 by NaOH. The pipette solution contained in mM: 115 Cs-aspartate, 20 TEA-Cl, 1 $MgCl_2$, 5 BAPTA, 3 Mg -ATP, 0.2 GTP, and 10 HEPES, pH 7.4 adjusted with CsOH (Ochi et al., 2016). At a holding potential of -80 mV, current-voltage relations were obtained by applying 500-millisecond depolarization steps in 10-mV increments at 0.2 Hz from -40 to 50 mV. The test potential initially depolarized from -80 to -40 mV to detect and inactivate the T-type Ca^{2+} channel current.

Statistical Analysis. Results are expressed as means $\pm$ S.E.M. The relaxation was presented as percentage of the evoked contraction. Data were analyzed using GraphPad Prism software. Two-tailed Student's *t* test was used for two group comparison. One-way analysis of variance with Newman-Keuls post-hoc test were used for statistical evaluation when more than two groups were compared. $P < 0.05$ was taken as statistically significant.

Results

HDAC Inhibitors Reversed the Vascular Contraction Elicited by High Extracellular K^+ in Mouse Aortas.

Exposure of aortic segments to high extracellular K^+ concentration at 60 mM induced a rapid contraction of mouse aortas, which was completely reversed by an L-type Ca^{2+} channel inhibitor verapamil at 10 μ M (Fig. 1A). SAHA at 3–300 μ M or TSA at 1–100 μ M induced dose-dependent relaxation in the arteries that were precontracted with high extracellular K^+ (Fig. 1, B and C). Three hundred micromolar SAHA and

Fig. 4. HDAC inhibitors reversed the vascular contraction elicited by phenylephrine (Phe) in mouse aortas. Shown were vascular relaxant effect of SAHA (A) and TSA (B) on phenylephrine (1 μM)-precontracted aortas with endothelium (+Endo) or without endothelium (–Endo). Representative traces (left) and data summary (right) of relaxing effect are shown. The data are expressed as the mean $\pm$ S.E.M. ($n = 5$).

100 μM TSA resulted in a relaxation of $45\% \pm 2\%$ and $47\% \pm 3\%$, respectively (Fig. 1, B and C). The effect of SAHA and TSA was endothelium-independent (Fig. 1, B and C).

Several other HDAC inhibitors were also tested, including hydroxamates LBH-589 and PDX-101, as well as cyclic peptide FK-228. Like SAHA and TSA, all these HDAC inhibitors caused dose-dependent relaxation in high extracellular K^+ -contracted arteries (Supplemental Fig. 1).

TSA Attenuated the High Extracellular K^+ -Elicited Cytosolic Ca^{2+} Rise and Inhibited L-Type Ca^{2+} Channel Current. Cytosolic Ca^{2+} rise in response to high extracellular K^+ was measured in the freshly isolated mouse vascular smooth muscle cells. Sixty millimolar extracellular K^+ stimulated Ca^{2+} influx in these cells. Three hundred micromolar SAHA or 100 μM TSA markedly attenuated the Ca^{2+} influx elicited by high extracellular K^+ (Fig. 2A).

Literature shows that hydroxamates such as TSA and SAHA are able to chelate multiple ions, including Zn^{2+} , Ni^{2+} , and Fe^{3+} (Hebbel et al., 2010; Griffith et al., 2011). In our experiments, 300 μM SAHA or 100 μM TSA had no

significant effect on 1 μM ionomycin-induced cytosolic Ca^{2+} rise in vascular smooth muscle cells (Fig. 2B), suggesting that the ability of these agents in chelating/buffering cytosolic Ca^{2+} was minimal, if any, in our experimental conditions.

Under high extracellular K^+ , L-type Ca^{2+} channel current is the major source of Ca^{2+} , which initiates contraction of arterial smooth muscle cells and regulates blood pressure (Zhang et al., 2007). Whole-cell patch-clamp was used to measure L-type Ca^{2+} channel current. Because of technical difficulty in making giga-ohm seal on the plasma membrane of freshly isolated mouse smooth muscle cells, we opted to use A7r5 rat smooth muscle cells for patch-clamp studies. In whole-cell recording, the cells were held at -80 mV and voltage steps ranging from -40 to $+50$ mV were applied for 500 milliseconds in 10 mV step increments. A typical U-shaped voltage-gated Ca^{2+} current was recorded in these cells (Fig. 2, C and D) (Sun et al., 2000). Application of 300 μM SAHA or 100 μM TSA substantially inhibited the voltage-gated Ca^{2+} current in these cells (Fig. 2, C and D).

Fig. 5. Relative contribution of L-type Ca^{2+} channels vs. other components in the phenylephrine-precontracted arteries. The aortas were precontracted with 1 μM phenylephrine (Phe), followed by 10 μM verapamil. After the relaxant response to verapamil became stable, 200 μM SAHA (A) or 100 μM TSA (B) was added to induce further relaxation. Representative traces (left) and data summary (right) of relaxing effect are shown. The data are expressed as the mean $\pm$ S.E.M. ($n = 5$). *** $P < 0.001$ vs. vehicle (DMSO) control.

Comparison of Inhibitory Potencies of TSA and SAHA on High Extracellular K^{+} -Induced Ca^{2+} Entry with Their Vasodilatory Potencies. We examined the effect of different doses of SAHA (3–300 μM) and TSA (1–100 μM) on Ca^{2+} influx induced by high extracellular K^{+} . The results showed that SAHA and TSA caused dose-dependent inhibition on the high extracellular K^{+} -induced Ca^{2+} influx with IC_{50} of 70 and 21 μM , respectively (Fig. 3, A and B, left panels).

Dose-dependent vasodilatory curves of SAHA and TSA on high K^{+} -precontracted mouse aortas were also plotted (Fig. 3, A and B, right panels). The vasodilatory potencies of SAHA and TSA were similar to their inhibitory potencies on Ca^{2+} influx with comparable IC_{50} values (Fig. 3).

HDAC Inhibitors Reversed the Vascular Contraction Elicited by Phenylephrine in Mouse Aortas. SAHA at 3–300 μM or TSA at 1–100 μM also induced dose-dependent relaxation of mouse aortas precontracted with an α -adrenergic receptor agonist phenylephrine at 1 μM (Fig. 4, A and B). Compared with the high K^{+} -contracted arteries, the relaxant effect of SAHA and TSA was greater in phenylephrine-contracted arteries. Three hundred micromolar SAHA and 100 μM TSA almost completely reversed the phenylephrine-induced contraction (Fig. 4, A and B). Several other HDAC inhibitors, including PDX-101, LBH-589, and FK-228, also showed similar dose-dependent relaxation in phenylephrine-contracted arteries (Supplemental Fig. 1). In addition, the relaxant responses to SAHA and TSA were not significantly altered by endothelium denudation, suggesting that SAHA and TSA acted on vascular smooth muscle cells to exert their relaxant effect (Fig. 4, A and B).

Relative contribution of L-type Ca^{2+} channels versus other components in the phenylephrine-precontracted arteries was

estimated. Application of 10 μM verapamil caused 42–55% relaxation in these arteries (Fig. 5, A and B). Subsequent addition of 300 μM SAHA or 100 μM TSA resulted in further relaxation to nearly 100% (Fig. 5, A and B). These data suggest that L-type Ca^{2+} channels accounted for 42–55% of the contraction and the remaining 45–58% was independent of L-type Ca^{2+} channels.

SOCE Did Not Participate in the HDAC Inhibitor-Induced Vascular Relaxation, but HDAC Inhibitors Reduced the Phenylephrine-Induced Cytosolic Ca^{2+} Rise. The mechanisms of vascular contraction in response to vascular agonists are complicated, involving Ca^{2+} release from intracellular Ca^{2+} stores, Ca^{2+} entry through store-operated Ca^{2+} influx and/or receptor-operated Ca^{2+} channels and/or L-type Ca^{2+} channels, as well as a sensitized contractile response to Ca^{2+} (Wang et al., 2008). We next explored whether HDAC inhibitors could act on other mechanism(s) independent of L-type Ca^{2+} channels to induce vascular relaxation.

Calcium release-activated calcium channel protein 1 (Orai1) is the pore-forming subunit of SOCE in vascular smooth muscle cells (Lewis, 2011). As expected, the selective Orai1 inhibitor RO2959 at 200 nM (Chen et al., 2013) greatly inhibited the SOCE in the acutely isolated mouse smooth muscle cells (Fig. 6A). However, RO2959 up to 5 μM failed to induce relaxation in these arteries (Fig. 6B).

In the absence of extracellular Ca^{2+} , application of 1 μM phenylephrine failed to elicit a cytosolic Ca^{2+} rise (Fig. 7A), suggesting no significant stimulating effect of phenylephrine on intracellular Ca^{2+} release. Subsequent Ca^{2+} addback (2 mM) to the bath solution resulted in a cytosolic Ca^{2+} rise, which was more marked in phenylephrine-treated cells than in DMSO-treated vehicle control (Fig. 7A, left panel),

Fig. 6. SOCE did not contribute to the HDAC inhibitor-induced vascular relaxation. (A) Effect of RO2959 on SOCE in the freshly isolated mouse smooth muscle cells. The cells were treated with 4 μM thapsigargin (TG) in a Ca^{2+} -free bath, which induced the first Ca^{2+} rise, followed by adding back 2 mM Ca^{2+} to the bath, which induced the second Ca^{2+} rise. The second Ca^{2+} rise was taken as SOCE. RO2959 at 200 nM was applied 5 minutes before TG. The representative traces (left) and data summary (right) of Ca^{2+} responses are shown. The summary data are expressed as the mean $\pm$ S.E.M. ($n = 5$ experiments, >25 cells per experiment). *** $P < 0.001$. (B) A representative trace showing lack of acute relaxant effect of RO2959 on 1 μM phenylephrine (Phe)-precontracted aortas. A representative time-control experiment (without RO2959) is displayed on top. $n = 4$ experiments.

suggesting that phenylephrine elicited extracellular Ca^{2+} entry. As there was no obvious intracellular Ca^{2+} release in response to phenylephrine (Fig. 7A), this Ca^{2+} entry may reflect receptor-operated Ca^{2+} entry rather than SOCE. Importantly, in the presence of 100 μM TSA or 300 μM SAHA, the phenylephrine-elicited extracellular Ca^{2+} entry was abolished (Fig. 7A).

Dose-dependent inhibition of SAHA (3–300 μM) and TSA (1–100 μM) on phenylephrine-induced Ca^{2+} influx was determined (left panels in Fig. 7, B and C). Dose-dependent vasodilatory curves of SAHA and TSA on phenylephrine-precontracted mouse aortas were also plotted (right panels in Fig. 7, B and C). The inhibitory potencies of SAHA and TSA on Ca^{2+} influx were similar to their vasodilatory potencies with comparable IC_{50} values (Fig. 7, B and C).

Lack of Nitric Oxide Involvement in TSA and SAHA-Induced Vascular Relaxation. Methylene blue is a guanylate cyclase inhibitor commonly used to abolish nitric oxide-mediated vascular dilation (Gruetter et al., 1981). Preincubation with 10 μM methylene blue for 15 minutes had no effect on SAHA-induced vascular relaxation in mouse aortas contracted with high extracellular K^{+} (Fig. 8A) or phenylephrine (Fig. 8B), suggesting lack of nitric oxide involvement in SAHA-induced vascular relaxation.

Discussion

The main findings of the present study were as follows: 1) Two commonly used HDAC inhibitors, SAHA and TSA, partially reversed the high extracellular K^{+} -induced contraction in mouse

aortas. The effect of SAHA and TSA was acute, dose-dependent, but endothelium-independent. 2) SAHA and TSA also caused dose-dependent relaxation in phenylephrine-precontracted mouse aortas. Three hundred micromolar SAHA and 100 μM TSA completely reversed the phenylephrine-induced contraction. Again, the effect of SAHA and TSA was acute and endothelium-independent. 3) In the freshly isolated mouse smooth muscle cells, SAHA and TSA markedly reduced the high K^{+} -induced cytosolic Ca^{2+} rise. Whole-cell patch-clamp studies demonstrated that SAHA and TSA acutely inhibited the voltage-gated L-type Ca^{2+} channel current in these cells. 4) SAHA and TSA also markedly reduced the phenylephrine-induced Ca^{2+} entry with inhibitory potency like their vasodilatory effect. 5) A potent Orai1 inhibitor, RO2959, failed to induce vascular relaxation in phenylephrine-induced contraction of mouse aortas. Taken together, the present study demonstrated that hydroxamate-based pan-HDAC inhibitors SAHA and TSA at intermediate micromolar concentrations could acutely inhibit L-type Ca^{2+} channels and consequently reduce the high extracellular K^{+} -induced contraction of aortas. SAHA and TSA also relaxed the phenylephrine-precontracted aortas, a minor part of which could be attributed to the inhibitory effect of these compounds on L-type Ca^{2+} channels.

It is well documented that exposure to high extracellular K^{+} at 60–100 mM causes vascular contraction (Eto et al., 2001). The underlying mechanism involves high K^{+} -induced membrane depolarization of vascular smooth muscle cells, which triggers an opening of voltage-gated Ca^{2+} channels, especially L-type Ca^{2+} channels. Subsequent Ca^{2+} influx through L-type Ca^{2+} channels elevates cytosolic Ca^{2+} level in vascular

Fig. 7. Action of HDAC inhibitors on phenylephrine-induced cytosolic Ca^{2+} rise and comparison with their vasodilatory effect. (A) The freshly isolated mouse vascular smooth muscle cells were treated with 1 μM phenylephrine (Phe) in a Ca^{2+} -free bath, followed by adding back 2 mM Ca^{2+} to the bath, which induced a Ca^{2+} entry. SAHA/TSA at 100 μM was applied 2 minutes before Phe. The representative traces (left) and data summary (right) of Ca^{2+} responses are shown. (B) Comparison of dose-response curves and IC_{50} values of SAHA on Phe-induced cytosolic Ca^{2+} rise in smooth muscle cells (left panel) with vasodilatory action of SAHA in Phe-contracted mouse aortas (right panel). (C) Comparison of dose-response curves and IC_{50} values of TSA on Phe-induced cytosolic Ca^{2+} rise in smooth muscle cells (left panel) with vasodilatory action of TSA in Phe-contracted mouse aortas (right panel). The summary data are expressed as the mean $\pm$ S.E.M. [$n = 5$ experiments, >25 cells per experiment in (A); $n = 3-5$ in (B and C)]. $^{**}P < 0.01$; $^{***}P < 0.001$.

smooth muscle cells, resulting in vascular contraction (Eto et al., 2001). In the present study, high extracellular K^{+} -induced aortic contraction was completely reversed by 10 μM verapamil, confirming the involvement of L-type Ca^{2+} channels. Importantly, SAHA and TSA partially but acutely reversed the high extracellular K^{+} -induced contraction of mouse aortas, suggesting a possible inhibitory action of these agents on voltage-gated Ca^{2+} channels. Two sets of experiments, cytosolic Ca^{2+} measurement and whole-cell patch-clamp, were performed to verify the action of SAHA and TSA on L-type Ca^{2+} channels. In cytosolic Ca^{2+} measurement, high extracellular K^{+} -evoked cytosolic Ca^{2+} rise was reduced by SAHA and TSA. Whole-cell patch-clamp experiments provide the direct evidence for inhibitory action of SAHA and TSA on L-type Ca^{2+} channel current in vascular smooth muscle cells. Interestingly, inhibitory potencies (IC_{50}) of SAHA and

TSA on high extracellular K^{+} -induced Ca^{2+} rise were similar to the vasodilatory potencies of these compounds in high K^{+} -contracted arteries. Taken together, these results provided unequivocal evidence for inhibitory action of HDAC inhibitors on the activity of L-type Ca^{2+} channels in vascular smooth muscle cells.

Our finding about the modulatory effect of HDAC inhibitors on voltage-gated Ca^{2+} channels has important physiologic and pathologic implication. SAHA and several other HDAC inhibitors are in clinical use for cancer treatment (Zwergel et al., 2016). In animal models, HDAC inhibitors also show beneficial cardiovascular effect by alleviating cardiac hypertrophy, hypertension, and myocardial infarction (Eom and Kook, 2014). In the present study, we show for the first time that HDAC inhibitors can acutely inhibit L-type Ca^{2+} channels and subsequently reduce the vascular tone. It is well

Fig. 8. Nitric oxide did not contribute to HDAC inhibitor-induced vascular relaxation. (A) The aortas were precontracted with 60 mM KCl in the absence (upper left panel) or presence (lower left panel) of 10 μM methylene blue (MB). After the contractile response became stable (~ 10 minutes), SAHA was added to induce relaxation. (B) The aortas were precontracted with 1 μM phenylephrine in the absence (upper left panel) or presence (lower left panel) of 10 μM methylene blue (MB). After the contractile response became stable (~ 10 minutes), SAHA was added to induce relaxation. Shown are representative traces (left) and data summary (right). The data are expressed as the mean $\pm$ S.E.M. ($n = 5$).

documented that L-type Ca^{2+} channels are closely associated with vascular tone and that an aggravated activity of L-type Ca^{2+} channels is a causative factor for pathologic progression of hypertrophy, hypertension, and myocardial infarction (Nayler, 1993). Therefore, our present results about the effect of HDAC inhibitors on L-type Ca^{2+} channels may provide a mechanistic explanation for cardio-protective effect of HDAC inhibitors against hypertrophy, hypertension, and myocardial infarction.

The mechanisms of vascular contraction in response to vascular agonists such as phenylephrine are complicated. These may include Ca^{2+} release from intracellular Ca^{2+} stores, Ca^{2+} entry through store-operated Ca^{2+} influx and/or receptor-operated Ca^{2+} channels and/or L-type Ca^{2+} channels, as well as a sensitized contractile response to Ca^{2+} (Nishimura et al., 1989;

Sun et al., 2000). In our study, SAHA and TSA could completely reverse the phenylephrine-induced contraction in mouse aorta. As expected, part of the phenylephrine-induced contraction, which amounts to 42–55%, was mediated by L-type Ca^{2+} channels (Nishimura et al., 1989), as indicated by its sensitivity to verapamil inhibition. However, in the presence of verapamil, SAHA and TSA could cause additional relaxation, which presumably resulted from other mechanisms independent of L-type Ca^{2+} channels. In our studies, phenylephrine failed to elicit significant intracellular Ca^{2+} release (Fig. 7A). Furthermore, inhibition of SOCE by an Orai1 inhibitor RO2959 failed to reverse the phenylephrine-induced contraction (Fig. 6B). These data suggest that the vascular relaxant effect of SAHA and TSA was independent of intracellular Ca^{2+} release or SOCE. Nevertheless, SAHA and TSA could inhibit phenylephrine-elicited

Ca²⁺ entry. Furthermore, inhibitory potencies (IC₅₀) of SAHA and TSA on phenylephrine-induced Ca²⁺ entry were similar to the vasodilatory potencies of these compounds in phenylephrine-contracted arteries, suggesting an important role of cytosolic Ca²⁺ in these events. Together, these data suggest a possible action of HDAC inhibitors on some Ca²⁺-signaling mechanisms independent of SOCE and intracellular Ca²⁺ release. These may include receptor-operated Ca²⁺ entry, Ca²⁺ sensitivity of contractile mechanism, or other alternative mechanisms.

One concern is that under oxidative stress, hydroxamate compounds, including SAHA and TSA, are able to produce nitric oxide (Samuni et al., 2009), which could account for the vasodilatory action of these compounds. However, all our experiments, including vascular tone studies and Ca²⁺ entry and patch-clamp experiments, were performed under normal physiologic saline without oxidative stress. Indeed, pretreatment of arteries with methylene blue, a guanylate cyclase inhibitor, had no effect on SAHA-induced vascular dilation, suggesting that the vasodilatory response to hydroxamate compounds in our experiments was not the result of hydroxamate-generated nitric oxide.

SAHA and TSA are pan-HDAC inhibitors that inhibit class I and class II HDAC proteins. However, their inhibitory potency to different classes of HDAC proteins varies greatly (Dokmanovic et al., 2007; Ko et al., 2013; Lobera et al., 2013). The reported IC₅₀ values of SAHA and TSA for class I and class IIb HDAC proteins range from nanomolar to sub-micromolar, whereas the IC₅₀ values for class IIa HDAC proteins range from low to intermediate micromolar (Dokmanovic et al., 2007; Ko et al., 2013; Lobera et al., 2013). In our studies, SAHA and TSA relaxed the extracellular high K⁺- and phenylephrine-precontracted aortas at intermediate micromolar concentration, which could inhibit class IIa HDAC proteins. However, considering the issue of drug permeation in cultured cells and vascular tissue, it is still uncertain which specific class of HDAC proteins might be involved.

In conclusion, we have demonstrated that HDAC inhibitors can cause acute inhibition of the activity of L-type Ca²⁺ channels, resulting in partial reversal of high extracellular K⁺-induced vascular contraction. HDAC inhibitors also acutely and potently reverse the phenylephrine-induced vascular contraction via L-type Ca²⁺ channel-dependent and -independent mechanisms.

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Authorship Contributions

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