Testosterone regulates the expression and functional activity of sphingosine-1-phosphate receptors in the rat corpus cavernosum

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Abstract

The bioactive lipid sphingosine-1-phosphate (S1P) regulates smooth muscle (SM) contractility predominantly via three G protein-coupled receptors. The S1P1 receptor is associated with nitric oxide (NO)-mediated SM relaxation, while S1P2 & S1P3 receptors are linked to SM contraction via activation of the Rho-kinase pathway. This study is to determine testosterone (T) modulating the expression and functional activity of S1P receptors in corpus cavernosum (CC). Adult male Sprague-Dawley rats were randomly divided into three groups: sham-operated controls, surgical castration and T supplemented group. Serum S1P levels were detected by high-performance liquid chromatography. The expression of S1P1-3 receptors and sphingosine kinases was detected by real-time RT-PCR. In vitro organ bath contractility and in vivo intracavernous pressure (ICP) measurement were also performed. T deprivation significantly decreased ICP rise. Meanwhile, surgical castration induced a significant increase in serum S1P level and the expression of S1P2-3 receptors by twofold ($P < 0.05$) but a decrease in the expression of S1P1 receptor. Castration also augmented exogenous phenylephrine (PE), S1P, S1P1,3 receptor agonist FTY720-P contractility and S1P2-specific antagonist JTE013 relaxation effect. T supplemented could restore the aforementioned changes. We provide novel data that castration increased serum S1P concentration and up-regulated the expression of S1P2-3 receptors in CC. Consistently, agonizing S1P receptors induced CCSM contraction and antagonizing mediated relaxation were augmented. This provides the first clear evidence that S1P system dysregulation may contribute to hypogonadism-related erectile dysfunction (ED), and S1P receptors may be expected as a potential target for treating ED.

Keywords: sphingosine-1-phosphate ● testosterone ● corpus cavernosum ● smooth muscle ● erectile dysfunction

Introduction

Penile erection is a neurovascular event involving the relaxation of corpus cavernosum smooth muscle (CCSM), which is maintained tonically contracted state in majority of the time [1–3]. Although many studies have investigated CCSM relaxation pathways, few have focused on the role of the contractile apparatus in erectile function (EF). Similarly, the major therapeutic treatments available for ED have primarily targeted CCSM relaxation pathways with particular emphasis on NO/cyclic guanosine monophosphate (cGMP) signalling [3–5].

Shingosine-1-phosphate (S1P), a member of a large family of lipid metabolites termed sphingolipids, represents one of the key latest additions to the list of ‘vasoactive’ substances that modulate vascular tone [6]. S1P is produced by sphingosine kinases (SphK) catalyzing the ATP-dependent phosphorylation of sphingosine. Thus far, two different SphK isoforms have been found in mammals named SphK1 and SphK2 [7]. SphK1 is predominantly cytosolic and pro-survival, while SphK2 functions mainly in the endoplasmic reticulum and stimulates apoptosis but its role is still poorly understood [6–8]. In plasma, S1P can reach a concentration of 0.1–4 μM [9]. Besides vasoactive potential, S1P is capable of regulating a wide array of biological processes such as cell proliferation, migration, survival, differentiation and others [6, 8, 10]. Many of these cellular responses are initiated by S1P binding to and
activating a set of five G protein-coupled S1P receptors (S1P1-5) [11]. In mammals, S1P1, S1P2 and S1P3 are found in all tissues, whereas S1P4 is restricted to lymphoid tissues and lung [12], and S1P5 to brain and skin [13]. In blood vessels, vascular endothelial cells (ECs) and SM cells express specific receptors for S1P that modulates vascular tone [14]. In particular, vasorelaxation is elicited by S1P through S1P1 receptors in ECs via endothelial nitric oxide synthase (eNOS) pathways [6, 8, 15], whereas S1P2 and S1P3 receptors in SM can elicit vasoconstriction responses through the activation of RhoA/ROK pathways [6, 8, 16, 17]. For a given blood vessel preparation, whether S1P stimulation causes vasodilatation or vasoconstriction may depend on multiple experimental variables, such as the animal species, the vascular bed, the S1P concentrations, S1P receptor subtype expression profile, as well as disease states. In general, higher concentration of S1P induces vasoconstriction in resistance vessels such as mesenteric, cerebral and coronary arteries but has little or no effect on conduit vessels such as aorta, carotid and femoral arteries [6, 8]. Apart from resistant arteries, S1P can contract SM of urinary bladder, uterus, gastrointestinal tract and bronchial tube [16, 18]. With regard to CCSM, di Villa Bianca et al. [19] found exogenous S1P did not directly relax or contract of human CCSM strips but potentiated acetylcholine response. The tissues used in their study were obtained from male to female transsexual patients with androandrogen and oestrogen pre-surgery treatment. Thus, we do not know S1P on intact CCSM. In fact, we recently reported that S1P caused strong in vitro and in vivo contraction of CCSM of normal rats and patients with various kinds of ED, while S1P receptor antagonist induced CCSM relaxation [20].

Androgens play a dual role in the erectile process by controlling both pro-erectile and anti-erectile signalling pathways [21, 22]. Indeed, testosterone (T) modulates NO and its generating enzyme NOS, which induce penile erection [23–25]. Recently, our serial studies together with other institutions have confirmed that T also regulates phosphodiesterase-5 (PDE5) [4, 5, 22, 26–28] which is mainly responsible for penile detumescence. In addition, castration up-regulates CCSM contractility process, including hyper-responsiveness to α-adrenergic agonist, increased SM myosin (SMM) phosphorylation and alteration of SMM isoform composition, as well as activation of RhoA/ROK signalling [3, 29, 30].

However, the CCSM, unlike most other SM, spends the majority of its time in the contracted state and relaxes only upon receiving erectogenic stimuli. A better understanding of androgens that regulate contraction in the CCSM is as critical as their modulation on CCSM

Fig. 1 Effect of testosterone (T) on the weight of androgen-sensitive organs. Left panel is bar graph of the average weight of the ventral prostates from all experimental groups. And right panel is bar graph of the average weight of seminal vesicle from all experimental groups. Values are expressed as means ± S.E. **P < 0.01 versus sham or castration + T. (n = number of samples from different animals).

Fig. 2 Effect of testosterone (T) on rat serum S1P levels. A: Serum S1P levels were detected by high-performance liquid chromatography (HPLC). The green curve is serum sample, while the blue one is serum sample mixed with 2.6 ng exogenous S1P. B: bar graph of the average serum S1P from all experimental groups. Values are expressed as means ± S.E. *P < 0.05 versus sham or castration + T. (n = number of samples from different animals).
relaxation mechanism. The goal of this study was to investigate T mod-
ulating the expression and functional activity of S1P receptors in CCSM.

Materials and methods

Animals and tissues

Adult male Sprague–Dawley rats weighing 300–350 g were used in this
study. All the rats were divided into three groups: sham, castration and
castration with T (testosterone propionate, Sigma-Aldrich; St. Louis,
MO, USA). As described in previous study [31], rats that underwent just
a perineal incision served as sham. Castration was performed through
the bilateral orchectomy under sodium pentobarbital anaesthesia. Part
of the castrated rats was injected subcutaneously with 30 mg/kg T per
week, and others were treated with vehicle (sesame oil) only. Rats were
killed 2 weeks post-surgery, blood was drawn from the heart for S1P
measurements and CC was collected for organ bath physiology studies
(placed in Krebs–Henseleit solution) or subsequent molecular analyses
(frozen in liquid nitrogen). The prostate and seminal vesicles were har-
vested and weighted after kill. All animal studies were approved by the
research committee of Zhongnan Hospital of Wuhan University.

High-performance liquid chromatography

S1P serum levels were determined with high-performance liquid chro-
matography (HPLC) as previously described with minor modification
[17]. Briefly, serum samples were extracted into chloroform/methanol,
and the obtained lipid samples were vacuum dried. The sphingolipids
were then derivatized with 9-fluorenylmethyl chloroformiate (FMOC-Cl),
and chromatographic detection of sphingolipids was performed with
reversed-phase chromatography on a 300 x 3.9 mm Delta-Pak C18
column (Waters Corporation, Milford, MA, USA).

Total RNA extraction and real-time reverse tran
criptase polymerase chain reaction
(real-time RT-PCR)

As previously reported [17, 32], total RNA was isolated from the frozen
tissues using TRizol reagent (Invitrogen, Carlsbad, CA, USA) according
to the manufacturer’s protocol. RNA concentration and purity were
determined using a ND-1000 Nanodrop spectrophotometer (Nanodrop
Technologies; Wilmington, DE, USA). For each sample, 1 μg of RNA
was converted to complementary DNA (cDNA) using reverse transcrip-
tase via the SuperScript II First-Strand Synthesis System (Invitrogen)

Fig. 3 Expression of S1P receptors and sphingosine kinases (SphKs) in rat corpus cavernosum from all experimental groups. The expression of all
studied molecules was quantified by real-time RT-PCR and normalized to expression of the ribosomal protein L19 (RPL19) housekeeping gene. A:
S1P1-3 receptors mRNA level, B: SphK1-2 mRNA level. Values are expressed as means ± S.E. *P < 0.05 versus sham or castration + T. (n = 8
different rats for each group).

Fig. 4 Effect of testosterone (T) on electrostimulation (ES)-induced erec-
tile response of rats. Penile erection was elicited by ES (width 5 ms,
duration 30 sec., 2.5 V) of the rat cavernous nerve at varying stimulation
frequencies (1–16 Hz). Erectile function was quantified by calculating
maximal intracavernous pressure (ICP)/mean arterial pressure
(MAP) ratio (x100). Values are expressed as means ± S.E. *P < 0.05
versus sham or castration + T. **P < 0.01 versus sham or castra-
tion + T. (n = 5 different rats for each group).
according to the manufacturer’s protocol. Primer pairs were designed using published cDNA sequences obtained via Entrez Nucleotide of the National Center for Biotechnology Information (NCBI) and the Primer Express program (Applied Biosystems, Foster City, CA, USA). Amplicons were deliberately kept at between 50 and 100 base pairs (bp) for all primer pairs to enable equal transcriptional efficiency.

RT products then were amplified in a 96-well plate in a 25 μl reaction volume with all samples run in triplicate, using the Model 7300 real-time thermocycler (Applied Biosystems). The following experimental protocol was utilized: denaturation (95°C for 10 min. to activate the polymerase) followed by an amplification program repeated for 40 cycles (95°C for 15 sec., then 60°C for 60 sec.) using a single fluorescence measurement. The following targets were amplified using SYBR Green for amplicon detection: S1P receptors 1–3 and SphK1-2. For relative quantification, the efficiency of amplification for each individual primer pair was determined using cDNA target and the 2^{ΔΔCt} method [33] in conjunction with the RQ Study Software version 1.2.3 (Applied Biosystems). Gene expression was normalized to expression of the RPL19 housekeeping gene.

**In vitro organ bath studies**

As previously reported [31, 34], the rat CC was mounted longitudinally in a 4 ml organ bath—Multi Myograph Model 800MS (Danish Myo Technology, Aarhus, Denmark) by securing to the two pins. One of the pins was attached to a force transducer which was calibrated to mg of force prior to the start of experimentation. The myograph was connected in line to a PowerLab 4/30 data acquisition system (ADInstruments; Colorado Springs, CO, USA) and in turn to a dual-core processor Pentium computer for real-time monitoring of physiological force.

The SM strips were equilibrated for at least 1 hr in Krebs buffer maintained at a mean temperature of 37 ± 0.05°C with continuous bubbling of 95% O₂ and 5% CO₂ with buffer changes every 15 min. The buffer had the following mM composition: NaCl 110, KCl 4.8, CaCl₂ 2.5, MgSO₄ 1.2, KH₂PO₄ 1.2, NaHCO₃ 25 and dextrose 11. The strips were continuously adjusted to a 400 mg resting tension and isometric tension recorded. After equilibration, the tissues were contracted with

![Fig. 5 Phenylophrine (PE)-induced contraction of rat corpus cavernosum smooth muscle (CCSM) in vitro. Panel A, B and C are PE-induced dose–response force tracings of rats CCSM from sham (A), castration (B) and castration + T (C) groups, respectively. For typical tracings, the x-axis represents time (min.), while the y-axis represents force (mg). Panel D is summary graph for curves of PE-induced dose–response contraction that resulted in sigmoid (S-shaped) curves after logarithmic transformation. Maximal response to KCl was taken as 100%, while the contractility of PE was evaluated as a percentage of this response. Values are expressed as means ± S.E.. (n = CC strips obtained from 6 to 7 different rats for each group).](image-url)
60 mM KCl. This degree of contractile response was taken as 100% and the force induced by different concentrations of the various agonists including PE, S1P (Cayman Chemical, Ann Arbor, MI, USA) and FTY720-P (Echelon Biosciences Inc, Salt Lake City, UT, USA), was expressed as a percentage of this value. Meanwhile, strips were precontracted with 1 μM PE (a dose that was determined to induce remarkable contraction) and allowed to reach a stable tension, and then, the relaxant effects of increasing doses (1–10 μM) of S1P2 receptor-selective antagonist JTE-013 (Tocris, Ellisville, MO, USA) were evaluated.

**In vivo studies**

As previously reported [4, 5], part of rats from each group was anaesthetized with pentobarbital (35 mg/kg) via an intraperitoneal (i.p) injection. Mean arterial pressure (MAP) via carotid artery and ICP were continuously monitored, using methods described previously [35]. Briefly, an incision was made in the perineum, and a window was made in the ischiocavernous muscle to expose both CC. The right crura were perforated with a 28-gauge needle connected to PE-50 tubing for ICP recording, while the left crura were perforated with a 30-gauge needle connected to PE-10 tubing for drug delivery. The cavernous nerve was identified ventrolateral to the prostate gland and carefully isolated. Direct electrostimulation (ES) of the cavernous nerve was performed with a delicate stainless steel bipolar hook electrode attached to the multijointed clamp. Each probe was 0.2 mm in diameter with a 1 mm separation between the two poles. Monophasic rectangular pulses were delivered by a signal generator (custom-made and with built-in constant current amplifier). MAP and ICP were recorded via pressure transducers connected in line to the PowerLab 4/30 data acquisition system connected in turn to computer for real-time analysis.
monitoring of pressure changes. Pressure transducers were calibrated to water prior to each experiment. After a stable baseline ICP was obtained, ICP rise induced by ES (width 5 ms, duration 30 sec., 2.5 V) at varying frequencies (1, 2, 4, 8, 16 Hz) was recorded. For another group of rats, intracavernous injection (ICI) of 50 l ethanol was carried out to test vehicle effects. Then, increasing doses (50, 250, 500 nmols) of JTE-013 alone were ICI with 30-min. intervals between with washout (saline flush). The ICP rise was quantified by calculating the ratio of maximum ICP/MAP.

Results

After a 2-week surgical castration, obvious atrophy of androgen-sensitive organs was observed in castrated rats. As shown in Figure 1, the weight of both prostate and seminal vesicles decreased by approximately eightfold and 10-fold in castrated rats and the mass reinstated completely after T supplementation.

Serum S1P concentrations were detected with HPLC. As displayed in Figure 2A, serum S1P peak appeared at 38 min. (green curve) which was confirmed by a synergic peak occurred exactly at the same time-point when sample mixed with exogenous S1P (blue curve). As shown in Figure 2B, T deprivation significantly increased serum S1P to 1.5 ng/ul, which was 0.23 ng/ul higher than that of sham. And the change was reversed with T injection.

To determine whether T affected the expression of S1P receptors in CC, the mRNA levels of S1P1-3 receptors in CC were detected. With real-time RT-PCR, we found castration significantly increased both S1P2 and S1P3 transcripts in CC (Fig. 3A), which were mainly related to constriction of SM. In contrast, S1P1 mRNA was found to be significantly decreased in CC from castrated rats (Fig. 3A), which was contributed to vasorelaxation. Additionally, critical enzymes mediating the biosynthesis of S1P in CC were examined. As shown in Figure 3B, lower SphK1 and higher SphK2 mRNA levels in CC were found in castrated rats as compared with controls. Altered expression

Fig. 7 S1P analogue (FTY720-P)-induced contractility of rat corpus cavernosum smooth muscle (CCSM) in vitro. Panel A, B and C are FTY720-P-induced dose–response force tracings of rats CCSM from sham (A), castration (B) and castration + T (C) groups, respectively. For typical tracings, the x-axis represents time (min.), while the y-axis represents force (mg). Panel D is a summary graph for the data shown in panel A-C. Maximal response to KCl was taken as 100%, while the contractility of FTY720-P was evaluated as a percentage of this response. Values are expressed as means ± S.E. **P < 0.01 versus sham or castration + T. (n = CC strips obtained from 7 to 8 different rats for each group).
of all these molecules was maintained or recovered to control levels when T was given back.

In line with our previous study [5], castrated rats were found to exhibit significant ED, with the ICP rise elicited by ES of the cavernous nerve significantly attenuated in the castration group at all stimulation frequencies when compared to controls (Fig. 4). The impaired EF can be partially attributed to increased CCSM tone mediated by vasoconstrictors. In this study, major penile neurotransmitter PE and new ‘vasoactive’ substance S1P were investigated. Consistently, cavernosal strips from castrated animals displayed heighted contractile responses to cumulative doses of PE, which was demonstrated in Figure 5 including typical force tracings (Fig. 5A–C) and the averaged results summarized in the graph of Figure 5D. As shown in Figure 5D, isolated CC strips from castrated rats almost reached five-fold KCl induced contraction at $10^{-4}$ M PE, whereas sham and T replacement group generated around threefold KCl induced contraction. Similar to our previous report [20], micromolar concentrations of S1P (Fig. 6A and D) and FTY720-P (Fig. 7A and D) dose-dependently increased sham rat CC tension with 20 lM S1P and 2 lM FTY720-P elicited maximal contraction reaching 15% and 20% KCl induced force, respectively. These tensions were enhanced by castration. Maximal contraction induced by 20 lM SIP (Fig. 6B and D) and 2 lM FTY720-P (Fig. 7B and D) increased by almost twofold and threefold in castrated rats. Again, T administration fully restored both S1P (Fig. 6C and D) and FTY720-P (Fig. 7C and D) contractility. S1P receptors functional activity and its modulation by androgens were...
The current study demonstrated, for the first time, that testosterone modulated the expression and functional activities of S1P receptors in CC. Castration increased serum S1P concentration and up-regulated the expression of S1P2-3 receptors while down-regulated the expression of S1P1 receptor in CC at mRNA level. Accordingly, CCSM contraction induced by agonizing S1P receptors and relaxation mediated by antagonizing S1P receptors were augmented at functional level.

In accordance with our previous studies [5, 31], the surgically castrated rat model was validated by the significant weight loss of the androgen-sensitive organs (prostate and seminal vesicle), which was recovered after T supplementation. Also in line with previous observations [29, 36–39], the current study showed T deprivation-produced ED which was confirmed by the dramatically attenuated ICP/MAP elicited by ES of the cavernous nerve of castrated rats. This hypogonadism-related ED can be partially attributed to increased CCSM tone mediated by vasoconstrictors. Indeed, our in vitro contractility studies observed castration enhanced the α1-adrenergic agonist PE-induced rat CCSM contractile responses. Similarly, Wingo et al. [30] showed that isolated CC strips from castrated rats were more sensitive to PE stimulation. Although α-adrenergic signaling pathway is the main contributor responsible for maintaining penis in the flaccid state [40], antagonizing this system with α-blocker such as yohimbine and phentolamine possessed unsatisfactory efficacy for the treatment of ED [41, 42]. Therefore, other vasoconstrictors may play more important roles in hypogonadism ED.

S1P represents one of the key new additions to the list of ‘vasoactive’ substances that modulate vascular tone, as well as many other biological processes including cell proliferation, differentiation, survival and migration [6]. In the current study, rat serum S1P level reached over 1 ng/μl and it was increased by castration. It is widely recognized that serum S1P is secreted by activated thrombocytes and erythrocytes [9]. Campelo et al. [43] found T inhibited platelet activation and aggregation which was likely dependent on endothelial NO synthesis. Li et al. [44, 45] also observed platelet aggregation was enhanced in castrated rats when compared with sham and dihydrotestosterone replacement suppressed the increased platelet aggregation in castrated animals, indicating T might modulate S1P level through inhibiting platelet activation. However, it is known that tissue S1P levels are much lower and there is a large concentration gradient maintained between serum and extra vascular compartment [9]. Although we did not measure the S1P level in CC due to the difficulty of measuring tissue S1P concentration with HPLC, the critical enzymes for S1P formation in CC were detected. Interestingly, T deprivation down-regulated SphK1 expression but up-regulated SphK2. Both SphK1 and SphK2 could alter S1P level. Allende et al. [46] and Zemann et al. [47] found S1P levels were reduced by 50% in SphK1−/− mice, while Kharel et al. [48] showed S1P levels were reduced by only 25% in SphK2−/− mice. Hence, SphK1 played a more important role than SphK2 in maintaining S1P level. Indeed, many studies reported SphK2 was related to inducing cell apoptosis [49, 50]. The reduced SphK1 expression in CC might be negative feedback to up-regulated serum S1P. The up-regulation of SphK2 might mainly contribute to the apoptosis in CC which was observed in our previous study [51], rather than the increase in serum S1P concentration. The increased serum S1P level could contribute to hypogonadism ED, although the tissue S1P concentration in CC remains to be elucidated.

Consistent with our previous report [20], the current study found exogenous S1P and its analogue FTY720-P potently and dose-dependently induced rat CCSM contraction. Moreover, castration enhanced both S1P and FTY720-P-mediated CCSM contraction, which could attribute to increased S1P receptors activity. Indeed, we found T deprivation up-regulated the expression of S1P2-3 receptors while down-regulated the expression of S1P1 receptor in CC at mRNA level. Although S1P1 receptor was thought to mediate vasorelaxation through eNOS pathways [6, 8, 15], S1P1 receptor-specific agonist SEW2781 was not observed to induce CCSM relaxation and
contraction in our study (data not shown). Additionally, we observed S1P2 receptor-specific antagonist JTE-013, independent of NO, potently relaxed rat CC pre-contracted with PE in a dose-dependent manner. And JTE-013 exhibited more efficacy in relaxing CC strips from castrated rats at the lower concentration of 1 μM. The contractility of S1P system was further confirmed by our in vivo study. In a dose-dependent manner, ICI of JTE-013 alone antagonizing S1P2 receptor induced ICP rise. Consistent with our in vitro experiment, castrated rats were more sensitive to JTE-013 with almost 10-20% receptor induced ICP rise. Consistent with our dose-dependent manner, ICI of JTE-013 alone antagonizing S1P2 receptor was observed in a range of 50–500 nmol. Therefore, the augmented expression of S1P2-3 receptors could mainly contribute to the hyper-responsiveness to S1P stimulation for castrated animals. Similarly, several other studies demonstrated that S1P system expression and functional activity were modulated by sex hormone. Hemmings et al. [52] found that the expression of S1P1 was reduced in mesenteric resistance arteries from aged female rats with lower oestrogen level and restored with oestrogen replacement. And ovariectomy reduced the maximum S1P-induced vasoconstriction observed in aged rats. Moreover, Guo et al. [53] found plasma S1P levels were significantly higher in women than in men, and 17β-estradiol treatment increased S1P level in EA.hy926 cells. T supplementation recovered all aforementioned alterations including the expression and functional activities of S1P receptors.

Several limitations were noted in the present study. Firstly, S1P itself in vivo effect was not determined in this study because the vehicle (0.3 M NaOH) was not suitable for intracavernous injection. Secondly, as specific S1P receptor agonists or antagonists are limited or commercially available and hard to dissolve these lipids, the effect of T on S1P receptors functional activities in CC is not fully evaluated. Additionally, Western blot was not performed because of the low affinity of antibody to S1P receptors. However, our functional studies further confirmed the changed expression of S1P receptors.

In conclusion, we provide novel data that T positively regulate S1P system at both mRNA and functional level. It is suggested that S1P system dysregulation may contribute to hypogonadism-related ED, and antagonizing S1P receptors in CC might be a new target for treating ED.

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Disclosures

The authors have nothing to disclose.

Conflict of interest

The authors confirm that there are no conflicts of interest.

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