Erectile Dysfunction: Key Role of Cavernous Smooth Muscle Cells

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Erectile dysfunction is increasingly affecting men, from the elderly to young adults, being a sexual disorder related to the inability to generate or maintain a penile erection. This disorder is related to psychosocial factors such as anxiety, depression, and low self-esteem, to organic factors such as the presence of preexisting conditions like hypertension, diabetes and dyslipidemia. The pathophysiology of the disease is related to changes in the neurotransmission of the autonomic or the non-cholinergic non-adrenergic nervous system, as well as the release of local mediators, such as thromboxane A₂ and endothelin, and hormonal action. These changes lead to impaired relaxation of cavernous smooth muscle, which reduces local blood flow and impairs penile erection. Currently, therapy is based on oral vasodilation, such as sildenafil, tadalafil, vardenafil and idenafil, or by direct administration of these agents into the corpus cavernosum or by intraurethral route, such as alprostadil and papaverine. Despite this, studies that consolidate the understanding of its pathophysiological process contribute to the discovery of new more efficient drugs for the treatment of erectile dysfunction. In this sense, in the present work an extensive survey was carried out of the mechanisms already consolidated and the most recent ones related to the development of erectile dysfunction.

Keywords: smooth muscle, corpus cavernous, erection, flacity, autonomic nervous system, NANC, endothelin

INTRODUCTION

Erectile dysfunction is characterized by the inability to achieve and/or maintain a suitable penile erection for a satisfactory sexual intercourse (NIH, 1993) and represents the sexual dysfunction most studied in men (Ückert et al., 2007). The first reports of this clinical disorder were found in ancient Egyptian writings dating back over 5,000 years, where reductions in both number and quality of erections of different Egyptians were described (Smith, 1974; Shah, 2002).

The international consultation committee for sexual medicine on definitions, epidemiology and risk factors for sexual dysfunction conducted an extensive analysis of the worldwide prevalence of erectile dysfunction. In this view, the prevalence of the disease was 1–10% in men under 40 years of age, 29% in those aged 40–49 years, 20–40% in men aged 60–69 years and 50–100% in men over 70 years of age (Nicolosi et al., 2003). In addition, the worldwide prevalence of erectile dysfunction is estimated at 322 million men by 2025 (Costa and Potempa, 2012).

Data regarding the erectile dysfunction incidence are less abundant. However, the number of new cases of the disease per year varies from 19 to 66 cases per 1,000 men, according to studies
conducted in the United States, Netherlands, and Brazil (Johannes et al., 2000; Moreira et al., 2003; Schouten et al., 2005).

Erectile dysfunction is, predominantly, a vascular and benign disease, but it affects both physical and psychological health and has a significant impact on the quality of life of men and their partners, mainly due to the reduction of self-esteem and the commitment of the interpersonal relationship. The prevalence of the disease increases with age and can be seen as a serious public health problem (Medeiros-Júnior et al., 2014). The etiology is multifactorial, and, among the risk factors, the presence of cardiovascular diseases, sedentary lifestyle, smoking, diabetes, depression, anxiety and obesity are prominent (Alves et al., 2012).

Currently, erectile dysfunction is not limited to the reduction of sexual activity but acts as an indicator of systemic endothelial dysfunction. From the clinical point of view, this disease can precede cardiovascular events and can be used as an initial marker to identify men with high cardiovascular risk. In this case, patients with ED and no medical history of cardiovascular disease should be screened for cardiovascular disease (Gandaglia et al., 2014; Yafi et al., 2016; Hatzimouratidis et al., 2019).

Therefore, since alterations in the cellular signaling of cavernous smooth muscle cells can lead to this disease, this review focused on both contraction and relaxation processes that regulate penile erection, as well as highlights different targets as new approaches toward the development of new drugs for erectile dysfunction treatment.

### PHYSIOLOGICAL DETERMINANTS OF PENILE FLACCIDITY

Sympathetic stimulation is the primary mechanism responsible for maintaining the penis in the flaccid state. In this case, noradrenaline (NA) released from noradrenergic neurons stimulates its receptors in penile vessels and cavernous smooth muscle cells from penis to induce contraction (Mills et al., 2001).

Pharmacological characterization of adrenergic receptors has shown that the expression of \( \alpha \) subtype is 10 times more abundant than \( \beta \) subtype in cavernous tissue. In addition, \( \alpha_{1A}, \alpha_{1B}, \alpha_{1D}, \alpha_{1L} \) and \( \alpha_{2} \) (\( \alpha_{2A,C} \)) adrenergic receptors are expressed in human corpus cavernosum (Maggi et al., 2000; Andersson, 2011). Interestingly, activation of all \( \alpha \)-adrenergic receptor subtypes promotes smooth cavernous muscle contraction, since it was evidenced that presynaptic stimulation of \( \alpha_{2} \)-receptors inhibits non-adrenergic non-cholinergic transmission (NANC) (Filippi et al., 2002). Other important mediators of cavernous contraction include endothelin-1 (ET-1), prostaglandin F\(_2\alpha\) (PGF\(_2\alpha\)), thromboxane A\(_2\) (TXA\(_2\)) and angiotensin II (ANG II) (Tejada et al., 1991; Becker et al., 2001; Andersson, 2011).

In addition, the functional regulation of \([\text{Ca}^{2+}]_{i}\) to initiate a contractile response in the cavernous smooth muscle depends on two types of couplings: an electro- and a pharmacomechanical. The electromechanical coupling leads to the contractile response through membrane depolarization directly associated with increased extracellular K\(^+\) concentration \([\text{K}^{+}]_{e}\) (Figure 1) or the action of K\(^+\) channel blockers (Rembold, 1996).

Despite the importance of electromechanical coupling of cavernous smooth muscle cell contraction, the condition of penile flaccidity arises mainly from the pharmacomechanical coupling triggered by the binding of agonists to G protein coupled receptors (GPCRs) and the activation of the inositol cascade through the \( G_{\beta,\gamma} \). In this scenario, adrenaline, and NA binding to \( \alpha_{1}, \text{ET}-1 \) to \( \text{ET}_{\alpha/B} \), PGF\(_{2}\alpha \) to FP, TXA\(_2\) to TP and ANG II to \( \text{AT}_{1} \) receptors, which triggers the activation of phospholipase \( C_{\beta_{1}} \) (PLC\(_{\beta_{1}} \)) pathway. In this signaling, PLC\(_{\beta_{1}} \) hydrolyzes the phosphatidylinositol 4,5-bisphosphate (PIP\(_{2}\)), producing inositol 1,4,5-trisphosphate (IP\(_{3}\)) and diacylglycerol (DAG) (Berridge, 2008). IP\(_{3}\) induces Ca\(^{2+}\) release from the sarcoplasmic reticulum (SR) by activating the IP\(_{3}\) receptors (IP\(_{R}\)). In addition, in SR are expressed ryanodine receptors (RyR), caffeine sensitive Ca\(^{2+}\) release channels, which are activated by Ca\(^{2+}\) previously released via IP\(_{3}\)R in a process called Ca\(^{2+}\)-induced Ca\(^{2+}\) release (CICR) (Dellis et al., 2006; Mchale et al., 2006).

Following this signaling cascade, Ca\(^{2+}\) and DAG activate the Ca\(^{2+}\) dependent protein kinase (PKC), which, in turn, leads to [Ca\(^{2+}\)]\(_{i}\) increase through phosphorylation and the direct activation of the voltage-dependent Ca\(^{2+}\) channels (CaV) present in plasma membrane (Fukata et al., 2001). The increase in [Ca\(^{2+}\)]\(_{i}\) favors the interaction of this ion with calmodulin protein (CaM), leading to 4Ca\(^{2+}\)·CaM complex formation. This complex activates myosin light chain kinase (MLCK), which phosphorylates the regulatory myosin light chain (rMLC), promoting the interaction of myosin and actin.
filaments, triggering the process of contraction of the cavernous smooth muscle (Figure 2) (Webb, 2003).

Although $[\text{Ca}^{2+}]_i$ increase is transient, cavernous smooth muscle cells are able to maintain the contracted state even after the reduction of the intracellular levels of this ion. In fact, studies have reported an alternative pathway that contributes to the maintenance of smooth muscle contraction, designated as a $\text{Ca}^{2+}$ sensitization pathway (Figure 3). This pathway involves the modulation of myosin light chain phosphatase (MLCP), mainly by the small GTP binding protein $\text{G}_\text{q/11}$ and its associated kinase (ROCK), a serine/threonine kinase (Hori and Karaki, 1998). In the inactivated state,
RhoA is bound to guanosine diphosphate (GDP) and the guanine disassociation inhibitory protein (RhoGDI), forming a complex found in cytoplasm (Jin and Burnett, 2006).

Several contractile agonists, such as ET-1 and ANG II, which normally increase \([\text{Ca}^{2+}]_i\), via GPCRs lead to the direct activation of the RhoA guanine nucleotide exchange factor (RhoGEF) by the G12/13 proteins, which activate RhoA by exchanging GDP for guanosine triphosphate (GTP) in this protein (Somlyo and Somlyo, 2003). Thus, RhoA-GTP is translocated to plasma membrane, where it remains anchored by geranilgeranilization, and activates the ROCK. The activity of RhoA is reduced by the action of Rho-GTPase activating protein (RhoGAP), which culminates with the GTP cleavage in GDP and the formation of the inactive RhoA-GDP/RhoGDI complex, returning to cytoplasm (Burridge and Wennerberg, 2004).

After activation by RhoA-GTP, ROCK directly phosphorylates rMLC and the regulatory subunit MYPT1 of MLCP, making it inactive, promoting the maintenance of the phosphorylated state of rMLC and, consequently, the contraction force. In addition, it can activate a Ca\(^{2+}\) dependent kinase protein, also known as zipper-interacting protein kinase (ZIPK). The ZIPK phosphorylates directly the rMLC, however its main target is the Thr\(^{696}\) residue of MYPT1 which, when phosphorylated, inhibits the action of MLCP (Murthy, 2006).

CPI-17 (PKC-dependent phosphatase inhibitor of 17 kDa) represents a substrate for both ROCK and PKC. When phosphorylated, CPI-17 binds to the PP1c catalytic subunit of MLCP to inhibit its enzymatic activity and prolong smooth muscle contraction (Kitazawa et al., 2000).

RhoA-GTP also stimulates phospholipase D (PLD), this enzyme is mainly associated with intracellular membranes, but is also found in the plasma membrane and is specific for phosphatidylcholine (PC), releasing phosphatidic acid (PA) that through the action of phosphohydrolase enzyme is dephosphorylated to DAG leading to sustained activation of PKC. Activation of PKC may also be dependent on \(G_{q/11}\)-PLC\(\beta1\), which forms DAG by PIP\(_2\) hydrolysis. PKC can phosphorylate the Thr\(^{38}\) residue of CPI-17, thereby increasing its inhibitory potency over PP1c by more than 1000-fold, inhibiting the action of MLCP (Murthy, 2006; Berridge, 2009).

Mitogen-activated protein kinases (MAPK) have been associated with increased smooth muscle contractility, proliferation, and chemotaxis. Among them, MAPK ERK1/2 and p38 are especially noteworthy, especially for their activity on vascular smooth muscle, however, molecular mechanisms that directly alter cavernous contractility are not fully understood (Berridge, 2009).

PHYSIOLOGICAL DETERMINANTS OF PENILE ERECTION

Relaxation of penile corpus cavernosum occurs in response to NANC and cholinergic neurotransmission, with nitric oxide (NO) as the most important neurotransmitter. The synthesis of NO is catalyzed by nitric oxide synthase (NOS) enzyme that in the presence of \(O_2\) converts l-arginine to l-citrulline and NO in both endothelial cells and nerve endings. Three distinct isoforms of NOS are expressed in human cavernous smooth muscle cells, the constitutive and calcium dependent isoforms include neuronal (nNOS) and endothelial (eNOS). In addition, the inducible isofrom (iNOS) was identified, especially when there is tissue injury (Burnett et al., 1993; Gonzalez-Cadavid et al, 1999).
In endothelial cells, eNOS activation may occur in response to shear stress induced by increased blood flow in the penile vessels, leading to phosphatidylinositol 3-kinase (PI3K) activation, which phosphorylates PIP2 and forms the 3,4,5-phosphatidylinositol trisphosphate (PIP3). PIP3 recruits and activates phosphoinositide-dependent kinase-1 (PDK-1) that, in turn, phosphorylates and activates Akt, also known as protein kinase B (PKB). Akt is a serine/threonine kinase that phosphorylates directly the eNOS on its Ser1177 residues, resulting in increased activity of this enzyme, a phenomenon that contributes to continuous NO production (Hurt et al., 2002; Sommer et al., 2002).

Other substances are responsible for modulating cavernous smooth muscle relaxation, such as vasoactive intestinal peptide (VIP) and calcitonin gene related peptide (CGRP), released by cholinergic nerve endings. The endothelium also releases relaxing prostanoids, such as prostacyclin (PGI2) and prostaglandins E types 1 and 2 (PGE1 and PGE2) (Molderings et al., 1992; Porst, 1996).

Although these substances are involved in cavernous smooth muscle relaxation, NO is considered the most important physiological mediator for penile erection. When there is a sexual stimulus, it is released by NANC neurons or endothelial cells in response to ACh and, as a soluble gas; it diffuses and activates its soluble guanylyl cyclase receptor (sGC) in cavernous smooth muscle cell (Figure 4) (Andersson, 2011). The sGC converts GTP into cyclic guanosine monophosphate (cGMP) that activates cGMP-dependent protein kinase (PKG). This kinase phosphorylates several substrates to promote cavernous muscle relaxation (Figure 4). These include: 1) activation of K+ channels that indirectly, by repolarization or hyperpolarization, can inhibit CaV; 2) direct inhibition of CaV, decreasing Ca2+ influx (Rembold, 1996); 3) increase in Ca2+ sequestration and extrusion; 4) decrease of [Ca2+]i by Na+/Ca2+ exchanger (NCX) activation that acts in a ratio of 3/1 (Blaustein, 1989); 5) inhibition of MLCK, reducing its affinity to 4Ca2+-CaM complex, preventing the phosphorylation of rMLC and, consequently, the contractile process (Rembold, 1996); 6) inactivation of IP3R that reduces Ca2+ release from RS (Woodrum and Brophy, 2001); 7) RhoA inactivation by phosphorylation of Ser188 residue, displacing it into the cytosol; 8) MLCP activation by phosphorylation of Ser695.
residue from MYPT1 subunit which prevents phosphorylation of Thr696 residue by ROCK; 9) MLCP activation by telokin phosphorylation (endogenous activator of MLCP), being an independent mechanism of RhoA (Murthy, 2006).

From the physiological point of view, NO derived from nNOS is responsible for the onset of erection, whereas NO produced by eNOS in response to shear stress contributes to the maintenance of penile rigidity during erection (Andersson, 2011).

The other relaxing mediators, such as prostaglandins E₃,₂ and prostacyclin act on EP₂, EP₄ and IP receptors, respectively, all coupled to Gₛ protein. The a-GTP subunit activates adenyl cyclase (AC), which converts adenosine triphosphate (ATP) into cyclic adenosine monophosphate (cAMP), responsible for activating cAMP-dependent protein kinase (PKA). This kinase phosphorylates the same targets of PKG to promote cavernous muscle relaxation, with the exception of IP₃R (Figure 4) (Andersson, 2011).

Due to cGMP and cAMP importance in [Ca²⁺]ᵢ reduction, highlights the essential role for the regulation of erectile function performed by the phosphodiesterases enzymes (PDEs), which hydrolyze cGMP and cAMP to their non-cyclic forms (GMP and AMP, respectively) and enclose the signaling cascade that results in cavernous smooth muscle relaxation. Each PDE family includes multiple isoforms that can act on a specific cyclic nucleotide or on both types. In penis, the predominant phosphodiesterase isofrom is 5 (PDE5), which acts specifically on cGMP (Mancina et al., 2005).

As the balance between the contraction and relaxation processes of the cavernous smooth muscle regulates the processes of penile flaccidity and erection, any deregulation in steps of these signaling pathways may compromise penile function and thus contribute to erectile dysfunction development.

CAVERNOUS SMOOTH MUSCLE CELLS AS TARGETS FOR THE PHARMACOLOGICAL TREATMENT OF ERECTILE DYSFUNCTION

Intracavernous and Intraurethral Therapies

Currently, they represent the last line of treatment and show, as advantages, speed to start, less than 10 min, and quality of penile erections, even in the absence of sexual stimulation. In this case, the man needs to receive previous training to be able to apply the intracavernous injection or to position the intraurethral suppository. The vasodilatory substances commonly used to induce penile erection are alprostadil, papaverine and phentolamine alone or in combinations of two or three of them with high success rates therapeutic use (90%) (Hatzimouratidis and Hatzichristou, 2005; Hatzimouratidis et al., 2019).

Alprostadil is the synthetic prostaglandin E₃, thus, when it binds to EP₂,₄ receptors it activates the AC signaling pathway, culminating in an increase in the intracellular concentration of cAMP ([cAMP]ᵢ), which through previously described mechanisms triggers corpus cavernosum relaxation and, consequently, penile erection. It is used in intracavernous injection therapy and as a suppository for intraurethral use (Moreland et al., 2003).

Papaverine non-selectively inhibits phosphodiesterases and thus increases [cAMP]ᵢ and the intracellular concentration of cGMP ([cGMP]ᵢ), with similar results to that seen with alprostadil, however, it is only used in intracavernous injection therapy. In addition to these substances, selective α₁-adrenoceptor antagonist phentolamine prevents vasoconstriction resulting from the activation of the PLCβ₃ pathway and thereby assists penile erection. It is used only in intracavernous injection therapy (Virag et al., 1991).

The main risk of using these drugs is priapism, which consists of a painful and prolonged penile erection (longer than 2 hours), independent of sexual desire and due to insufficient penile blood drainage. In cases of priapism, emergency medical attention is indicated in order to perform blood aspiration of the corpus cavernosum, the application of phenylephrine, a selective α₁-adrenergic agonist, which promotes local vasoconstriction through the activation of the PLCβ₃ pathway and, in more severe cases, the creation of shunts between the corpus cavernosum and the glans or spongy body through surgical procedures (Linet and Ogrinc, 1996).

Oral Therapy

Primary pharmacotherapy for erectile dysfunction treatment involves the use of PDE5 inhibitors, such as sildenafil, which is the prototype of the group, tadalafil, vardenafil and iodenafil. Mechanically, PDE5 inhibitors increase [cGMP], and thus initiate the cascade of intracellular events that result in relaxation of the corpus cavernosum and promote penile erection. However, previous sexual stimulation is essential to increase intracellular NO levels and, consequently, to generate cGMP (Yafi et al., 2016).

PDE5 inhibitor use improves the sexual performance of the men, without alteration of the libido. Recently, the discovery that the use of these inhibitors reduces the refractory period, a time when a temporary physiological flaccidity occurs immediately after ejaculation, has promoted an increase in the demand for these drugs by young and potent men (Ekmekcioğlu et al., 2005; Hatzimouratidis et al., 2019).

Despite the great therapeutic success of these drugs, about 30–40% of the men affected by erectile dysfunction do not respond to this first line of treatment. Additionally, PDE5 inhibitor use should be performed with caution in patients with cardiovascular impairment, such as uncontrolled hypertension and unstable angina. In addition, the use of these drugs is contraindicated in patients who use nitrates because of the increased risk of severe hypotension (Eardley and Sethia, 2003).

The major side effects related to therapy with PDE5 inhibitors include headache, nasal congestion, and facial flushing. In addition, visual and auditory changes, including macular degeneration, anterior ischemic optic neuropathy, hearing loss and tinnitus are recent reasons for precautions. These complications appear to be related to the mild inhibitory action of these drugs on PDE6 (Zelefsky et al., 2014; Hatzimouratidis et al., 2019).
**Targets Studied as an Alternative to the Use of Oral Therapy**

Despite the use of oral therapy being the first line of treatment for erectile dysfunction, there are patients who are refractory to treatment, creating a field of research for new drugs. In this sense, there are different targets in the contractile machinery of the penile corpus cavernosum.

In clinical trials, the use of agents that act as activators of the soluble guanylyl cyclase enzyme have been shown to improve the development of penile erection. Additionally, improvement in erectile dysfunction was also observed with the therapeutic use of Rho-kinase inhibitors. However, in terms of clinical efficacy and adverse effect profile, there was no superiority when compared to PDE inhibitors (Sanofi-Aventis, 2010; Bayer, 2017).

Another approach considered in the function of cavernous smooth muscle cells is to promote the opening of potassium channels, which would lead to a reduction in the intracellular concentration of calcium and, consequently, to relaxation. In this sense, the use of calcium-sensitive Maxi-K ion channel gene (hSlo cDNA) was evaluated in clinical phase I studies, demonstrating a good safety of use, however, there was no improvement to the use of PDE inhibitors, limiting clinical progress (Melman, 2006).

**CONCLUSION**

Balance among flaccidity and penile erection depends on the timing between the contraction and relaxation processes of cavernous smooth muscle cells. Thus, \([\text{Ca}^{2+}]_i\) is the central point of regulation of smooth muscle tone. In this view, the increase in \([\text{Ca}^{2+}]_i\) leads to contraction of these cavernous cells and, consequently, maintains the penis in the flaccid state, while relaxation, which promotes penile erection, is mediated by the reduction of \([\text{Ca}^{2+}]_i\). In this sense, the imbalance of these mechanisms is related to ED development.

Currently, the main therapeutic lines for ED present as targets steps of contraction and relaxation signaling in cavernous smooth muscle cells. Therefore, further research targeting different components of electro- and pharmacomechanical couplings of cavernous smooth muscle cell arise as new targets for the development of promising drugs for the treatment of this disease.

**AUTHOR CONTRIBUTIONS**

IS, EF, and LV have made effort on researching data and writing the article, FC and BS were the advisors, contributing with the correction and review of the article.

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