Abstract

Purpose: To evaluate the expression of endothelial and inducible NOS in addition to the miRNA-27b in the corpus cavernosum and peripheral blood of healthy rats, diabetic rats, alcoholic rats and rats with both pathologies.

Methods: Forty-eight Wistar rats were divided into four groups: control (C), alcoholic (A), diabetic (D) and alcoholic-diabetic (AD). Samples of the corpus cavernosum were prepared to study protein expressions of eNOS and iNOS by immunohistochemistry and expression of miRNA-27b in the corpus cavernosum and peripheral blood.

Results: Immunohistochemistry for eNOS and iNOS showed an increase in cavernosal smooth muscle cells in the alcoholic, diabetic and alcoholic-diabetic groups when compared with the control group. Similarly, the mRNA levels for eNOS were increased in cavernosal smooth muscle (CSM) in the alcoholic, diabetic and alcoholic-diabetic groups and miRNA-27b were decreased in CSM in the alcoholic, diabetic and alcoholic-diabetic groups.

Conclusion: The major new finding of our study was an impairment of relaxation of cavernosal smooth muscle in alcoholic, diabetic, and alcoholic-diabetic rats that involved a decrease in the nitric oxide pathway by endothelium-dependent mechanisms accompanied by a change in the corpus cavernosum contractile sensitivity.

Key words: Alcoholism. Diabetes Mellitus. Erectile Dysfunction. Rats.
Introduction

Erectile dysfunction is characterized by a persistent inability to achieve and/or maintain an erection sufficient for satisfactory sexual performance\(^1\). In the last 25 years there has been a revolution in the treatment of Erectile Dysfunction (ED) when the role of nitric oxide (NO) in the physiology of erection can be understood and phosphodiesterase 5 (PDE5) inhibitors have been used in the treatment of ED\(^2\).

It is estimated that the overall prevalence of erectile dysfunction (ED) is approximately 20\% of the male population\(^3\). In 1995 it was estimated that there were 152 million men with ED in the world and 322 million were projected with this disease by 2025\(^4\). Prevalence increases with age and comorbidities such as type II diabetes, obesity, cardiovascular diseases, hypertension, dyslipidemia, and benign prostatic hyperplasia (BPH)\(^5\)-\(^7\).

Nowadays it is widely accepted that the release of NO into the cavernous tissue is the most important event triggering erection; it arises as a neurotransmitter of non-adrenergic and non-cholinergic neurons (NANC) and as vasodilator derived from the endothelium, acting in order to increase intracellular levels of cGMP in the cavernous vascular smooth muscle\(^8\)-\(^13\). It is a consensus that NO derived from nitricergic nerve endings (produced by neuronal nitric oxide synthase - nNOS) is responsible for rapid and initial vasodilation, whereas endothelial NO (produced by endothelial nitric oxide synthase - eNOS) contributes to the maintenance of erection\(^13\). Diabetics have a higher incidence of ED, probably due to vascular and neurological changes as a result of the disease, however, this dysfunction can occur even with normal glycemic indexes. Pathological alterations in the cavernous arteries, ultrastructural alterations in the smooth muscle cavernosa and decrease in endothelium relaxation dependent on the smooth muscle cavernosa were observed in samples of cavernous tissue of diabetic patients\(^14,15\).

Alcoholism has also been associated with ED; several mechanisms are described, among which neurological, endocrine, psychological and supposed vascular alterations\(^15\). However, findings suggest that a decrease in the relaxation response of the corpus cavernosum in rats with chronic alcoholism occurs due to the decrease of the endothelial response to the cholinergic stimulant\(^17\).

In search of clarifying molecular events associated with ED present studies have analyzed the role of microRNAs (miRNA). MicroRNAs have recently been recognized as a class of short, highly conserved RNA molecules (around 22 nucleotides), which are highly conserved (among distant species), whose function seems to be to regulate gene expression at the post-transcriptional level\(^18\). The present study aims at evaluating the gene and protein expressions of iNOS and eNOS isoforms, as well as microRNA-27b in the corpus cavernosum and in the peripheral blood of rats submitted to model of alcoholism and diabetes.

Methods

A total of 48 male Wistar rats (Rattus norvegicus) from the University of São Paulo - Ribeirão Preto Campus, after approval by the Ethics Committee of our institution, were used. They were divided into 4 groups and followed for 4 weeks after the adaptive period: control group (C), alcoholic group (A), diabetic group (D), alcoholic-diabetic group (AD), all groups consisting of 12 animals each.

For the animals in the ethanol group (A, AD), the model of “semi-voluntary alcoholism” was used, in which the ethanol solution at 20%
was the only liquid available to these animals, following a model proposed by Tirapelli in 2006, where there was an adaptive phase that consisted of the supply of ethanol in increasing weekly concentrations of 5, 10 and 20%, starting the experimental phase after the third week of treatment\textsuperscript{19}.

For the immunohistochemistry, the corpus cavernosum of control (n = 6), alcoholic (n = 6), diabetic (n = 6) and alcoholic-diabetic rats (n = 6) were immediately removed and fixed for 24 h in ice-cold 0.1 mol/l PBS (pH 7.4), containing 4% paraformaldehyde, followed by cryoprotection in 15% of sucrose for 4 h and 30% sucrose overnight at 4 C. Immunohistochemical analyses of longitudinal sections (10 µm) of the corpus cavernosum were carried out via avidin-biotin-peroxidase (Novostain Super ABC Kit - Universal, NCL-ABCu, Novocastra Laboratories Ltd, Newcastle upon Tyne, UK) - (universal Kit mach 4 BIOCARE). The longitudinal sections were incubated with 3% H2O2, followed by antigen retrieval with a moist heat steam cooker Optistream Plus (Krups North America, Inc., Millville, New Jersey, USA) with 10 mM citrate buffer at pH 6.0 for 35 minutes. Then, the sections were incubated for 24 hours in a primary antibody: ETA and ETB diluted 1/300 in PBS solution of bovine serum albumine (BSA). Subsequently, the blocking of the endogenous biotin was performed (Biotin Blocking System, Dako North America, Inc., Carpinteria, USA) and only then the sections were incubated with secondary antibody HRP kit MACH 4-Universal Polymer (M4BD534, Biocare Medical) and then with avidin-biotin-peroxidase kit same (1/200 in PBS). Color was developed through the addition of diaminobenzidine (Sigma Chemical, St. Louis, USA).

The sections were dehydrated in ethanol, cleared with xylene and mounted under the cover slip with Permount liquid (Fisher Scientific Company LLC, Fair Lawn, New Jersey, USA).

The slides for the immunohistochemical study were analyzed using the Zeiss microscope Axioskop 2 plus model in magnification of 400 times. The number of cells with positive staining for eNOS and iNOS isoforms were measured by using a camera (Axio Cam, Zeiss, Germany) and the program Axiovision 4.6 (Zeiss, Germany). Analysis of expression profile of the miRNAs-155 and 199

The expression profile of the miRNA-27b was analyzed in blood and cavernous tissue samples from each animal. Total cellular RNA was extracted using Trizol Reagent (Invitrogen, Carlsbad, CA) and RNA was reverse transcribed to single-stranded cDNA, using a High Capacity Kit (Applied Biosystems, Foster City, CA) according to the manufacturer’s protocol. For the quantitative analysis of the miRNA-27b, we used the commercially available system TaqMan Assay-ondemand (Applied Biosystems). Reverse transcription was performed using 5ng total RNA for each sample in 7,5µL of the total reaction mixture. The cDNA obtained was diluted 1:4 and 4.5µL was used for each 10µL of the quantitative real-time polymerase chain reaction mixture using the TaqMan Master Mix (Applied Biosystems). All reactions were carried out in duplicate and analyzed with the 7500 Sequence Detection System apparatus (Applied Biosystems). Data were analyzed using the ABI-7500 SDS software. The total RNA absorbed was normalized on the basis of the Ct value for U6. The variation in expression among samples was calculated by the 2-∆∆Ct method, with the mean ∆Ct value for a group of 6 samples from control rats being used as a calibrator.

Statistical analysis

For the evaluation of all studies in this research (protein expression and gene expression), statistical analysis was performed using the Kruskal-Wallis test and Dunn’s multiple comparison post-test. We used the
GraphPad Prism program 6:00 version for Windows (GraphPad Software, San Diego - California USA) and considered statistically significant p<0.05.

### Results

**Immunohistochemistry**

From the global evaluation of the labeling pattern by immunohistochemistry of the expressions of eNOS and iNOS in the cavernous bodies, in all the groups studied, low tissue expression was observed in the animals in all groups, being slightly higher in the animals of the AD group (Figures 1 and 2).

**Figure 1** - Representative immunohistochemical photomicrograph of the global expression of the eNOS in rat penile tissue sections from alcoholic-diabetic rats. Observe diffuse endothelial marking in the erectile tissue. Vascular or cavernous space (CE). (x400).

**Expression profile of genes and miRNAs**

We found no significant difference between the groups analyzed. There was a non-significant increase in the expression profiles of the eNOS and iNOS in both D and AD groups (Figures 3 and 4).

**Figure 2** - Representative immunohistochemical photomicrograph of the global expression of the iNOS in rat penile tissue sections from alcoholic-diabetic rats (x400).

**Figure 3** - Representation of mean values (± standard error) of eNOS gene expression in cavernous tissue samples of the groups studied. There was no significant difference in the groups (p=0.0968, Kruskall Wallis test).

**Figure 4** - Representation of mean values (± standard error) of iNOS gene expression in cavernous tissue samples of the groups studied. There was no significant difference in the groups (p=0.0629, Kruskall Wallis test).
A significant decrease in the expression of miRNA-199 was observed in the tissue of the corpus cavernosum and blood of the rats in the AD group when compared to the C group (Figure 5).

Figure 5 - Representation of mean values (± standard error) of microRNA-27b expression in the blood and cavernous tissue samples of the groups studied. There was a significant difference between the control group and the alcoholic group (Kruskall Wallis test, *p <0.05, Kruskall Wallis test, Dunn’s post-test).

Discussion

Penile erection is a complex neurovascular process dependent on the relaxation of the smooth muscle of the corpora cavernosa and NO, which has been shown to be released from the nerves, is the major relaxing agent in several mammals\(^{20,21}\).

Lizarte et al. investigated the effect of diabetes mellitus and chronic ethanol consumption on the relaxation mediated by nitric oxide in the smooth muscle of the cavernous bodies of rats for four weeks\(^{22,23}\). When performing the analysis of immunohistochemistry and gene expression by real-time PCR, high expressions of eNOS and iNOS were observed in the smooth muscle fibers of the rats in the three groups, ethanol, diabetic and diabetic associated to the chronic consumption of ethanol, when compared to the animals in the control group. These studies demonstrated a decrease in the acetylcholine-induced relaxation capacity in the musculature of the corpora cavernosa via the endothelium-dependent mechanism suggesting, therefore, that the chronic consumption of ethanol induces endothelial dysfunction. Compared to the described findings, similar results were demonstrated in our study in the analysis of the eNOS and iNOS gene expressions of the animals in the diabetic and diabetic alcoholic groups, and also an increase in the eNOS and mainly iNOS expressions in the animals in the alcoholic diabetic group.

Our results demonstrated, through the time PCR technique, increased gene expressions of eNOS and iNOS in the group of diabetic animals and also in the group of diabetic animals associated with alcoholism. In the results of the eNOS and iNOS protein expression analysis by immunohistochemistry, diffuse expression was observed in the cavernosal tissue of both eNOS and iNOS where eNOS presented greater expression in the alcoholic-diabetic group and iNOS also presented increased expression in the alcoholic-diabetic group but a higher expression still in the group of animals submitted to alcoholism alone.

Recent studies have shown that specific classes of miRNAs affect the function of endothelial cells that are involved in many aspects of vascular biology, producing different factors that regulate smooth muscle cell proliferation and vascular tone. Among
these, miRNA-27b has been reported to have a significant role in the angiogenesis of endothelial cells. Our results demonstrated low gene expression of miRNA-27b in the blood and corpus cavernosum of alcoholic, diabetic and alcoholic-diabetic groups, when compared to control group. We know, however, that the suppression of miRNA-27b by ethanol may be responsible for the inhibition of angiogenesis of the endothelial cells, suggesting that the suppression of its expression inhibits the formation of endothelial cells, thus promoting an imbalance in the regulatory mechanisms of nitric oxide that are essential for the occurrence of angiogenesis. As a result, chronic alcoholism plays an important role in the pathogenesis of erectile dysfunction.

Our results demonstrated a possible regulation of the eNOS and iNOS isoforms by microRNA-27b, since they showed opposite expression levels, that is, the gene and protein expressions of eNOS and iNOS were increased in the diabetic and alcoholic-diabetic groups while the expression of microRNA-27b was low in the same groups.

## Conclusions

The major new finding of our study was an impairment of relaxation of cavernosal smooth muscle in ethanol, diabetic, and ethanol-diabetic rats that involved a decrease in the nitric oxide pathway by endothelium-dependent mechanisms accompanied by a change in the corpus cavernosum contractile sensitivity. Early diagnosis and treatment of complex diseases such as erectile dysfunction, alcoholism, and diabetes may benefit from miRNAs studies designed to identify disease-specific molecular markers that provide a fingerprint of the condition or to identify potential therapeutic targets. Rapid advances in the field of genomics offer new diagnoses and potential for screening even for diseases such as erectile dysfunction, alcoholism, and diabetes.

## References

10. Burnett AL, Lowenstein CJ, Bredt DS, Chang
Expression profiles of eNOS, iNOS and microRNA-27b in the corpus cavernosum of rats submitted to chronic alcoholism and Diabetes mellitus

Cunha JP et al.

Acta Cir Bras. 2017;32(1):38-45


