Modulation of Oxidative Stress Associated with Experimentally-Induced Benign Prostatic Hyperplasia in Rats by Zapoteca portoricensis Root Extracts †

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Abstract: Background: The existence of oxidative stress in the pathogenesis of benign prostatic hyperplasia (BPH), characterized by elevation in markers of oxidative stress/lipid peroxidation (8-hydroxyguanosine, malondialdehyde and 8-hydroxynonenal) and reduction in antioxidant status (catalase, superoxide dismutase, glutathione peroxidase and reduced glutathione) is scientifically documented. We hypothesize that a good treatment regimen for BPH should return the pro-oxidant/antioxidant status to normal; hence, pro-oxidant/antioxidant status is an indirect indicator of treatment response. In this study, the effect of crude methanol extract (CME) of Zapoteca portoricensis root and its methanol (MF) and ethyl acetate (EAF) fractions on the pro-oxidant/antioxidant status of experimentally-induced BPH was investigated. Methods: Forty-five Wistar albino rats (7 weeks, 180–200 g) used in this study were divided into nine groups (n = 5). Group 1 served as normal control. BPH was induced in groups 2–9 by daily subcutaneous administration of dihydrotestosterone (400 μg/mL) and estradiol (80 μg/mL) for 28 days. Group 2 served as BPH-control (was left untreated) while group 3 received dutesteride (Avodart®). Groups 4 and 5, 6 and 7, and 8 and 9 received, by gavage 200 and 400 mg/kg/d b.w. of CME, 200 and 400 mg/kg/d b.w. of MF, and 200 and 400 mg/kg/d b.w. of EAF, respectively for 14 days. Results: There were increased prostatic specific (PSA) and malondialdehyde but reduced antioxidant status in BPH-control relative to normal control. At 400 mg/kg/d b.w., CME, MF and EAF decreased prostatic specific antigen by 55.91%, 57.54% and 56.75%, respectively comparable to 58.80% by dutesteride. In addition, the results of histological assessment of prostate tissues of the experimental rats fed extracts demonstrate an improved prostate status. Conclusion: The extracts returned the pro-oxidant/antioxidant status modified by BPH to normal. These findings may justify the plant’s folkloric use and suggest that extracts can be exploited further as potential source of entities for managing BPH.

Keywords: Zapoteca portoricensis; prostatic specific antigen; benign prostatic hyperplasia; oxidative stress; antioxidant

1. Introduction

Benign prostatic hyperplasia (BPH) also called benign enlargement of the prostate (BEP) or prostate Adenoma, is a common event in aging and constitutes a lot of health burden and prevalent among aging
It is a multi factorial disease characterized by a non-malignant, uncontrolled proliferation of the smooth muscles, stromal and epithelial cells within the transition zone of the prostate gland. The etiology is not clear so far, however, it is the commonest non-cancerous form prostate cell growth and initial development usually after 40 years of age with ~40% of males [2–5]. Prevalence statistics is well documented in the developed world but not so in Africa, especially Nigeria A variety of factors chiefly age, genetics, lifestyle, hypertension, obesity, diabetes and insulin resistance have been linked to the development of BPH The prostate, an exocrine gland found in the male reproductive system, produces, secretes and controls the flow of seminal fluid expelled at the time of sexual climax together with the spermatozoa The testosterone metabolite- Dihydroxytestosterone (DHT) and oestrogen are believed to play a role in the etiology of BPH. Studies have shown that as men age the active testosterone in the blood decreases leaving a higher proportion of estrogen. Thus, the accumulation DHT in the prostate gland encourages the growth of prostate cells. In addition, the development of BPH, an age dependent disease, has been associated with increased lipid peroxidation, oxidative stress, inflammatory process and decreased levels of antioxidants. Studies have shown an increased malondialdehyde (MDA) levels, a product of lipid peroxidation and an indicator oxidative stress, a decrease in serum levels of antioxidants such as glutathione (GH) glutathione peroxidase(GSH-Px),and glutathione reductase(GR), Superoxide dismutase (SOD) and non-enzymatic antioxidants vitamins C and E, in the plasma and erythrocytes The use of surgical procedures, alpha-blockers/adrenoceptor antagonists (Terazosin and Doxazosin), 5-alpha-reductase inhibitors (finasteride and dutasteride) and some forms of combination therapies in the management of BPH achieve reduction in mass of the prostate gland or relaxation of the muscle tone The associated side effects cost and risks of surgery however, have led to increased search for alternatives in managing this disease. Antioxidants therapy and plant-derived dietary polyphenolic compounds, such as flavonoids with cancer cells and chemo preventive potential seems to be a promising therapy Phyto-therapeutic preparations most commonly used in BPH include 5-beta- sterol from Hypoxis roperi (African star grass), Pygeum from Prunies africanaum, the roots of Urtica dioica and Serenoa repens from Saw palmetto [12–14]. Zapoteca portoricensis (jacq), HM Hernandez (Fabaceae) is a perennial seasonal shrub and a native of West Africa, West Indies and the Atlantic coast of America. The peoples of eastern and southern Nigeria have found the different plant parts useful in traditional medicinal practices [15] for diarrhea, convulsion and tonsillitis [16,17]. Its antiulcer, antimicrobial and anti-inflammatory activities have been reported. The root extracts have been shown to have trypanocidal activity against Trypanosoma brucei rhodensiense Reported phytochemical analysis of the root and other plant parts indicated the presence phytochemicals such as saponins, resins, glycosides, flavonoids, alkaloids, terpenoids and steroids The traditional use of Zapoteca portoricensis root extract in the management of microbial and inflammatory disorders and its folkloric use in the management of BPH motivated this study.

2. Methods

Plant materials:

The roots of Zapoteca portoricensis were collected from Orba and Eha-Alumona, in Nsukka zone of Enugu state, Nigeria. The roots were identified and authenticated by Mr. A. Ozioko of the Bioresource development and Conservation Program (BDCP) Research Centre Nsukka, Enugu state, Nigeria. The root samples were air dried for three weeks to constant weight at room temperature (29 °C–35 °C) and ground into uniform coarse form using a milling machine. The methanolic extract was prepared by soaking 2000 g of dried pulverized roots samples in 1.5 L of methanol for 72 h. It was filtered using Whatman Number 4 filter paper and the filtrate was concentrated using Rotary evaporator at regulated temperature. The methanol extract obtained was fractionated by column chromatography using 1.3 L of methanol and 1.3 L of ethyl acetate as solvents.

Animals:
Adult male rats Wistar strain (80–120 g) obtained from the animal holding unit of the Department of Zoology and Environment Biology, University of Nigeria, Nsukka were used in the study. The animals were housed under standard conditions (25 ± 2 °C and 12 h light/dark cycle). The rats were fed twice daily with standard pellets (Grand Cereals Ltd., Enugu Nigeria) and had unrestricted access to clean drinking water. The guide for the care and use of laboratory animals procedures were followed in this study [20].

Experimental Design-Animal Grouping

For this study, Forty five male albino rats divided into nine groups of five rats each representing groups receiving different doses of crude methanol extract, methanol and ethyl acetate fractions of the plant root sample. Group 1 is the normal control (Not induced and untreated), Group 2 is the positive control (Induced and Untreated), and Group 3 is the standard control (induced and treated with standard drug, Dutasteride/Avodart). Groups 4 and 5 were induced with BPH and treated with 200 mg/kg and 400 mg/kg body weight of crude methanol extract respectively. Groups 6 and 7 were induced with BPH and treated with 200 mg/kg and 400 mg/kg body weight of methanol fraction respectively. Groups 8 and 9 were induced with BPH and treated with 200 mg/kg and 400 mg/kg body weight of ethyl acetate fraction respectively.

Induction of Benign Prostatic Hyperplasia (BPH)

Experimentally developed BPH model was created in the male Wistar rats by subcutaneous administration of dihydrotestosterone (DHT) and estradiol (300 μg/mL and 80 μg/mL respectively) dissolved in olive oil for 28 days [21–23]. Before induction blood was drawn by ocular puncture for initial prostatic specific antigen (PSA) assay.

Treatment of Benign Prostatic Hyperplasia (BPH) with Plant Extracts

The crude methanol extract, methanol and ethyl acetate fractions were dissolved in tween-80 and normal saline. The dissolved standard drug, Dutasteride (Avodart) was administered orally to group 3 once daily while crude methanol extract and fractions were administered orally to groups 4–9 once daily. The treatment lasted for 14 days. At the end of the treatment with plant extracts and fractions blood was drawn by ocular puncture into clean EDTA bottles for final PSA assay and determination of biochemical parameters.

Determination of biochemical indices

The serum prostatic specific antigen (PSA) levels expressed in ng/mL were analysed with a PSA ELISA kit according to the manufacturer’s instructions (Biocheck, Inc., South San Francisco, CA, USA. Catlog number: BC1019) while the activity of serum acid phosphatase(ACP) was assayed using the colorimetric method of Tietz [24] as outlined in Randox Kit(UK). Similarly, lipid peroxidation was estimated by measuring spectrophotometrically the malondialdehyde (MDA) concentration as described by Wallin et al. [25], superoxide dismutase (SOD) activity was assayed by the method of Arthur and Boyne [26] as contained in the Randox Kit, UK used and the catalase (CAT) activity was assayed by the method of Sinha [27]. In addition, reduced glutathione concentration was determined by the method of King and Wootton [28] and vitamin C concentration was determined using the method of Tietz, [29].

Statistical analysis

Results were expressed as mean ± standard deviation (SD) and test of significance was determined at p < 0.05. Laboratory data were analysed using one-way analysis of variance (ANOVA) in Statistical Product and Service Solutions (SPSS), version 18.
3. Results and Discussion

This study investigated the effect of crude methanol extract (CME) of *Zapoteca portoricensis* root and its fractions on PSA levels, ACP activities, the oxidative stress and antioxidant status of albino rats experimentally induced with benign prostatic hyperplasia (BPH).

<table>
<thead>
<tr>
<th>Table 1. Effect of crude methanol extract and fractions of <em>Zapoteca portoricensis</em> roots sample on Prostatic specific antigen (PSA) of BPH induced rats.</th>
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<tr>
<td><strong>Group</strong></td>
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<td>Group 9</td>
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* n = 5; Data represent mean ±SD (n = 5); Values with “*” are significantly different compared to BPH-control while values with letters of the alphabets ‘a,b,…n,’ as superscripts are significantly different compared to normal control (p < 0.05). Group 1 = Normal control (not BPH-induced and no treatment). Group 2 = BPH-control (BPH-induced but no treatment). Group 3 = Standard control (induced and treated with Avodart®). Groups 4 and 5 = Induced and treated with 200 and 400 mg/kg of CME, respectively. Groups 6 and 7 = Induced and treated with 200 and 400 mg/kg of MF,respectively. Groups 8 and 9 = Induced and treated with 200 and 400 mg/kg of EAF, respectively.

3.1. Effect of Crude Methanol Extract and Fractions of *Zapoteca Portoricensis* Roots Sample on Prostatic Specific Antigen (PSA) of BPH Induced Rats

The results obtained showed a significant (p < 0.05) decrease in PSA levels and ACP activities for all the test groups at different concentrations (Table 1). The decrease in PSA levels by the extract is comparable to that obtained from the standard drug-Dutasteride (Avodart). This is an indication of its protective effect against the development of BPH. The result agrees with a recent report that treatment of BPH-induced rats with ethanol extract of *Z. portoricensis* stem has positive outcome by reducing PSA levels and prostate weight in rats PSA is generally elevated in BPH and other Prostate disorders and is a reliable marker for BPH. A reduced PSA is associated with reduced prostate hyperplasia as a direct consequence of 5α reductase inhibition or anti-inflammatory action. Many plants associated with reduced BPH have been documented [31,32]. BPH is caused by an increase in dihydrotestosterone (DHT), a metabolite produced when testosterone is converted to DHT by an enzyme 5α reductase. Consequently, inhibitors of 5- alpha reductase which block the production of DHT will slow down the development of BPH.

<table>
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<tr>
<th>Table 2. Effect of crude methanol extract and fractions of <em>Zapoteca portoricensis</em> roots sample on oxidative stress and antioxidant indices of benign prostate hyperplasia (BPH)-induced rats.</th>
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<td><strong>Group</strong></td>
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<tr>
<td>Group 1</td>
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<table>
<thead>
<tr>
<th>Group</th>
<th>SOD Activity</th>
<th>CAT Activity</th>
<th>GSH Activity</th>
<th>PSA Activity</th>
<th>ACP Activity</th>
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</thead>
<tbody>
<tr>
<td>4</td>
<td>2.68 ± 0.21 b</td>
<td>70.48 ± 2.61 bc</td>
<td>1.53 ± 0.11 a</td>
<td>18.64 ± 1.92 a</td>
<td>0.43 ± 0.09 a</td>
</tr>
<tr>
<td>5</td>
<td>2.58 ± 0.25 a,b</td>
<td>71.64 ± 4.24 b,c,d,e</td>
<td>1.66 ± 0.14 a</td>
<td>20.46 ± 0.81 a</td>
<td>2.25 ± 1.81 a</td>
</tr>
<tr>
<td>6</td>
<td>2.54 ± 0.13 a,b</td>
<td>72.47 ± 3.10 b,c,d,e</td>
<td>1.67 ± 0.22 a</td>
<td>20.22 ± 1.70 a</td>
<td>0.47 ± 0.04 a</td>
</tr>
<tr>
<td>7</td>
<td>2.71 ± 0.25 b</td>
<td>74.44 ± 2.83 b,c,d,e</td>
<td>1.71 ± 0.27 a,b</td>
<td>57.09 ± 6.63 a</td>
<td>0.44 ± 0.04 a</td>
</tr>
<tr>
<td>8</td>
<td>2.18 ± 0.16 a,c</td>
<td>67.58 ± 6.38 a,b</td>
<td>1.48 ± 0.09 a</td>
<td>19.05 ± 1.17 a</td>
<td>0.30 ± 0.04 a</td>
</tr>
<tr>
<td>9</td>
<td>2.24 ± 0.51 a</td>
<td>72.06 ± 2.18 b,c,d,e</td>
<td>1.55 ± 0.05 a</td>
<td>19.85 ± 0.85 a</td>
<td>0.36 ± 0.08 a</td>
</tr>
</tbody>
</table>

n = 5; Data represent mean ±SD (n = 5); Values with “**” are significantly different compared to BPH-control while while values with letters of the alphabets ‘a,b,…n,’ as superscripts are significantly different compared to normal control (p < 0.05).

3.2. Effect of Crude Methanol Extract and Fractions of Zapoteca Portoricensis Roots Sample on Oxidative Stress and Antioxidant Indices of Benign Prostate Hyperplasia (BPH)-Induced Rats

A significant (p < 0.05) decrease in malondialdehyde concentration, a significant (p < 0.05) increase in superoxide dismutase activity and a non-significant (p > 0.05) increase in catalase activity and glutathione concentration was observed in all the test groups. However, there was no significant difference in vitamin C concentration in all the test groups (Table 2). These results are indicative of the oxidative stress/lipid peroxidation and antioxidant potentials of Zapoteca portoricensis root extracts and thus its protective effect against BPH. Oxidative stress (OS)/Lipid peroxidation is considered to be one of the mechanisms that trigger the chain of reactions involved in the development and progression of prostatic hyperplasia (BPH). This is characterized by the observed increase in oxidative stress/lipid peroxidation and decreased levels of antioxidants indices. The development of BPH, an age dependent disease, has been associated with increased lipid peroxidation, oxidative stress, inflammatory process and decreased levels of antioxidants [33–35]. In Lipid peroxidation Polyunsaturated fatty acids peroxides generate malondialdehyde (MDA) and 4-hydroxyalkanals upon decomposition. This process is an established mechanism of cellular injury, and is used as an indicator of oxidative stress. Superoxide dismutase (SOD) decomposes superoxide anion into hydrogen peroxide and oxygen at very high rates. Superoxide radical is involved in diverse physiological and pathophysiological processes. Catalase (CAT) is antioxidant enzymes ubiquitously present in aerobic cells and catalyses the decomposition of hydrogen peroxide to water and oxygen. High concentration of hydrogen peroxide is deleterious to cells, and its accumulation causes oxidation of cellular targets such as DNA, proteins and lipids, leading to mutagenesis and cell death [38]). Glutathione has antioxidant properties since the thiol group in its cysteine moiety is a reducing agent and can be reversibly oxidized and reduced. In cells, glutathione is maintained in its reduced form by glutathione reductase (GR), and in turn reduces other metabolites and enzymes, as well as react directly with oxidants. Vitamin C (ascorbate), a water-soluble vitamin involved in anti-oxidative processes in cellular metabolism, plays an important role in synthesis of collagen and nor epinephrine by maintaining the necessary enzymes in their active forms. It is also a co-factor for the hydroxylase involved in the hydroxylation of proline residues of collagen, which is vital to wound healing. It is probable that the various phytochemical constituents of the extracts, such as the flavonoids, saponins, phenols and terpenoids [41] are involved in scavenging free radicals from the tissues, thus reducing oxidative stress.

4. Conclusions

The extracts and fractions of the plant Zapoteca portoricensis root exhibited some protective effect against the development of BPH, in addition to oxidative stress/lipid peroxidation and antioxidant capacity as shown in the increased levels of SOD, CAT and GSH and decreased levels of PSA, ACP and MAD. These observations are indications of modulatory effects of Zapoteca portoricensis root extracts on induced benign prostate hyperplasia (BPH) in male albino rats and could be a source of new agent for managing BPH.

Conflicts of Interest: The authors declare no conflict of interest in this work.
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