In Vitro Effect of Pomegranate Extract on Trichomonas Tenax

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Abstract

The incidence of T. tenax in patients with acute ulcerative gingivitis has been demonstrated in several published reports about it. Metronidazole was known as the most effective drug for human trichomoniases, however, drug resistance and toxicity appeared. This study was designed to investigate the in vitro inhibitory activity of P. granatum ethanol extract on the growth and motility of Trichomonas tenax in comparison to metronidazole. Pomegranate ethanol extract group was divided into four groups with concentration of 12.5, 25, 50, and 100 mg/ml, respectively, metronidazole group and blank control. Metronidazole group was given 10 μg/ml of the drug. Each group had 4 wells with 125 μl T. tenax (2×10^4/ml). At 12 hr, 24 hr, 48 hr and 72 hr after drug treatment, the anti-T. tenax effect of pomegranate ethanol extract was tested by microscope counting method. The results showed that the better effect on anti-T. tenax 60% pomegranate ethanol extract group with concentrations of 12.5 mg/ml and 25 mg/ml showed higher anti-T. tenax (P<0.01). The ethanol extract of pomegranate granules has a remarkable effect on T. tenax, and among the groups, 60% ethanol extract shows the best anti-T. tenax activity.

Keywords: Trichomonas tenax; Punica granatum; Metronidazole; Herbal medication

Introduction

The human oral cavity is home to numerous microorganisms. Trichomonas tenax (Trichomonas buccalis) is a regular guest of human oral cavity microorganism [1]. It is an anaerobic species that lives as a commensal in the mouth of human oral cavity. It is frequently associated with pyogenic organisms in pus pockets or at the base of teeth. There are studies that relate to its prevalence in patients with Marginal Chronic Periodontitis [2]. Transmission is through saliva, droplet spray, and kissing or use of contaminated dishes and drinking water [3]. World widely, its prevalence in the mouth ranges from 4 to 53% [4,5].

The detection of T. tenax in the human oral cavity is an indication of hygiene poor oral, so that its incidence increases significantly in patients with periodontal problems, this being three to four times higher than in periodontal healthy subjects [6].

Since the organism is believed to enter the respiratory tract by aspiration from the oropharynx and then cause bronchopulmonary trichomoniasis, the importance of oral infections has been increased recently [7].

The development of drug resistance in human pathogens against commonly used treatment has necessitated a search for new therapeutic agents from other sources. Recently, there has been considerable interest in the use of plant materials as an alternative method to control pathogenic microorganisms [8,9], and many compounds of plant products have been shown to be specifically targeted against resistant pathogens [10].

Punica granatum, which belongs to the family of Puniceae, is commonly known as pomegranate, grenade, granats and punica apple [10]. Punica granatum has been used extensively as a traditional medicine in many countries [9] for the treatment of dysentery, diarrhoea, helminthiasis, acidosis, hemorrhage and respiratory pathologies [11,12]. In addition, P. granatum is reported to have antioxidant [9,10] anti-atherosclerotic [13,14], antibacterial [15,16] and antiviral [17] properties. The constituents of P. granatum include galacto-chins, delphinidin, cyanidin, gallic acid, ellagic acid, pelargonidin and sitosterol, which are very well known for their therapeutic properties [18]. Punica granatum peel is used to treat infections found in human sexual organs as well as mastitis, acne, folliculitis, pile, allergic dermatitis, typanitis, scalds, diarrhoea, dysentery and as an antioxidant. In addition, it is reported that the extracts of P. granatum have antimicrobial activity against Salmonella [18].

Pomegranate components have properties that could promote oral health, including reducing the risk of gingivitis. However, to date, no studies regarding the anti-T. tenax activity of P. granatum extract have been conducted. Therefore, the goal of this study is to evaluate the anti-T. tenax activity of the extracts of P. granatum peel in vitro.

Materials and Methods

Patients

The periodontiums of 51 patients were clinically examined, and diagnosis and classification of the periodontium was done according to the Periodontal Screening and Recording (PSR) I, in agreement with the Military Academy of Periodontology and Dental Association criteria. Twenty patients were diagnosed with gingivitis (EG1), 22 with periodontitis (EG2) and 9 presented a healthy periodontium (CG). The patients were also asked about the use of medications and systemic conditions which might predispose them to the development of periodontal disease.

Sample collection

Samples of saliva and dental biofilm/calculi were collected from all patients in the morning, before any oral hygiene. After determining the frontal mandibular area most affected by periodontal disease, PSR, dental biofilm/calculi samples were collected by scraping the area with sterile periodontal curettes. Unstimulated saliva sam-

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samples were collected as recommended by Navazesh. All samples were placed in sterile Petri dishes and diluted with saline at room temperature (25 to 28°C). Immediately after dilution, the samples were examined under a light microscope.

For the removal of dental calculus, as well as debris and plaque subgingival periodontal curettes used tarterctomos previously sterilized and, taking the material with the latter the bottom of the periodontal pockets of patients with PMC were also taken plaque samples tooth around the neck of the teeth of patients, employing previously sterile curettes. Dental calculus removed from the experimental group, was ground prior to planting and microscopic observation, using previously sterilized glass rods.

Transport of samples

The samples were placed in vials containing transport medium (sterile Ringer solution) and taken to the laboratory for further planting and microscopic observation.

Inoculation of the samples

Once the samples were taken to the laboratory, 0.1 ml was taken. Ringer’s solution containing the inoculum, using sterile 1 ml pipettes, inoculating the above-mentioned amount in the broth selective Kupferberg, used for the growth of T. tenax (“Kupferberg Trichomonas Broth.”Difco Laboratories, Detroit, Michigan, USA), to which were added 0.1 g of chloramphenicol to prevent the growth of bacteria and other microorganisms. Two plantings were made for each patient, an aerobically and anaerobically another, using the jug designed for this purpose (Gas Pak). Seeded culture media were taken to the oven at 37°C for 72 hours.

Microscopic observation

For the identification of T. tenax microscopic observations were made three times for each patient in order to determine what opportunities exist in the ability to view the scourge. These observations were made in the first instance on the same day of sampling, taking a drop of inoculum containing transport medium with the previously sterile platinum loop and placed onto a glass slide. The two remaining microscopic observations were performed at 72 hours of selective media incubated in an oven, taking a drop in the previously sterile platinum loop, both the stock that was planted under aerobic conditions as was shown in the anaerobic placing in each case on the surface of its respective blade slide.

In each of the cases described above, was placed on the drop of a slide coverslip microbial suspension, then the observation for the cool, using the light microscope (Leitz), focusing first with a low magnification lens (10 x) and then with higher magnification lenses (20 x 40 x).

Extraction of plant material

Preparation of the plant extract: Firstly fresh pomegranates (500 gram) were obtained (in order to prepare fresh extraction) from a public market. The peels of pomegranate were separated and oven dried at 33°C for 7 days. The dried peels were powdered in an electric grinder and stored in plastic bags for the next step. A 100 gm sample of powder was extracted using 200 ml methanol (99.9%) in an electric blender for 30 min. This suspension was filtered three times per day for 30 days. New methanol dissolvent was used each time. Then methanol was removed in a rotary evaporator to produce a dry powder. The final material was dissolved in methanol for obtaining concentrations of 4 mg/ml, 8 mg/ml and 12 mg/ml of dry plant powder [17-19].

Metronidazole

It was supplied as 500 mg tablets (Rhone Poulenc Rorer, France). Tablets were dissolved in distilled water, and then diluted in incubation medium to yield 12.5 μg/ml, 25 μg/ml and 50 μg/ml and 100 ug/ml [20].

Growth inhibition assay

The effect of P. granatum on the growth of the T. tenax trophozoites was studied as follows: 2x105 trophozoites were incubated in Selective Kupferberg (KTB) medium with drugs in different concentrations: p.
granatum and metronidazole (12.5 μg/ml, 25 mg/ml, 50 mg/ml and 100 mg/ml), for 12, 24, 48, and 72 h at 37°C. In addition, controls were included (cultures containing only the parasites) and submitted to the same procedure used for the experimental cultures.

Evaluation of the drug efficacy was done by:

1. Counting the number of trophozoites using the haemocytometer (Neubauer cell-counter chamber).
2. Calculation of the percent of inhibition of multiplication according to the equation:

   \[ \text{Percent inhibition of growth} = \frac{a - b}{a} \times 100 \]

   Where:
   
   a = Mean number of trophozoites in control tubes and
   b = Mean number of trophozoites in test tubes [21].
3. Calculation of the percent of motility of trophozoites which is the ratio of motile to total number of parasites counted per 10 High Power Field (HPF).
4. The Minimal Lethal Concentration (MLC) of p. granatum extract and metronidazole was determined.

Results

The observation in the cool of the sheets prepared allowed us to visualize where they were considered positive, the flagellate in some cases moving in a circular motion and other times not moving and isolated. Usually, the flagellar morphotypes of T. tenax can be seen with four flagella, but sometimes can be seen only with one or two flagella (Figures 1-4).

The present study was carried out to investigate the in vitro activity of Punica granatum peel (EEPDP) on the growth and motility of T. tenax, compared to the standard drug metronidazole. The results showed that the degree of growth inhibition was dependent upon the concentration of P. granatum and metronidazole.

Effect of P. granatum extracts in specified times and concentrations on Trichomonas tenax were assessed. The inhibitory effect of extract on Trichomonas was assessed by counting the live parasites 1, 2 and 24 hours after exposure with extracts.

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NTC = Non Treated Culture control

Table 1: Percentage of motility of T. tenax after exposure to various concentration of ethanol extract in comparison to normal control.

Table 2: Percentage of motility of T. tenax after exposure to various concentration of metronidazole in comparison to normal control.
Microscopic observation in the cool of the samples from dental calculus and subgingival plaque by using light microscopy is essential if you want to achieve faster discovery of *T. tenax*, being able to visualize usually with one or two flagella. However, provided the samples should be inoculated in the culture media most appropriate and allow for finding more sure of this species [28,29].

**Conclusion**

The incidence of *T. tenax* was higher in patients and this study suggests that EEPGP might be used as an antiparasitic agent in controlling oral *Trichomonas tenax* infections.

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**References**