Phytochemical Aspects, Cytotoxicity and Antimicrobial Activity of the Methanolic Extract of Tropical Fruit Pulps on Clinical Isolates of *Escherichia coli*

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Abstract: *Escherichia coli* is naturally present at the gastrointestinal tract of humans, but is also involved in diseases such as urinary tract infection. The current scenario of bacterial resistance to antimicrobials raises the demand for new drugs, and natural products represent interesting sources of bioactive compounds. Here we investigated the cytotoxicity and antimicrobial potential of 80% methanolic extracts of *Spondias tuberosa* (umbu), *Spondias purpurea* (seriguela), and *Theobroma grandiflorum* (cupuaçu) fruit juice pulps against clinical isolates of uropathogenic *E. coli*. Phytochemical aspects of the pulps were elucidated, and their antioxidant properties were analyzed. The minimal inhibitory concentration (MIC) of the extracts was of 500 µg/mL, and no toxicity was observed against BGM cells. Vitamin C and total carbohydrates were not in accordance to the levels determined in the legislation. We used HPLC to confirm the presence of flavonoids. To the best of our knowledge, this is the first report of the antimicrobial potential of these fruit pulps. Our data open doors for more studies with chromatographic fractions and isolated flavonoids from these pulps.

Keywords: plant extract; uropathogenic *E. coli*; antimicrobial; umbu; seriguela; cupuaçu.

1. Introduction

   Escherichia coli is a relevant Gram-negative commensal microorganism actively involved in colonizing the gastrointestinal tract of humans, precluding the entry of pathogenic species. However, *E. coli* may reach other body sites and behave as a pathogen, causing diseases such as urinary tract infection (UTI) [1-3]. UTI is a common bacterial infection, especially in women, and results in decreased quality of life because of the symptoms, which include dysuria, increased urinary frequency, suprapubic pain, and hematuria [2, 4, 5]. Uropathogenic *E. coli* (UPEC) strains are mostly detected in uncomplicated cystitis cases, but might also be found at the kidneys, bladder, and even at the blood on complex cases [5-7].

   The treatment of UTI is largely dependent on antimicrobial drugs, but bacterial resistance makes it technically difficult [2, 8]. Several genetically regulated mechanisms have been described in UPEC, including efflux pumps and β-lactamases [6, 8-10]. A recent WHO

Global Surveillance of Antimicrobial Resistance reported high rates of *E. coli* resistance to third-generation cephalosporins and fluoroquinolones, the standard choices for the treatment [11]. Incorrect prescription and use of antimicrobials are widely recognized as the main causes of bacterial resistance [5, 6]. Thus, beyond the education of prescribers and patients on this topic, there is an urgent need for new antimicrobial drugs. Plant-derived metabolites are promising compounds in the antimicrobial research context. Reports on resistance to natural products are rare, and many of them present broad-spectrum activity [12]. Roots, stems, and leaves of plants, which are not always edible, are commonly explored for phytochemical and pharmacological studies with microorganisms.

Our group has been prospecting antimicrobial extracts of edible parts of plants such as fruits, which have extensive culinary and cosmetic use but are far less investigated for biological properties such as antimicrobial, antioxidant, and wound healing. Here we report the cytotoxicity of the methanolic extracts of *Spondias tuberosa* (umbu), *Spondias purpurea* (siriguela), and *Theobroma grandiflorum* (cupuaçu) fruit juice pulps, and their antimicrobial activity on UPEC strains. Their pulps are widely used for preparing juices, ice-cream, and other food preparations that are economically important in several Brazilian cities, especially in the northeast (siriguela/umbu) and north (cupuaçu) regions [13]. Non-edible parts of these plants are popularly used on the treatment of inflammatory and infectious diseases, but no investigation has been conducted so far with the fruit pulps. Phytochemical analyses indicated the presence of tannins and flavonoids, and a comparative analysis of their antioxidant potential was conducted. To the best of our knowledge, such evidence are being provided for the first time, making our work even more relevant concerning the treatment of UTI.

2. Materials and Methods

2.1. Pulps samples.

The pulp samples used in this study belong to a brand widely sold in Brazil. Samples were purchased (10 units each) from local markets at Minas Gerais State and consisted of integral frozen pulps in vacuum-sealed plastic containers, which require adding water and culinary sugar prior to consumption, as recommended by the manufacturer. Products were of the same batch code. All pulps were stored at -20 °C until used, and defrosted in a refrigerator prior to the tests.

2.2. Biochemical assays with the whole pulps.

2.2.1. Carbohydrates.

Total carbohydrates were determined in each sample using the phenol-sulphuric method [14]. A calibration curve was prepared using glucose as a standard.

2.2.2. Vitamin C quantitative detection.

Vitamin C (ascorbic acid) content was determined in triplicate, as previously described [15]. A total of 35 mL of starch-sulphuric acid solution was added to 25 g of each pulp and mixed. The solution was titrated with 0.1M iodine solution while stirring until the first stable blue color was seen.

2.3. Preparation of methanolic extracts.

The extracts were prepared at home temperature with 80% methanol solution through magnetic stirring at maximum speed for 24 h. The solution was then rotavaporized to dryness under vacuum at 50 °C. The final product was weighed and stored at 4 °C until used.

2.4. Qualitative phytochemical screening.

The extracts were analyzed through classical methods [16]. Shinoda method for flavonoids was conducted as follows: the dried extracts were suspended in enough quantity of water and mixed with fragments of magnesium ribbon. HCl was dropped until the pink color was seen, indicating a positive result. For tannins, the suspended extracts were boiled in water for 10 minutes, filtrated in Whatman paper (n#10), and 0.1% ferric chloride was dropped on the filtrate. A brownish-green or a blue-black color indicated a positive result. For saponins, olive oil was added to the filtrate and agitated using a vortex at maximum speed. The formation of an emulsion showed a positive result.

2.5. Antioxidant activity.

The antioxidant activity of the extracts at 1 mg/mL was analyzed in duplicate by the β-carotene bleaching assay, as described [15]. The antioxidant chloroform solution was prepared with β-carotene, linoleic acid and Tween 40, and readings were taken in a spectrophotometer at 470 nm at 0, 15, 30, 45, and 60 min intervals.

2.6. High-Performance Liquid Chromatography coupled to diode array (HPLC-DAD) analysis.

We conducted HPLC analysis to detect polyphenols in the samples following the extract preparation method and equipment program as described by our group [17]. Aliquots of 20 µL of the extracts were injected in a C18 column (Shim-pack ODS) of a SPD 20A HPLC-DAD system (Shimadzu), and fractions were separated with gradient elution consisting in water and methanol at a flow rate of 0.5 mL/min. Polyphenols were detected at 254 nm.

2.7. Bacterial strains.

Clinical UPEC isolates were from the collection of the clinical laboratory of Pitágoras College. The strains were cultured in nutrient broth (Difco) overnight and were tested for identity confirmation using VITEK 2 system (version R04.02, BioMérieux SA, Marcy-l'Étoile, France) with identification cards, following the manufacturer's instructions.

2.8. Minimal inhibitory concentration (MIC) assay.

The MIC of the extracts was determined in untreated sterile 96-well polystyrene microtiter plates using an adapted CLSI protocol published by our group [18]. Bacterial cultures were prepared in Mueller Hinton broth (Difco, Becton Dickinson, USA) in 1 McFarland scale by adjusting the optical density to 1 at 600 nm wavelength. The wells received each of the extracts serially diluted, creating a final concentration of the bacterial inoculum equal to 0.5 McFarland scale and final concentrations of drugs ranging from 1 mg/mL to 7.8 µg/mL (final volume of 200 µL). MIC was established as the lowest concentration in which

resazurin (0.1g/L) staining was unaltered (no color modification from blue to pink) in all strains. The extracts were used as a negative control.

2.9. Cytotoxicity of the extracts.

The cytotoxic effect of the extracts was tested in triplicate using immortalized fibroblast-like BGM cells, as described previously [17]. The cells were treated with 20 μL of each extract in concentrations ranging from 500 to 7.81 μg/mL. The plates were incubated for 24 h. Cell viability was assessed by resazurin staining (0.1 g/L). The plates were then incubated for four h, and readings were taken in a fluorimetric microplate reader (λ_{ex}570 nm, λ_{em}590 nm). Untreated cells were used as the control group.

3. Results and Discussion

3.1. Biochemical analyzes, antimicrobial activity, and cytotoxicity.

Vitamin C was not detected in the pulps, and the total carbohydrate levels were lower than the ranges determined by the Brazilian laws (Table 1). All the extracts were effective against the clinical isolates of *E. coli*. The extracts were not cytotoxic to BGM cells: cell viability was not statistically different from the untreated control (data not shown).

Table 1. Antimicrobial and biochemical parameters of the fruit pulps.

Parameter	*T. grandiflorum*	*S. purpurea*	*S. tuberosa*
MIC$^{\alpha}$	500 μg/mL	500 μg/mL	500 μg/mL
Vitamin C	ND	ND	ND
Reference Value	X > 18 mg/100g	Δ	X > 12.9 mg/100g
Carbohydrates	3.74 g/100g	5 g/100g	ND
Reference Value	X > 6 g/100g	Δ	X > 2.4 g/100g

Δ – Reference values for *S. purpurea* are not available in the current legislation. ND: not detected. MIC: Minimal inhibitory concentration. α: Values for all bacterial strains.

3.2. Phytochemical profile of the pulps.

The qualitative tests indicated the presence of flavonoids and tannins in all samples, but saponins were not detected in any of the pulps. HPLC analysis confirmed the presence of polyphenols on the samples at 254 nm (Fig.1). The identification of the flavonoids is being conducted for further studies.
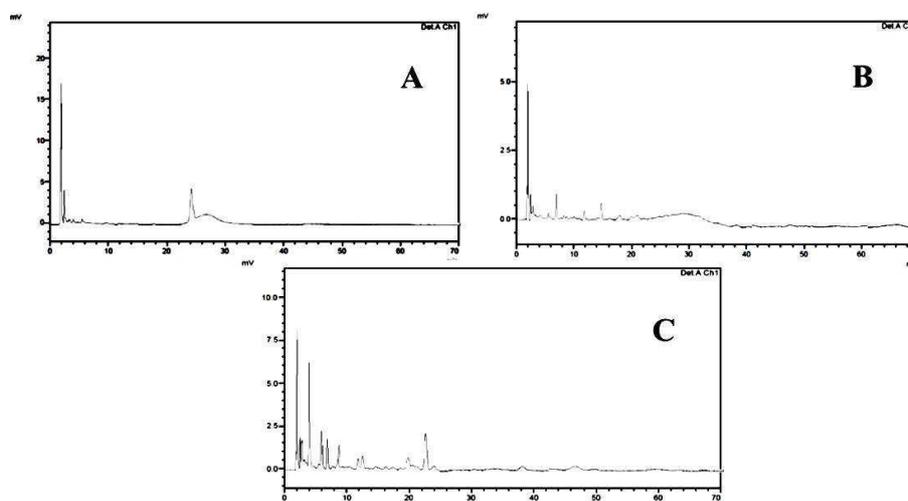

Figure 1. HPLC chromatograms of the extracts. A: *T. grandiflorum*, B: *S. purpurea*, C: *S. tuberosa*.

3.3. Antioxidant activity.

The extracts presented interesting antioxidant activity (Fig. 2), and all were statistically significant when compared to the untreated β-carotene solution used as control (p<0.05). *T. grandiflorum* presented 48.21% of antioxidant protection, whereas *S. tuberosa* and *S. purpurea* presented 67.64% and 68.17% of antioxidant protection, respectively. Differences between *S. tuberosa* and *S. purpurea* results were not statistically significant, although their protective effect was superior to *T. grandiflorum* (p>0.05).

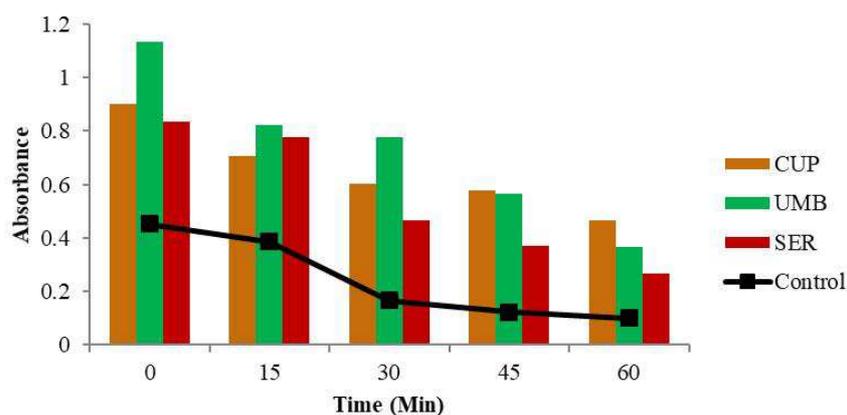**Figure 2.** Antioxidant potential of the pulp extracts. CUP: *T. ganrdifolium* sample (cupuaçu); UMB: *S. tuberosa* (umbu); SER: *S. purpurea* (seriguela).

3.4. Discussion.

S. tuberosa provides its fruits in dry seasons under considerable salt and hydric stress [19]. The pulp is rich in flavonoids, minerals, ascorbic acid, and carotenoids, and its exotic flavor reached from juices to ice-creams [19, 20]. *T. grandiflorum*, as the name suggests, is genetically close to *Theobroma cacao* (chocolate), albeit there are phytochemical differences [21]. A chocolate-like product can be prepared using *T. grandiflorum* seeds [22]. *T. grandiflorum* is mostly explored for juice, ice-cream, and butter, used in both food and cosmetic industries [21, 22]. *S. purpurea* is rich in phenolic molecules and is traditionally used for gastrointestinal disorders and parasitic infections [23]. Fruit pulps, however, remain poorly explored. To the best of our knowledge, this is the first study on the antimicrobial activity of *S. tuberosa*, *T. grandiflorum,* and *S. purpurea* fruit pulps hydromethanolic extract. The leaf and stem bark extracts of these plants are traditionally used for several conditions such as diarrhea and other intestinal disorders, diabetes, and infectious diseases, and some evidence has suggested their antimicrobial potential [19-25]. The juices of these fruits are rich in several phytochemical molecules, and thus, we hypothesized that the pulps could present antimicrobial potential as well.

The antimicrobial activity of the 80% methanolic extracts of the fruit pulps was assessed using broth microdilution, and a 500 μg/mL MIC was found for all extracts. Although some strains were susceptible to the extracts in lower concentrations (data not shown), MIC values were analyzed considering all the microorganisms. As the clinical isolates are from different patients with different backgrounds concerning the immune system and the use of antimicrobial drugs, some variability on MIC values amidst the isolates were expected. Previous studies of our group with fruit pulp-based extracts reached lower MIC values, such as for cashew apple and açaí pulps [15, 17]. The nutritional quality of the pulps is probably the

most plausible explanation for these differences and can be influenced by several factors such as soil nutrition, time of harvesting, and industrial processing steps. These factors, especially the last one, might also explain the unexpected absence of vitamin C, an important immunomodulatory and antioxidant nutrient.

One might ask why we did not perform minimal bactericidal concentration (MBC) or minimal biofilm eradication concentration (MBEC) tests. Previous findings from our group are in agreement with the observation of others that MIC and MBC values of crude phytoextracts are rarely equal or close, suggesting a bacteriostatic profile [26]. A possible explanation is that bacteria may use harmless phytomolecules as carbon and nitrogen sources to keep growing, in spite of the presence of antimicrobial secondary metabolites as tannins and flavonoids.

We conducted an MBC experiment, but the highest concentration of the extracts (1000 µg/mL) was ineffective in inhibiting bacterial growth on agar plates (data not shown). MBEC can be up to 1000 times higher than MIC values [15, 17]. Therefore, an antibiofilm activity of the crude extracts could not be expected. However, preliminary antimicrobial experiments conducted at our laboratory with chromatographic fractions, which can concentrate on structurally similar secondary metabolites, resulted in considerably lower MIC values [Alcântara et al., unpublished data]. These findings open doors for more studies with the pulps.

The extracts were efficient antioxidants. The presence of phytomolecules such as flavonoids and tannins helps to explain this result, as confirmed by HPLC (Fig.1). Carbohydrate levels were unexpectedly low in all pulps (Table 1), and at least two hypotheses may explain this result. First, the maturation level of the fruit used for manufacturing was inadequate. Old fruits present lower levels of carbohydrates, as well as total acidity [27]. Second, the pulps could have been prepared with an increased proportion of water, leading to a final product diluted to some extent. This may have affected vitamin C levels as well.

Most of the studies on antimicrobial properties of natural herbal products explore leaves and stem barks extracts, generally following ethnopharmacological data [12, 13]. Nevertheless, leaves and stem barks may have cytotoxic effects in fibroblast-like cells due to the presence of secondary metabolites. The extracts had no toxic effect on BGM cells in our tests. Such tests help to predict adverse reactions resulting from the use of the tested items as drugs. Exploring edible fruit as sources of antimicrobial compounds can be an interesting pathway, as the presence of toxic secondary metabolites is unlikely [23, 27].

4. Conclusions

The three pulp extracts were effective against clinical isolates of UPEC, and more studies with chromatographic fractions are being conducted in order to reach even lower MIC values. This study, nevertheless, is not without limitations. It is possible that artisanal pulps of fresh fruit could provide lower MIC values, as the manufacturing processes can impact on the level of bioactive molecules. Moreover, *in vivo* studies with animal models of urinary infection are necessary to investigate the antimicrobial potential of the extracts considering the hepatic metabolism and the immune system. These limitations, however, has no implication on our measuring of the antimicrobial activity of the extracts on UPEC strains.

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Conflicts of Interest

The authors declare no conflict of interest.

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