ANTIMICROBIAL POTENTIAL OF EXTRACTS FROM STEVIA REBAUDIANA LEAVES AGAINST BACTERIA OF IMPORTANCE IN DENTAL CARIES

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ABSTRACT
In recent years, the antimicrobial activity of Stevia rebaudiana Bertoni leaf extracts against a large number of microorganisms has been evaluated, but not its activity against microorganisms of importance in dental caries. The aim of this study was to evaluate the antibacterial activity of Stevia rebaudiana Bertoni leaf extracts against cariogenic bacteria. Extracts were obtained from the dried Stevia rebaudiana Bertoni leaves in hexane, methanol, ethanol, ethyl acetate and chloroform. The antimicrobial activity of the 5 extracts against 16 bacterial strains of the genera Streptococcus (n= 12) and Lactobacillus (n= 4) was evaluated by the well diffusion method. Minimal inhibitory concentrations (MIC) of the extracts in hexane, methanol, ethanol, ethyl acetate and chloroform on the 16 bacterial strains were respectively 30 mg/ml, 120 mg/ml, 120 mg/ml, 60 mg/ml and 60 mg/ml. The zones of inhibition present at the MIC were variable, ranging from 9 mm to 17.3 mm. Our results suggest that inhibition zones with a hexane extract are similar to those obtained with ethanol and methanol, but the minimal inhibitory concentration (30 mg/ml) is lower. For the four Lactobacillus species, the inhibition zones obtained between 12.3 and 17.3 mm were somewhat larger with ethyl acetate and chloroform extracts, suggesting they were the most susceptible microorganisms.

Keywords: antiInfective agents - stevia - dental caries

INTRODUCTION
In physiological conditions, most bacteria in the oral cavity are compatible with health. However, under certain circumstances of the oral environment and the condition and behavior of the host, these bacteria may reveal their potential virulence and cause disease¹. Dental caries is considered to be an alteration of the oral ecosystem with predominance of pathogenic flora. The main microorganisms associated to caries production are, in order of frequency: 1) Streptococcus mutans (mainly serotype c) and to a lesser extent S. sobrinus and S. gordonii and 2) Lactobacillus and Actinomyces species¹². Their participation in the dental caries generating process has led to the development and implementation of measures for prevention and/or control¹³. Among other strategies implemented to eliminate and/or control the microorganisms involved in dental caries, chemical and antimicrobial substances have been used. Antimicrobial biomolecules of


POTENCIAL ANTIMICROBIANO DE EXTRACTOS OBTENIDOS DE HOJAS DE STEVIA REBAUDIANA BERTONI SOBRE BACTERIAS DE IMPORTANCIA EN CARIES DENTAL

RESUMEN
En los últimos años se ha evaluado la actividad antimicrobiana de extractos obtenidos de hojas de Stevia rebaudiana Bertoni sobre un gran número de microorganismos. Sin embargo, no existen evaluaciones en microorganismos de importancia en caries dental. El objetivo de este estudio fue evaluar la actividad antibacteriana de extractos de hojas de Stevia rebaudiana Bertoni sobre cariogenic bacteria. Extractos fueron obtenidos de las hojas de Stevia rebaudiana Bertoni convertidas en polvillo en hexano, metanol, etanol, etanol acetato y cloroformo. La actividad antimicrobiana de los 5 extractos sobre 16 cepas bacterianas de los géneros Streptococcus (n= 12) y Lactobacillus (n= 4) fue evaluada por el método de difusión en pozo. Las concentraciones mínimas inhibitorias (CMI) de los extractos en hexano, metanol, etanol, acetato de etilo y cloroformo sobre las 16 cepas bacterianas fueron respectivamente 30 mg/ml, 120 mg/ml, 120 mg/ml, 60 mg/ml y 60 mg/ml. Las zonas de inhibición determinadas a las CMI fueron variables, variando entre 9 mm y 17.3 mm. Nuestros resultados sugieren que los halos de inhibición con un extracto de hexano son similares a los obtenidos con metanol y etanol, sin embargo, la CMI (30 mg/ml) es menor. Para las cuatro especies de Lactobacillus las zonas de inhibición obtenidas entre 12.3 y 17.3 mm fueron algo mayores con extractos de acetato de etilo y cloroformo, sugiriendo que fueron los microorganismos más susceptibles.

Palabras claves: agentes antiInfectivos - stevia - caries dental
natural origin are also currently being explored for use in adjuvant therapy. Plants have been widely used around the world as traditional remedies in the treatment of diseases. It is estimated that 66% to 75% of the world population currently uses plant-based medicines. The main aim of research into medicinal plants is to identify plants that possess pharmacological activity, and thus discover new substances or molecules having antimicrobial activity that could be transformed into medications by different chemical processes and used to control or prevent infectious diseases.

Research by Katsura et al. reports the bactericidal activity of bakuchiol against S. mutans, S. sanguinis, S. salivarius, S. sobrinus, L. acidophilus, L. casei among other microorganisms of the oral cavity. Another study reports the high inhibitory capacity of isopanduratin A against S. mutans, S. sobrinus, S. sanguinis and S. salivarius.

Stevia rebaudiana (Bertoni), one of the 407 species of the genus Stevia and one of the only two whose leaves contain a sweetening substance, has been used by native people as a sweetener and for medicinal purposes. It is a shrub originally from Paraguay and Brazil, which occasionally grows wild. It was described botanically in 1905 by naturalist Moisés Santiago Bertoni, as an herbaceous plant 40 to 80 cm tall of the family Compositae. In addition to being a non-caloric sweetener known in many parts of the world, it has hypoglycemi, antioxidant and antihypertensive action. Another advantage is that no toxic or genotoxic activity has been found in the complete extracts obtained from Stevia rebaudiana Bertoni leaves.

There are studies reporting antimicrobial activity of extracts obtained from Stevia rebaudiana on fungi and Gram-positive and Gram-negative bacteria. None of these studies includes an evaluation of Stevia rebaudiana extracts on the microorganisms involved in dental caries. The aim of this study was to evaluate the antibacterial activity of extracts in hexane, methanol, ethyl acetate and chloroform from Stevia rebaudiana Bertoni leaves against bacteria that are important in dental caries and oral health.

**MATERIALS AND METHODS**

**Bacteria**

The following 16 bacterial strains were used to evaluate the antimicrobial activity of the extracts obtained from Stevia rebaudiana: Streptococcus mutans ATCC 25175, Streptococcus mutans ATCC 31989, Streptococcus mitis 804 NCTC 3165, Streptococcus salivarius NCTC 8606, Lactobacillus acidophilus ATCC 4365, Streptococcus rattus FA 1 (G), Streptococcus mutans C67-1, Streptococcus cricetus AHT, Streptococcus mutans Ingbrit, Lactobacillus plantarum 748, Lactobacillus casei 475, Lactobacillus brevis, Streptococcus mutans 35FS3, Streptococcus mutans 35FS1, Streptococcus mutans 29FS2 and Streptococcus sobrinus CIO 428. In order to reconstitute and confirm their viability for the assays, the lyophilized strains were re-suspended in brain heart infusion (BHI) broth and incubated at 37 ºC for 48 hours in anaerobic conditions (H2:CO2:N2; 10:10:80) so that they would grow well, after which they were transferred to BHI agar for isolating.

**Extracts**

Dry Stevia rebaudiana leaves (Agricultura Colombiana) were powdered in a mill until 800 grams were collected. Extracts were obtained in ethanol, methanol, ethyl acetate, chloroform and hexane using the cold soaking technique. Sixty grams of powdered Stevia rebaudiana leaves were soaked in 250 ml of each solvent and placed in a mechanical shaker (Precision Reciprocal Shaking Bath, Precision Scientific; USA) at 150 rpm and 37ºC for 24 hours. Then they were filtered through filter paper (Sonderpapier Filtrak, GMBH; Germany) to remove any leaf residue. The extracts were immediately filtered again through 0.45 µm filters (Filter Bottle Top, Sigma Chemical Company; USA) to remove bacteria and ensure that they were free from contamination. They were concentrated at low pressure in a rotary evaporator (Buchi B-169, Vacuum-System; Germany) until they were dry. A microbiological test was performed on each concentrated extract to ensure that there was no bacterial contamination at the end of the process. Finally, the concentrated extracts were used to prepare 4 concentrations (15 mg/ml, 30 mg/ml, 60 mg/ml and 120 mg/ml).

**Microbiological Assays**

The antimicrobial activity of the extracts on bacteria was tested using the well diffusion method described by Dobner et al. A pure culture of each bacterium to be tested was used to prepare a suspension in trypticase soy broth and adjusted by tur-
bidimetry to 0.5 on the Mac Farland scale. From this suspension, 100 µl were taken and added to 20 ml of Mueller Hinton agar (liquid and sterile), mixed and poured into Petri dishes. The agar was allowed to solidify, and after waiting 20 - 30 minutes, a sterile Pasteur pipette was used to make 4 to 5 0.5 cm wells in the agar in each dish. Thirty µl of each of the 5 extracts was placed in the wells (this volume fits exactly into the wells without overflowing onto the surface of the culture medium). Vancomycin 180 µg/ml and Azithromycin 150 µg/ml were used as positive controls and each of the solvents as negative controls. The dishes were incubated at 37 ºC for 24-72 hours under anaerobic conditions (H2:CO2:N2; 10:10:80).

After the incubation, the presence or absence of inhibition zones was determined. Their diameter was measured and the minimal inhibitory concentrations (MIC: lowest concentration of the extract that produces an inhibition zone of at least 6 mm) determined. Each assay was performed in triplicate and the average reported in mm.

RESULTS

Final yield (dry weight -expressed as a percentage-that was finally obtained from processing 60 grams of S. rebaudiana leaves) for each extract in ethanol, methanol, ethyl acetate, chloroform and hexane was, respectively, 22%, 16%, 10%, 14 % and 0.9%. None of the 5 negative controls had antimicrobial activity. The positive controls (Vancomycin 180µg/ml and Azithromycin 150 µg/ml) had variable inhibitory activity on the 16 strains included in the study, with values ranging from 18 mm to 25 mm.

The MIC of the extracts in hexane, methanol, ethanol, ethyl acetate and chloroform on the 16 bacterial strains of the genera Streptococcus and Lactobacillus were, respectively 30 mg/ml, 120 mg/ml, 120 mg/ml, 60 mg/ml and 60 mg/ml.

Table 1 shows the results obtained with extracts in hexane, methanol, ethanol, ethyl acetate and chloroform on the 16 microorganisms studied. The inhibition zones of the MIC for the 5 extracts on the bacterial strains were variable, ranging from 9 mm to 17.3 mm. The hexane extract, which had the lowest MIC, produced the following inhibition zones, in mm: 10.0, 9.3, 9.0, 10.0, 12.0, 11.6, 10.3, 10.6, 10.3, 9.6, 12.6, 9.6, 13.6, 14.3, 14.0 and 13.3. The inhibition zones of the 5 extracts were slightly higher for the 4 Lactobacillus strains than for the 12 Streptococcus strains, as the lowest value for inhibition zone was 12.3 mm and the highest was 17.3 mm.

<table>
<thead>
<tr>
<th>Bacteria</th>
<th>Hexane 30 mg/ml</th>
<th>Methanol 120 mg/ml</th>
<th>Ethanol 120 mg/ml</th>
<th>Ethyl acetate 60 mg/ml</th>
<th>Chloroform 60 mg/ml</th>
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</thead>
<tbody>
<tr>
<td>S. mutans ATCC 25175</td>
<td>10.0</td>
<td>9.3</td>
<td>9.0</td>
<td>10.0</td>
<td>12.0</td>
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<td>S. mutans ATCC 31989</td>
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<td>9.0</td>
<td>10.3</td>
<td>10.3</td>
<td>11.6</td>
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<td>S. mutans C67-1</td>
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<td>9.3</td>
<td>10.0</td>
<td>10.6</td>
<td>12.3</td>
</tr>
<tr>
<td>S. mutans Ingbritt</td>
<td>10.0</td>
<td>9.0</td>
<td>9.3</td>
<td>11.0</td>
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<tr>
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<tr>
<td>S. mutans 35FS1</td>
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<td>9.3</td>
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<tr>
<td>S. mutans 29FS2</td>
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<td>10.0</td>
<td>11.0</td>
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<tr>
<td>S. sobrinus CIO 428</td>
<td>10.6</td>
<td>9.3</td>
<td>10.3</td>
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<td>8.6</td>
<td>9.3</td>
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<tr>
<td>S. salivarius NCTC 8606</td>
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<td>9.0</td>
<td>10.0</td>
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<td>S. rattus FA1(G)</td>
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<td>13.0</td>
<td>13.0</td>
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<td>9.0</td>
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<tr>
<td>L. brevis</td>
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</table>

The values of the inhibition zones at the minimal inhibitory concentration are expressed in mm.
DISCUSSION
New substances with pharmacological potential have been searched for and applied since ancient times. During the last twenty years, there has been a revival of interest in research into natural products, as they make up 50% of the drugs used clinically in developed countries, of which 25% come from higher plants.

This is the first study to determine the antimicrobial activity of 5 extracts obtained from *Stevia rebaudiana* leaves on 16 Gram-positive bacterial species of the genera *Streptococcus* and *Lactobacillus*, which are important for dental caries and oral health. Existing research to date has only proved the antimicrobial activity of extracts of *Stevia rebaudiana* leaves on fungi, rotavirus virus and very different bacteria from those evaluated in this study.

The results of this study match those of Tadhani et al., Ghosh et al. and Taware et al. regarding the antimicrobial activity of extracts of *Stevia rebaudiana* leaves on fungi, rotavirus virus and very different bacteria from those evaluated in this study.

Noticeable in the study by Abou-Arab et al. is the lack of antimicrobial activity of the extracts in hexane, water, ethyl acetate and chloroform against three Gram-positive microorganisms (*Listeria monocytogenes*, *S. aureus* and *Bacillus cereus*). In the study by Jayaraman et al., the extracts in water and chloroform obtained from *S. rebaudiana* leaves have no antimicrobial activity against *S. aureus*. The differences in the susceptibility of Gram-positive microorganisms to *Stevia rebaudiana* leaf extracts might be due to particularities of the genus and species and to differences in the organization of the cell wall.

The results of this study show antimicrobial activity of all the extracts at MICs between 30 mg/ml and 120 mg/ml. The ethanol and methanol extracts which have the same MIC (120 mg/ml) have similar inhibitory activity. The inhibition zones for the hexane extract are similar to those for ethanol and methanol, nevertheless, the MIC (30 mg/ml) is lower. The inhibition zones for the 4 *Lactobacillus* species are slightly higher in the ethyl acetate and chloroform extracts (60 mg/ml), suggesting that they are the most susceptible microorganisms. Their greater susceptibility to the different extracts may be due to their cell wall structure or to the presence of a substance or synergic mixture in the extracts that can penetrate the bacteria easily and produce greater damage.

Further studies are needed on the isolation, characterization and identification of substances present in the extracts, and to determine the antibacterial activity against a wide range of microorganisms that are important in other oral infections. These substances could subsequently be used in toothpastes, mouthwashes or other oral products with antibacterial potential. In addition, before any extracts or substances obtained from *S. rebaudiana* leaves are used therapeutically, it must be ensured that they are not toxic to eukaryotic cells.

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