

Effects of yogurt containing probiotics on salivary *Streptococcus mutans* and *Lactobacillus* sp. counts in children

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
30/12/2017	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
20/03/2018	Completed	<input type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
13/02/2018	Other	<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Probiotics are live bacteria and yeasts promoted as having various health benefits. Fermented foods contain several probiotic bacteria that provide benefits especially on a digestive level. Previous research has analyzed the effect of probiotics in the mouth and have shown that certain probiotic microorganisms could reduce the saliva levels of the bacteria Mutans streptococci and Lactobacilli sp., which play an important role in the development of tooth cavities, but this effect has not been widely assessed for two probiotics (*Lactobacillus casei* and *Lactobacillus acidophilus*) available in some commercial yogurt brands in Colombia ('Original' Yogurt, manufactured by Alpina S.A.). Therefore, the aim of this study is to assess the effects of *Lactobacillus casei* and *Lactobacillus acidophilus* available in yogurt on the saliva levels of Mutans streptococci and Lactobacilli sp. in a population of Colombian children aged 6-12.

Who can participate?

Healthy children aged 6-12

What does the study involve?

Participants are randomly allocated into two groups. Those in group A drink 'Original' yogurt containing probiotics (*Lactobacillus casei* and *Lactobacillus acidophilus*) and group B drink 'Yogo-Yogo' ordinary yogurt without probiotics. Every child receives 150 g of the assigned yogurt after their lunch for five week days during three months. The children are encouraged to maintain their normal oral hygiene habits and continue to brush their teeth daily.

What are the possible benefits and risks of participating?

The children who take part in this study are given the benefit of drinking a beverage with many nutrients which either had probiotics or did not. There are no risks because the yogurt used is available in all supermarkets in the city of Bucaramanga (Colombia).

Where is the study run from?

The charity restaurant "Maria Reina de las Misiones" (Colombia)

When is study starting and how long is it expected to run for?
November 2013 to November 2017

Who is funding the study?
Universidad Santo Tomas (Colombia)

Who is the main contact?
Prof. Martha J. Rodriguez
marthajuro@gmail.com

Contact information

Type(s)

Public

Contact name

Mr Jose Pieruccini

ORCID ID

<https://orcid.org/0000-0002-5098-5130>

Contact details

Calle 147 No. 25 - 30 Palmas del Campo Torre C Apto. 402
Floridablanca
Colombia
681004
+57 (0)3017850598
josepieruccini@ustabuca.edu.co

Type(s)

Scientific

Contact name

Prof Martha Rodríguez

ORCID ID

<https://orcid.org/0000-0001-7339-7836>

Contact details

Torres del Club, Carrera 21 # 36-83
Floridablanca
Colombia
681004
+57 (0)315 8169925
marthajuro@ustabuca.edu.co

Additional identifiers

Protocol serial number

GIFODONTP42014

Study information

Scientific Title

Effects of yogurt containing probiotics on salivary Streptococcus mutans and Lactobacillus sp. counts in Colombian children aged from 6 to 12 years: a randomised controlled clinical trial

Study objectives

Lactobacillus casei and Lactobacillus acidophilus probiotics which are available in a yogurt called 'Original' distributed by Alpina S.A. reduce salivary Streptococcus mutans and Lactobacillus sp. counts in six to twelve year old Colombian children.

Ethics approval required

Old ethics approval format

Ethics approval(s)

The Research Ethics Committee of the Dentistry School at the Universidad Santo Tomas in Bucaramanga (Colombia), 22/05/2014

Study design

Single-centre single-blind randomised placebo-controlled trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Healthy children not being under antibiotic therapy the two previous months to the start of the intervention

Interventions

The children were randomised by block randomisation into group A (intervention arm: 'Original' probiotics yogurt containing Lactobacillus casei and Lactobacillus acidophilus) or group B (placebo control arm: 'Yogo-Yogo' ordinary yogurt without probiotics). Every participant received 150g of the assigned intake after their lunch during weekdays only, for a three month period.

Microbiological examinations were performed at baseline and at the end of the intervention period. Samples of stimulated whole saliva were collected directly into sterile sample bottles. In the laboratory, samples were vortexed for 30 seconds and serially diluted from 10-1 to 10-3. Two inoculums for each dilution were plated on selective agar, Mitis Salivarius-Bacitracin agar for Streptococcus mutans and Rogosa agar for Lactobacillus sp. The agar plates were coded and anaerobically incubated at 37°. The number of CFU were identified based on their morphology, size, and color with the help of a digital colony counter 24 hours after incubation for Rogosa agar plates and 48 hours for Mitis Salivarius-Bacitracin agar plates. Biochemical identification tests were also used for identification of microorganisms. CFU of individual species were estimated by a single trained investigator.

Intervention Type

Supplement

Primary outcome(s)

Streptococcus mutans and Lactobacillus sp. Colony Forming Units (CFU) counts; microbiological examinations performed at baseline and at the end of the intervention period

Key secondary outcome(s)

Salivary pH measured using a calibrated digital pH-meter and buffering capacity at baseline and at the end of the intervention period

Completion date

24/11/2017

Eligibility

Key inclusion criteria

1. Healthy children
2. Both genders
3. Age 6-12 years old
4. Lactose tolerant

Participant type(s)

Healthy volunteer

Healthy volunteers allowed

No

Age group

Child

Lower age limit

6 years

Upper age limit

12 years

Sex

All

Key exclusion criteria

1. Dry mouth or xerostomy
2. Morbid obesity
3. Antibiotic therapy

Date of first enrolment

16/07/2017

Date of final enrolment

16/08/2017

Locations

Countries of recruitment

Colombia

Study participating centre

Comedor Infantil Maria Reina de las Misiones (a charity restaurant for children)
Bucaramanga
Colombia
681002

Sponsor information

Organisation

Universidad Santo Tomás

ROR

<https://ror.org/01x628269>

Funder(s)

Funder type

University/education

Funder Name

Universidad Santo Tomás

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Prof. Martha J. Rodriguez (marthajuro@gmail.com or marthajuro@ustabuca.edu.co).

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
	Participant information sheet				

