MICROBIAL POPULATIONS IN ATOLE AGRIO: A TRADITIONAL MEXICAN FERMENTED MAIZE BEVERAGE

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Introduction. Most of Mexican fermented foods are beverages, within these those elaborated from maize, are common because it is a staple grain. Traditional fermented foods are produced by natural fermentation, where no inoculum is added. In Southeastern Mexico, maize fermented products as pozol and atole agrio are usually consumed. Atole agrío is a fermented non-alcoholic beverage elaborated with young non nixtamalized maize. The preparation process is carried out in the household and in small-scale (1). The microbiota of this product is not well defined.

The aim of this study was to identify the main stages of elaboration of atole agrío from Villahermosa, Tabasco; to characterize the microbial population and its changes during fermentation process and to identify some lactic acid bacteria (LAB), enterobacteria and total coliforms isolated from raw material and from the different stages of elaboration of atole agrío.

Methods. Atole agrio was prepared in a traditional way as described in Valderrama (2). Total mesophilic bacteria, LAB, ALAB (amylolytic lactic acid bacteria), yeasts, molds, coliforms and *Enterobacteriaceae* were counted by the plate count technique from raw materials, during fermentation and from end products, as well as the pH. LAB, ALAB, enterobacteria and coliforms were isolated, purified and preserved in 20% glycerol. The isolated strains were identified by phenotypic methods (API and Vitek2).

Results. Atole agrío is made by two different methods of fermentation that include a solid and a liquid process. The main identified stages of process were: dehulling, cleaning, corn grinding and fermentation. To prepare atole agrío, a certain amount of fermented mass and water were heated to obtain a beverage of the desired thickness. Aditionally, sugar, honey or species can be added. The results showed higher levels of LAB (10⁸-10⁹ CFU/ml) than ALAB (10⁶-10⁷ CFU/ml) throughout the fermentation. After 12h, the level of total mesophilic bacteria was 7x10⁹, yeasts and molds 3x10⁹ and *Enterobacteriaceae* and coliforms 6x10⁵ CFU/ml. The pH decreased from 7.5 to 4.5. Growth of the microbial groups in the solid fermentation was similar as in the liquid one. At the end of both fermentations, atole agrío was boiled for 30 minutes

and the bacterial growth was, for all the groups less than the method's sensitivity (10 CFU/g).

Some strains of LAB and ALAB were identified throughout fermentation (Table 1). Species of enterobacteria identified from raw material and atole agrío were: Serratia marcescens, Enterobacter cloacae, Klebsiella pnuemoniae, Pseudomonas aeruginosa, Morganella morganii and Raoultella terrigena.

Table 1. Species of LAB and ALAB identified from atole agrío.

LAB strains	ATOLE AGRIO		SPECIES
	LIQUID	SOLID	
	(1)	(s)	
IL5ℓ2	ı		Lactobacillus delbrueckii
IIL6ť2	I		Lactococcus lactis ssp lactis
IS2ť3	s		Lactobacillus plantarum
ALAB strains			
IL2A2	I		Lactobacillus delbrueckii
IL4A2	I		Lactococcus lactis

Conclusions. Atole agrío is made by two different methods of fermentation that include a solid and a liquid process. Despite the presence of other microbial groups, LAB and ALAB may have an important role in this beverage. A deeper description of microbial diversity involved in atole agrío fermentation must be carried out.

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