



Biotecnia

ISSN: 1665-1456

Universidad de Sonora, División de Ciencias Biológicas y de la Salud

Alvarez-Ainza, Maritza Lizeth; García-Galaz, Alfonso; González-Ríos, Humberto; Moreno-Ibarra, Griselda Macrina; Torre-Martínez, Mayra De la; Zamora-Quñones, Karina Alejandra; Acedo-Félix, Evelia
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Biotecnia, vol. 23, núm. 1, 2021, pp. 21-27
Universidad de Sonora, División de Ciencias Biológicas y de la Salud

DOI: <https://doi.org/10.18633/biotecnia.v23i1.1221>

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Characterization and selection of native yeast isolated from natural fermentation for the production of the artisanal beverage bacanora

Caracterización y selección de levaduras nativas aisladas de la fermentación natural de la producción artesanal de la bebida bacanora

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ABSTRACT

Bacanora is a Mexican distilled beverage made from agave, similar to tequila and mezcal, produced with an artisanal process. Therefore, the organoleptic characteristics of different bacanora vary highly. The aim of this study was to characterize and select native yeast involved in this artisanal process of fermentation. An initial pre-selection based upon the phenotypic characteristics of 580 strains of yeast was done, and then based on fermentation characteristics with agave juice. After pre-selection, three *Saccharomyces cerevisiae* and three non-*Saccharomyces* were chosen for the fermentative characterization. The *S. cerevisiae* yeasts produced 25.51 ± 1.19 g/L of ethanol at the end of the fermentation process, with a volumetric ethanol productivity of 1.76 ± 0.17 g/L/h, while for the three non-*Saccharomyces* strains the parameters were 15.56 ± 1.68 g/L and 0.484 ± 0.06 g/L/h, respectively. In the mixed culture of three *S. cerevisiae*, all remained at the end, but one of them predominates, and of the non-*Saccharomyces* yeasts, *Torulaspora delbrueckii* was the predominant yeast, according with PFGE results. The results showed that the native yeasts are adapted to the agave juice used for bacanora production, and these yeasts would contribute to improve the ethanol yields and the organoleptic characteristics of bacanora.

Keywords: Bacanora; alcoholic fermentation; native yeast; *S. cerevisiae*; non-*Saccharomyces*.

RESUMEN

Bacanora es una bebida Mexicana derivada de agave, similar al tequila o mezcal producida de manera artesanal, por lo que sus características organolépticas varían altamente entre bacanoras. El objetivo de este estudio fue caracterizar y seleccionar levaduras nativas involucradas en el proceso artesanal de fermentación. Se realizó una pre-selección inicial basada en las características fenotípicas de 580 levaduras, y después una selección basada en sus características fermentativas en jugo de agave. Después de la preselección se eligieron tres *S. cerevisiae* y tres no-*Saccharomyces*. Al evaluar las características fermentativas el cultivo con *Saccharomyces*

cerevisiae produjo al final de la fermentación, 25.51 ± 1.19 g/L de etanol, con una productividad volumétrica de etanol de 1.76 ± 0.17 g/L/h, mientras que para el cultivo con las levaduras de no-*Saccharomyces* se obtuvieron 15.56 ± 1.68 g/L y 0.484 ± 0.06 g/L/h respectivamente. En el cultivo mixto de *S. cerevisiae*, todas se mantuvieron al final, siendo una la predominante y en las levaduras no-*Saccharomyces*, *Torulaspora delbrueckii* fue la predominante esto según el seguimiento con PFGE. Las levaduras nativas están adaptadas al jugo de agave utilizado para la producción del bacanora y además pueden contribuir a mejorar el proceso fermentativo, así como, las características organolépticas sensoriales de esta exquisita bebida.

Keywords: Bacanora; fermentación alcohólica; levaduras nativas; *S. cerevisiae*; no-*Saccharomyces*.

INTRODUCTION

Bacanora is a Mexican distilled beverage made from agave, similar to tequila and mezcal. The organoleptic characteristics of the beverage vary highly from producer to producer, due to the artisanal process used for production (Núñez, 2001; Aguirre-Rivera *et al.*, 2016). According to the Official Mexican Standards for bacanora, the drink must be made only from *Agave angustifolia* Haw and produced in the area of origin denomination for bacanora (AODB) (Alvarez-Ainza *et al.*, 2009; D.O.F., 2005). The process is similar for mezcal and tequila production. First, the agave core is cooked in ovens with mezquite-wood, to hydrolyze the inulin, then milled to obtain the must called "saite" and then fermented by native yeasts. Finally, fermented agave must be distilled and standardized to the particular touch of each producer (Moreno, 1998; Núñez, 2001; Yanez, 2003; De la Cruz, 2003).

From the Mexican distilled beverages from agave (tequila, mezcal and bacanora), only tequila batches keep a constant sensorial quality, since it is produced in large facilities with high technology. Mezcal and bacanora, on the other hand, have large variations in sensory quality, since they are produced using artisanal process and the spontaneous fermentation driven by the combined action of various species

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Recibido: 12 de febrero de 2020

Aceptado: 25 de agosto de 2020

of yeasts and bacteria. Furthermore, only a specific specie of agave is used for production of tequila, another kind for bacanora, and several different species for mezcal (D.O.F., 2004; D.O.F., 2005; Aguirre-Rivera *et al.*, 2017). In the tequila industry, most companies use spontaneous fermentation, in many cases due to the natural-fermentation policy of the company. Different *S. cerevisiae* strains utilized for wine, beer or whisky fabrication, and even bakery yeast, are used as started in some tequila industries, although this yeast is able to ferment the tequila must, this is not the best suited to carry out this process (Gschaedler *et al.*, 2004).

Starter cultures are required for keeping the typical quality of wine, however, pure cultures of yeasts from the same area are more effective (Querol *et al.*, 2003). It is believed that these yeasts, called "selected local yeast", are better because they are completely adapted to the climatic conditions, the raw material and they are partially responsible of the unique characteristics of the beverage (Mas *et al.*, 2002; Querol *et al.*, 2003). For example, in the study of Wondra and Berovic (2001), 29 native *S. cerevisiae* strains were evaluated during wine making, and found that only one of these strains produced an excellent wine, while 11 wines did not have good quality. The main desirable phenotypic characteristics to select a yeast culture, is resistance to adverse factors such as high concentrations of ethanol and low pH. In addition, other desirable characteristics are a short lag phase during fermentation, complete consumption of fermentable sugars and glycerol, and production of killer toxins. On the other hand, undesirable characteristics are production of H_2S , volatile acids, and some of intermediate metabolites like acetaldehyde and pyruvate (Mas *et al.*, 2002).

In addition, several studies have identified the importance of *S. cerevisiae* due to the role that it has on the modification of beverages. This could be depending on the inoculation process in the case of wine (Heart & Fleet, 1983), or on how this yeast can be found at different points of tequila production (Lachance, 1995).

In another study, Alvarez-Ainza *et al.* (2015) observed a high variability between the *S. cerevisiae* yeast isolated in every one of the municipalities during the natural fermentations process on the bacanora production. Moreover, this study showed how some non-*Saccharomyces* yeasts exhibited *Saccharomyces*-like characteristics, which might have an important participation on the fermentation process. In relation to this, Arellano *et al.* (2008) evaluated the kinetic parameters and the formation of volatile compounds from native *S. cerevisiae* and *Kloeckera sp.* strains, isolated from agave juice from tequila production, and observed that all *S. cerevisiae* strains as well as *Kloeckera* species had a similar behavior, however, between these species the production of aromatic compounds was different. Furthermore, Alvarez-Ainza *et al.* (2013) evaluated the main compounds of bacanora produced in the area with origin denomination, and found a great variability in the products; they recommend the use of a correct yeast culture as started for the fermentation process, so this variability could be corrected. In a Lopez-Alvarez

et al. (2016) study, the fermentation process in the elaboration of tequila was evaluated, carried out by comparing an ethanol yield in a *Kluyveromyces marxianus* native strain and in a bakery strain of *S. cerevisiae*, the results showed that the *K. marxianus* strain reached 96 % yield.

These studies support the idea that the *Saccharomyces* and non-*Saccharomyces* species play an important role in the fermentation process for tequila production, so they recommend the evaluation of mixed cultures. Therefore, the aim of this work was to select native yeast from a large group of *S. cerevisiae* and non-*Saccharomyces* strains isolated during the fermentation process of bacanora, and to evaluate the performance of the selected native yeast as started-mixed cultures for production of bacanora.

MATERIALS AND METHODS

Yeast Strain

Two hundred and five *S. cerevisiae* strains and 375 non-*Saccharomyces* yeast strains isolated from the fermentation tank of bacanora facilities, at different stages during the production of this beverage, were used (Collection of the laboratory of Microbiology in the Centro de Investigación en Alimentación y Desarrollo A. C., Hermosillo, Sonora, México). The yeasts were stored at $-70^{\circ}C$ in a YEPD medium (yeast extract 10 g/L, peptone 20 g/L and dextrose 20 g/L) with 30 % (v/v) glycerol.

Analysis of some phenotypical characteristics of yeast

Utilization of 1 % (w/v) glucose, fructose and inulin was assessed according to García-Galaz *et al.* (2004). Growth at 37 and 42° C was evaluated in PDA (Potato Dextrose Agar, Difco) plates acidified with tartaric acid 10 % (w/v) and incubated 48 h (D.O.F., 2004). Tolerance to ethanol was evaluated according to Lachance (1995), using 3, 5, 8, 10, 15 and 20 % (v/v) ethanol. Growth at different pH was evaluated on PDA plates, the initial pH was adjusted with tartaric acid in a pH range from 2.5 to 7.5, in 0.5 increments, and incubated at 30° C for 48 h. H_2S production was evaluated on Bismuth Sulfite Agar (Difco) plates and incubating for 5 days at 30° C (Caridi *et al.*, 2002). Killer factor was also evaluated using the killer-sensitive strain CECT 1018 and the Killer strain CECT 1891 (Caridi *et al.*, 2002) as references. Yeasts were cultivated in YEPD broth 1.8 % (w/v), with fructose instead of glucose, absorbance at 600 nm was followed until stationary phase and the length of the lag phase was determined.

Evaluation of mixed cultures as starter for bacanora production

Juice from *Agave angustifolia* Haw. The juice from agave was obtained from the saite of a producer; the saite was mixed with water and all the juice was filtered and collected. The final sugar concentration was adjusted to 12° Brix (55 ± 3 g/L reducing sugars) and stored at $-10^{\circ}C$.

Pre-inoculum and inoculum. The selected *S. cerevisiae* and non-*Saccharomyces* (*Torulaspora delbrucekii*, *Pichia membranefaciens* and *Kluyveromyces marxianus*) strains were

evaluated. Each strain was inoculated in diluted fermentation medium, from cultures of each yeast stored at -70°C in a YEPD medium, to obtain the pre-inoculum and the inoculum, the diluted agave juice (1:2 from an original of 12° Brix) was incubated for 12 h at 30°C in 500-mL Erlenmeyer flasks with 100 mL of agave juice, with low stirring (200 rpm). Two inoculums were used, one with three strains of *S. cerevisiae* (*S. cerevisiae* key 46, *S. cerevisiae* 591 and *S. cerevisiae* 1101) and another with three species of non-*Saccharomyces* (*Torulaspora delbrueckii* key 1227, *Pichia membranefaciens* key 807 and *Kluyveromyces marxianus* key 251).

Fermentation. Juice from *Agave angustifolia* Haw was sterilized in the fermentation tank (121°C , 15 min), the sugar concentration was adjusted at 12° Brix. The agave juice was supplemented with ammonium sulfate (1 g/L). Fermentations were carried in 3 L bioreactor (Applikon, The Netherlands) at 30°C and 200 rpm, using 2.5 L of the agave juice; air was not supplemented. Yeast was inoculated to an initial concentration of 20,000,000 cells/mL. Two fermentations were performed for each culture. The fermentation course was followed for 24 h and sampling was performed every hour during the first 12 h and each 2 h the remaining 12 h.

Analytical methods

Biomass concentration was estimated by total count using a Neubauer chamber or dry weight. Total reducing sugar was determined in previously hydrolyzed ferment yeast-free must by DNS method (Miller, 1959). Viability was estimated after dying cells with methylene blue (Díaz-Montañón et al., 2008; Segura-García et al., 2015). Ethanol concentration was determined using an enzymatic method measurement in an YSI biochemical analyzer 2700 (Yellow Spring Instrument). Karyotype technique was determined by pulsed clamp electrophoresis (PFGE) according to Alvarez-Ainza et al., (2015) with some modifications, using the zymolase-20T enzyme at the first step of the process to degrade the cell wall (25 µg/mL, Seikaguru Corporation). *S. cerevisiae* YPH80 was used as molecular weight (Molecular Weight log Biolabs Inc.). The gel was stained with ethidium bromide and photograph with a GelDoc camera (BioRad).

Statistics analysis

The phenotypic characteristics analysis was conducted by BioNumerics software (Applied Maths 1998-2002), using UPGMA (Unweighted Pair Group Method with Arithmetic Mean) and the DICE coefficient to obtain a dendrogram. The specific growth rate (μ), substrate consumed rate (r_s), ethanol production rate (r_p), biomass and ethanol yields ($Y_{x/s}$, $Y_{p/s}$) as well as a maximal biomass production (X_{max}) kinetic parameters were obtained by fitting a three-parameter logistic model for the fermentation with *Saccharomyces*, and the Gompertz model for the fermentation with non-*Saccharomyces*, based on the highest predictive fit obtained (R^2), using the statistical package NCSS 2007 (Kaysville, Utah, USA). The three-parameter logistic model states that: $Y = A / (1 + B \cdot \exp(-CT))$, where Y is the number of organisms in a given time; EXP is

the exponential function; A is the asymptote; B denotes the displacement in the X axis; C is the growth rate, and T is time. The Gompertz model states that: $Y = A \cdot \exp(-\exp(-B(T-C)))$, where Y is the response variable; EXP the exponential function; A, B, and C are the parameters of equation, and T is time.

RESULTS AND DISCUSSION

Analysis of some phenotypical characteristics of yeast

All *S. cerevisiae* strains fermented glucose and fructose, but only 43 % (88 strains) fermented inulin. Of the non-*Saccharomyces* yeast, 8.2 % did not ferment glucose, 7.4 % were unable to ferment fructose and 57.3 % could use inulin. Of the *S. cerevisiae* strains, 94 % were able to grow at 37°C and 64 % at 42°C , and for the non-*Saccharomyces* yeast 97.4 % were able to grow at 37°C and 95.8 % at 42°C . All yeasts were able to grow and resist ethanol concentrations in the range of 3-15 %, and 95 % of *S. cerevisiae* yeast, as well as 95.7 % of the non-*Saccharomyces* strains, were able to resist 20 % ethanol. Only 8 *S. cerevisiae* and 170 non-*Saccharomyces* strains were unable to grow at pH 2.5. Regarding H₂S production, 17 % of *S. cerevisiae* and 16.5 % of non-*Saccharomyces* yeast strains resulted positive. The killer phenotype was present only in three *Saccharomyces* yeasts, 14 were killer-sensible and 257 were killer-neutral.

The individual carbohydrate fermentation was performed in order to assess the ability of yeast strains to use these sugars individually. In this study, glucose, fructose and inulin, were assessed, which in theory are the agave juice sugars for the bacanora production (Aguirre-Rivera, et al., 2017); when cooked, agave “piñas” hydrolyze the carbohydrate reserve, those are the predominating with other fructans identified as new fructans or agavins (Mancilla-Margalli and Lopez, 2006). The bacanora process is done by a thermal treatment, however during the cooking process, all the fructans present in the “piñas” cannot be hydrolyzed, therefore, we also assessed whether these yeasts strains ferment inulin. In this study, it is noted that not all the non-*Saccharomyces* used glucose with a fermentative process, it is well known that this species uses most the oxidative metabolism (Heard and Fleet, 1985; Ciani et al., 2016a; Ciani et al., 2016b). The yeast tolerance or ability to grow at 37 and 42°C was important because in the region where the beverage is produced, temperatures of up to 42°C are reached in summer, based on the reports for tequila and whisky fermentation (Díaz-Montañón et al., 2008; Walker and Hill, 2016). Optimum temperature for most yeast is 30 - 35°C , but this temperature may increase by 2 or 3 degrees during the process, and in summer, it has been observed that during tequila fermentation temperatures up to 40°C are reached (Díaz-Montañón et al., 2008). The yeast ability for ethanol tolerance was performed by the recommendations of Lachance (1995). In that work, most of the *Saccharomyces* strains resisted up to 11 % ethanol, and in the present work almost all yeasts resist 20 %. Moreover, in fermented musts of cane and agave juice between 4 and 6.5 % ethanol are reached, so the native strains involved in the process of developing bacanora have this great quality (Heard and Fleet, 1985). The

pH of the "saite" is between 5 and 6, in some cases even lower (Fraizer and Westhoff, 1998), so in this study the survival of yeast at different pH was determined. Moreover, this study also evaluated the H_2S production on each of the strains because it is non-desirable (Mas et al., 2002; Caridi et al., 2005). Some bacanora-producers state that the "saite" sulfur aroma (rotten eggs) during the fermentation generates economic losses, since the batch is discarded (personal communication with producers).

The killer phenotype was present only in three *S. cerevisiae* strains, 14 *S. cerevisiae* were killer-sensitive and the rest of the *S. cerevisiae* and all the non-*Saccharomyces* (257 strains) were killer-neutral. Figure 1 shows the corresponding dendrogram analysis result of the phenotypic characteristics, and it shows the suitable characteristics recommended by Alvarez-Ainza et al. (2009) for bacanora production. Cluster 8 has the suitable characteristics recommended and it is constituted by 40 yeasts, 30 of them are *S. cerevisiae*, and 10 are non-*Saccharomyces* strains. Killer phenotype might represent an antagonist mechanism between yeasts during spontaneous fermentation process (Romano et al., 2003). The native populations might be a relatively high number of killer-resistant strains to the toxin and low presence of strains with killer-killer phenotype. Moreover, killer native populations in each area could put in interrogative the inoculation of a killer-sensitive strain selected to observe its best oenological characteristics. It is then recommended to select strains that show the killer-neutro o killer-killer phenotype, because it favors when are used as starter culture (Rodríguez et al., 1998).

Figure 1 shows the results of the analysis performed by the BioNumerics program, where the coefficient Dice and UPGMA were used to obtain a dendrogram, where a group or cluster with the suitable characteristics recommended by Alvarez-Ainza et al. (2009) for the bacanora production, can be observed. This group is cluster 8, constituted by 40 yeasts, 30 of *S. cerevisiae* and 10 of non-*Saccharomyces* strains. After the analysis of the phenotypic characteristics, the growth curves analyses were done. These analyses showed that the latency phase was diverse in the different kinds of yeast, where the non-*Saccharomyces* show latencies phases of up to 5 hours and the *S. cerevisiae* strains show no more than 2.5 h. These results help to determinate the yeasts that are adapted to the medium with fructose, fermentable carbohydrate present in the agave juice (data not shown).

Later, three non-*Saccharomyces* yeasts with a latency phase under 2 h and different velocity of growth, were chosen for evaluation in mixed cultures. Likewise, 3 different species of non-*Saccharomyces* were chosen to be evaluated in mixed cultures: *Kluyveromyces marxianus*, *Pichia membranefaciens* and *Toluraspora delbrueckii*.

Evaluation of mixed cultures as starter for bacanora production

The time course of the *Agave angustifolia* Haw juice fermentation with *Saccharomyces* or non-*Saccharomyces* strains mixed cultures is shown on Figure 2, and the

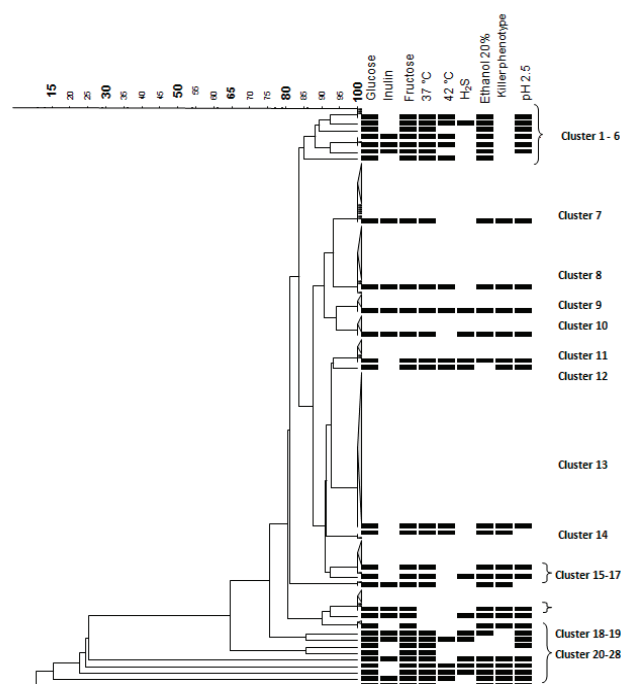


Figure 1. Phenotypic characteristics dendrogram of *S. cerevisiae* and non-*Saccharomyces* yeasts, obtained by the BioNumerics program using the coefficient Dice and UPGMA.

Figura 1. Dendrograma de las características fenotípicas de levaduras *S. cerevisiae* y no-*Saccharomyces* obtenido por el programa BioNumerics, utilizando el coeficiente Dice y UPGMA.

kinetic parameters on Table 1, where the P-value of the statistical analysis carried out with variance of kinetic parameters of the culture obtained using a logistic (*S. cerevisiae* culture) or Gompers model (non-*Saccharomyces* culture). The *Saccharomyces* mixed culture grew faster, and the maximal biomass had also greater numbers than the non-*Saccharomyces*, which yielded a maximal biomass of 8.27 ± 0.24 g/L, with reducing sugars depleted between 18-20 h. On the other hand, the mixed non-*Saccharomyces* strains culture yielded a maximum biomass of 3.87 ± 1.14 g/L and reducing sugars were not depleted.

The *Saccharomyces* mixed culture (*S. cerevisiae* key 46, *S. cerevisiae* 591 and *S. cerevisiae* 1101) was more efficient than non-*Saccharomyces* (*Torulaspora delbrueckii* key 1227, *Pichia membranefaciens* key 807 and *Kluyveromyces marxianus* key 251) to produce ethanol. The first produced up to 25.51 ± 1.19 g/L with a volumetric productivity of 1.76 ± 0.17 g/L/h, while non-*Saccharomyces* strains produced 15.56 ± 1.68 g/L of ethanol with a volumetric productivity of 0.484 ± 0.06 g/L/h. The kinetic parameter evaluated in the fermentations on this study, indicates the degree of adaptation of the microorganisms to the culture. The yeasts in mixed cultures were able to grow and achieve transformation of sugars into ethanol; it should be noted here that in the case of mixed cultures, there is also competition between the inoculated strains and the best adapted will be the predominant.

Similar results were reported by Arellano et al. (2008),

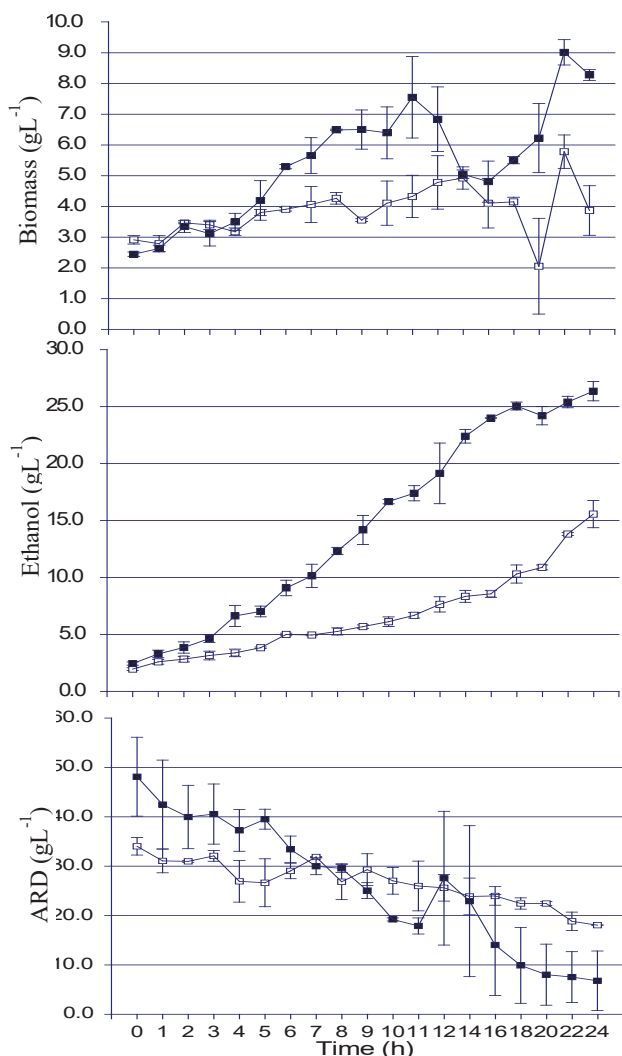


Figure 2. Fermentation kinetic profiles of cultures with *S. cerevisiae* (□) and non-*Saccharomyces* (■). ARD: reduction sugar concentration profile; Ethanol: ethanol concentration profile; Biomass: biomass concentration profile (Mean \pm standard error).

Figura 2. Perfil cinético de la fermentación de cultivos de *S. cerevisiae* (□) y no-*Saccharomyces* (■). ARD: perfil de reducción de la concentración de azúcares; Etanol: perfil de la concentración de etanol; Biomasa: perfil de la concentración de biomasa. (Media \pm error estándar).

they determined the kinetic parameters of wild yeast isolated from agave juice during tequila fermentation. In order to do this, they inoculated *S. cerevisiae* and *Kloeckera sp* isolates as monocultures in agave juice, and their results were in alcohol produced from 24 to 32 g/L being the *S. cerevisiae* specie the best alcohol producer. Similarly, Díaz-Montaña *et al.* (2008) study with *Agave tequilana* Weber juice, shows *S. cerevisiae* and *Kloeckera* with results between 21-43 g/L the ethanol produced at the end.

Di Serio *et al.* (2003) evaluated bakery yeast on YEPD medium, and other medium made with molasses and different salts, reporting different results with respect to this study. Values reported by Di serio *et al.* (2003) on the maximum growth rate (1.26 ± 0.11 per h) and biomass yield per sub-

Table 1. Kinetics parameters of the fermentation of agave juice inoculated with mixed cultures of *S. cerevisiae* and non-*Saccharomyces* strains.

Tabla 1. Parámetros cinéticos en la fermentación de jugos de agave inoculados con cultivos mixtos de cepas de *Saccharomyces cerevisiae* y no-*Saccharomyces*.

Parameter	<i>S. cerevisiae</i> mixed culture	Non- <i>Saccharomyces</i> mixed culture	P value
μ (per h)	0.29 ± 0.09	0.24 ± 0.06	0.6418
rs (g/L/h)	3.84 ± 0.12	1.03 ± 0.22	0.0008*
rp (g/L/h)	1.76 ± 0.17	0.48 ± 0.03	0.0000*
Yx/s (g/g)	0.07 ± 0.02	0.24 ± 0.03	0.0733
Yp/s (g/g)	0.45 ± 0.06	0.47 ± 0.03	0.8584
Biomass (g/L)	8.27 ± 0.24	3.86 ± 1.14	0.0331*
Total Alcohol (g/L)	25.5 ± 1.19	15.56 ± 1.68	0.0179*
Fermentation time (h)	18	>24	

μ : specific growth rate; rs , substrate consumed rate; rp , ethanol production rate; Yx/s , biomass yield; Yp/s , ethanol yield. Each value represents the average \pm standard deviation of duplicate determinations of two fermentations. * Statistical difference: $P < 0.05$.

strate consumption (0.723 ± 0.069 g/g), are higher than our results. This means that the YEPD medium and the molasses are more complete than the agave juice, or that the agave juice has inhibitors that affect the yeast, like some maillard compounds produced in the agave cooking, such as furfural. On the other hand, the results obtained are encouraging because the Bacanora producer obtained less than 20 g/L of ethanol in their fermentations, and in longer time with mixed *S. cerevisiae* cultures the production was 27.5 % higher.

The karyotypes of the yeast found at the end of the fermentation with mixed cultures (Figure 3), show that using *S. cerevisiae*, the strain with the key 1101 predominated at the end, but the other two also developed. In the non-*Saccharomyces*-mixed culture, *Torulaspora delbrueckii* predominated, and some *Kluyveromyces marxianus* strains were occasionally found, while *Pichia membranefaciens* was not found at the end, probably because this yeast is most sensible to alcohol or other products generated in the process. The yeast of *Saccharomyces* and non-*Saccharomyces* species are often found in alcoholic fermentations, for the production of wine, whiskey and other agave beverages (Esteve-Zarzoso *et al.*, 2001; Romano *et al.*, 2003; Fiore *et al.*, 2005; Díaz-Montaña, 2008; Fleet, 2008; Walker and Hill, 2016). The participation of the adequate *Saccharomyces* yeast has an important impact in ethanol production during fermentation, while the non-*Saccharomyces* yeasts in the final aroma; *Torulaspora delbrueckii* can be used to modulate sensory characteristics, such as acidity (Bely *et al.*, 2008). Also, the use of *S. cerevisiae* is recommended to prevent native yeast growth on must, and the reduction of fermentation time (García *et al.*, 2017). García *et al.* (2017) found that sequential cultures have produced more different wines providing organoleptic properties associated with the non-*Saccharomyces* strains used. Nolasco-Cancino *et al.* (2018) evaluated *S. cerevisiae* and non-*Saccharomyces* yeasts in single and mixed cultures in maguey juice (*Agave angustifolia*) used for mezcal production; the authors con-

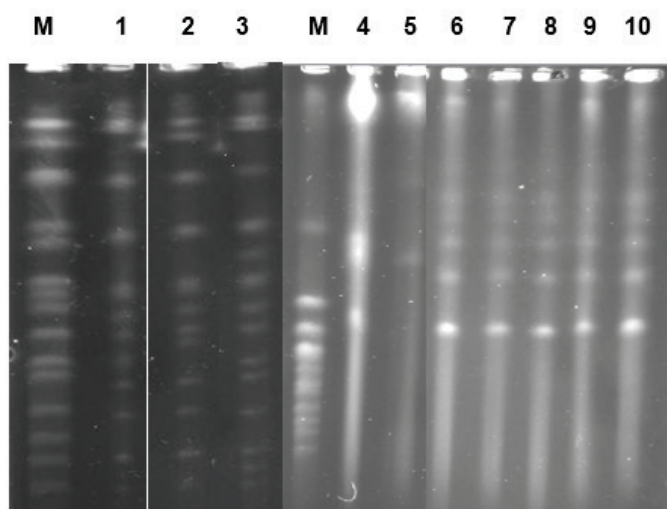


Figure 3. Karyotypes of yeasts found at the end of each culture probes fermentation.

M: molecular weight of *S. cerevisiae* YPH80; 1-3, *S. cerevisiae* strains isolated at the end of the fermentation; 4, *K. marxianus* control; 5, *P. membranefaciens* control; 6, *T. delbrueckii* control; 7-10, strains isolated at the end of the fermentation from the no-*Saccharomyces* culture.

Figura 3. Cariotipos de las levaduras encontradas al final de la fermentación de cada uno de los cultivos probados.

cluded that these cultures revealed their possible influence on volatile compound production, suggesting that the yeasts combination could contribute to improve the organoleptic characteristics of artisanal mezcal.

CONCLUSIONS

The native *S. cerevisiae* yeast isolated and selected from the *Agave angustifolia* Haw juice fermentation, can be used as starter mixed culture and the concentration of ethanol increases compared to the spontaneous fermentation in the production facility. Since *Torulaspora delbrueckii* ferments agave juice and predominated along the process in our study, it could be used together with *S. cerevisiae* to improve the fermentative process and enhance the sensory characteristics of bacanora but further studies are required.

ACKNOWLEDGEMENTS

The authors would like to thank and devote this manuscript to Evelia Acedo-Felix† who made this work possible. The authors also thank to the producers involves in the Bacanora production, and Gutiérrez-Franco, L. E. by proofreading.

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