Lactic acid Production by a Mixed Culture of Lactic Bacteria Based on Low Value Dates Syrup and Their Metabolic Uses

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Abstract

The common varieties of dates represent about 30% of national production. They are generally reserved for cattle feeding and sometimes thrown into the wild. Among these varieties, palm dates very rich in carbohydrates (glucose, fructose and sucrose) and contains practically most of minerals (Ca, Fe, K, Na, etc.). The use of fermentation technology is necessary to save the genetic heritage and enhances by-product of dates in various bio-products.

Lactic acid is a natural organic acid. The fermentation substrates most commonly used are glucose and fructose. The lactic acid has a major importance in food and pharmaceutical industries. It is used also in drink industries, medicines etc. Algeria till today continues to import this organic acid.

The objective of this study is the use of dates syrup as a substrate for the production of lactic acid, this study allowed us to have interesting results. Mixed culture of *Streptococcus thermophilus* and *Lactobacillus bulgaricus*, grown in batch fermentation. Two culture mediums were tested, the un-enriched one and the one enriched with Tween80 (1 g/l), MgSO4 (1 g/l), MnSO4 (1.5 g/l), from the results obtained by fermentation trials, the enriched medium seems best suited with a yield of lactic acid 49 g/l, where comes from its economical and commercial interest.

Keywords: Lactic acid; *Streptococcus thermophilus*; *Lactobacillus bulgaricus*; Date syrup; Fermentation; Metabolic rate

Introduction

Phoenicians heritage spreads on more than 85,000 hectares and reckons with more than 13 million palm trees, producing an average of 400,000 tons and allowing Algeria to the 5th place among the world producers of dates [1].

The Phoenicians Algerian sector provides date varieties of high quality and excellent commercial value. Other varieties are less appreciated, less commercialized and less competitive, in return they are very rich in nutrients, sugars such as: glucose, fructose and sucrose, vitamins and minerals [2,3]. Subsequently, they provide a very high energy. The use of transformation technology shows that it’s more necessary to use low-value dates for livestock feed [4].

Before adding the dates have subject of many tests in other contents. They are exploited as fermentation substrate for various metabolites production such as citric acid, ethanol and even biomass production (yeast bread) [5,6].

Lactic acid or its derivatives are indeed widely used by the pharmaceutical industry, food processing and chemical industries for the synthesis of varnishes and plastics [7]. The biotechnology evolution and the integration of microbiology and biochemistry enabled to master fermentation techniques and processes by involving an appropriate substance for lactic acid production from molasses, carob and glucose [8].

The objective of this work consists to use a second quality date’s syrup as a fermentation culture medium in the purpose of producing lactic acid by a mixed culture of *Streptococcus thermophilus* and *Lactobacillus bulgaricus* in different experimental conditions.

Materials and Methods

Materials

Vegetable material: The vegetable material used in current experiments was a half soft variety, known as palm dates badly exploited, cultivated in the area of Adrar (South-western of Algeria). The choice of this variety is justified by its availability and important nutritive value, especially the one of reducing sugars.

Biologic material: The biologic material used is cultures of *Streptococcus thermophilus* and *Lactobacillus bulgaricus*, derived from Orolait of Mascara (Algeria). The mixed culture is homofermentative (homolactic) which means that more than 90% of metabolites are produced by the lactic fermentation. The interest of mixed cultures of *Streptococcus thermophilus* and *Lactobacillus bulgaricus* is to start the phenomenon of symbiosis that is characterized by metabolic interactions between the two strains. This positive interaction demonstrated by comparing the lactic acid production in pure and mixed cultures of two species. The amount of lactic acid produced by the mixed culture is greater than the sum of acids produced by each of the pure cultures [9].

Experimental protocol

Preparation of date syrup: The dates are washed themselves, destoned and grounded. Two and half liters of hot water at 80-85°C were added to 1 kg of date, homogenized and through a cloth. The syrup obtained was centrifuged at 15000 rpm for 10 minutes to separate the cellulose debris. The collected supernatant was used as culture medium. The syrup is sterilized during 20 minutes at 120°C.

Batch fermentation: Batch cultures was performed in 250 ml Erlenmeyer with a working volume of 100 ml, inoculated in the presence of MRS medium (*Lactobacillus bulgaricus*) and M17 medium (*Streptococcus thermophilus*) and incubated for 42 h at 85°C, similar cultures were equally conducted in a reactor having a capacity of five liters provided with all the accessories. The temperature of fermentation was 40°C and the agitation speed was 200 rpm.

Preparation of the culture medium: The used culture medium was based on the MRS medium with Tween80 (1 g/l), MgSO4 (1 g/l), MnSO4 (1.5 g/l), used for the batch culture fermentation experiments, the enriched medium seemed best suited with a yield of lactic acid 49 g/l, where comes from its economical and commercial interest.

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is maintained constant to 42°C and agitation is of 300 turns per minute. 
\( \text{pH} \) was fixed at 6.1 because the development of mixed culture needs a 
\( \text{pH} \) between 5.8 and 6.5).

The biomass evolution, sugar consumption and lactic acid 
production are followed at regular time intervals.

Analytical methods

Biochemical analysis: The content of water is determined by 
drying 10 ml of syrup at 105°C for 18 hour [10]. The content of ashes 
is determined by incineration of one gram of syrup at a temperature of
600°C for 3 hours [11]. The reducing sugars, the sucrose and the total
sugars were determined by the method of Bertrand, reported by Ikeda 
et al. [12]. The total nitrogen is determined by the method of Kreutzer 
et al. [13]. The mineral salts are determined according to the methods
advocated by Nomikos et al. [14].

Kinetics of growth: We take off every two hours until 28 hours 10 ml
of medium of fermentation. We make a reading in a spectrophotometer
to a length of wave 620 nm.

Lactic acid: Concentration of the lactic acid was determined by
acidity titration with NaOH [15].

Residual sugar: The content of residual sugar was determined by
the method of Bertrand, reported by Ikeda et al. [9].

Data processing

The calculation of the kinetic parameters of fermentations has been
performed using Kaleidagraph Software, is a specific program for the
graphs, the histograms and of statistical calculation.

Results and Discussions

Biochemical composition of syrup of dates

The date syrup which has been the subject of our work has a high 
water content 88.69 g/l (Table 1), we agree that a product with a high
water content facilitates lactic acid bacteria proliferation and helps for
a better substrate -enzyme contact since free water is the nutrients
carrier, knowing that \textit{Lactobacillus bulgaricus} needs a minimum water
activity of 0.992 to grow while \textit{Streptococcus thermophilus} needs a
minimum of 0.983.

Dates syrup will be very rich in sugars i.e. 56.8 g/l among them
a rate of 4.16 g/l for sucrose [16]. It indicates the grade of 84 g/l of
total sugars in the soluble extract of date's syrup. These differences are
explained by climatic variations. Reducing sugars are a carbon source
that can satisfy the ferments requirements.

The protein fraction is considerable; therefore it can serve as
a nitrogen source. An ash content of 0.9 g/l indicates its richness of
minerals including potassium (280), sodium (180 mg/100 ml of MF),
phosphorus (28 mg/100 ml of MF) and calcium (146 mg/100 ml of MF)
and a very small amount of magnesium (0.11 mg/100 ml of MF) and
manganese (0.19 mg/100 ml of MF).

The date's syrup has an acidic \( \text{pH} \) 5.9 close to neutrality which
indicates its good nutritional quality (no bacterium contamination)
suitable for the proper development of lactic acid bacteria. Similarly
Ikeda [17] stated that the majority of common varieties (dates of
average quality) have \( \text{pH} \) ranging between 5.3 and 6.3.

The analysis applied on the date syrup shows that it is poor, with
minerals such as Magnesium, Manganese and fatty-acids, so the
addition of these elements (growth factors) is necessary to the syrup in
order to import this quantity of lactic acid.

Finally, the biochemical analysis of date's syrup shows that it can
constitute a fermentation medium of good quality.

Results of the fermentation kinetics

The production curves of lactic acid in enriched and un-enriched
mediums looked the same, the production in the un-enriched begins
with an initial rate of 0.12 g/l to achieve 26 g/l of lactate in 28 h of
fermentation (Figure 1). In parallel, in the medium enriched with
Tween80, MgSO\(_4\) and MnSO\(_4\), the lactic acid rate evolves gradually to
achieve the end of fermentation 49 g/l.

Meanwhile, the biomass evolution in the enriched medium starts
with an initial concentration (OD=0.04) and after 28 h of fermentation
it reaches a maximum value (OD=2.40) (Figure 2). In the case of non-
enriched one we observe a low initial concentration of biomass and in
the end of fermentation optical density reaches a maximum value 1.15.

However, decreasing of sugar rates is very faster in the medium
supplemented with MgSO\(_4\), MnSO\(_4\) and Tween80. Where the mixed
culture consumes about half quantity of initial total sugars, during 28h
of fermentation from an initial quantity of 56.8g/l of sugars, it remains
31.3 g/l which implies an amount of 25.5 g/l (Figure 3). In parallel 11.7
g/l as the quantity of sugar consumed in non-enriched medium.

### Table 1: Biochemical composition of syrup of dates.

| Constituants                   | Date syrup          |
|-------------------------------|---------------------|
| Content in water in percentage| 88.69               |
| pH                            | 5.9                 |
| Total sugars in percentage of M.F | 56.8               |
| Reducing sugars in percentage of M.F | 26.8              |
| Sucrose in percentage of M.F  | 4.16                |
| Proteins in percentage of M.F | 1.02                |
| Ashes in percentage of M.F    | 0.9                 |
| Potassium in mg/100 ml of M.F | 280                 |
| Phosphor in mg/100 ml of M.F  | 28                  |
| Sodium mg/100 ml of M.F       | 180                 |
| Calcium in mg/100 ml of M.F   | 146                 |
| Zinc in mg/100 ml of M.F      | 0.19                |
| Copper in mg/100 ml of M.F    | 0.17                |
| Magnesium in mg/100 ml of M.F | 0.11                |
| Manganese in mg/100 ml of M.F | 0.19                |
Comparing the results for both fermentations indicated that the addition of growth factors in culture medium has a positive and beneficial effect on fermentation, since the growth rate and lactic acid production increase after the enrichment of date syrup.

According to Nomikos et al. [14], the addition of Tween80 allows better cells excretion of lactate by creating pores in the membrane and plays the role of surfactant which makes a good contact between seed and nutrients, and for more it is considered as a source of carbon and energy for electrons.

According to Beauregard et al. [15], the presence of manganese in the medium is specific and cannot be replaced by other minerals, because the Mg²⁺ enters in the structure and functioning of enzymes.

By the same author, the addition of Manganese stimulates the proliferation of Streptococcus thermophilus and it is essential also for Lactobacillus bulgaricus, as Mn²⁺ is the activator of various metabolic reactions, it participates in the transfer of lactate across the cytoplasmic membrane to be released in the medium.

From these results, the enriched medium seems the best for its interesting results. Indeed, a production of 49g/l of lactic acid obtained with medium based on date syrup enriched with Tween80, MgSO₄, and MnSO₄ is higher comparing to the results found by Ikeda et al. [18].

The kinetic parameters of fermentation

Changes over time of specific growth rates (µ), lactate production (Qₗ) and substrate consumption (Qₛ) are presented in figures 4-6.

In unenriched medium, the specific growth speed (µ) after 3 hours of fermentation is 0.38 per hour (Figure 4). For the culture enriched with Tween80, MgSO₄, MnSO₄, the specific growth rate begins with a top speed of 1 h⁻¹ then decreases with time to the point of cancellation, which corresponds to 18 hours of culture time.

Basing on these results, we can say that the earlier addition of manganese, magnesium and Tween80 in fermentation helps in improving culture; enhancing and promoting cell proliferation and metabolic reactions. The decreasing in specific growth rates in the two fermentations can be explained by the disappearance of the culture medium, one or more essential elements to the maintenance and cell growth, it can be due also to the realizing of culture medium by using substances that have been fatal for the growth; this causes despite the abundance of sugar (carbon source) cell lyses and death of cells.

Specific speed of sugar consumption is very important 16 g/g. h in un-enriched fermentation, whereas it is 7 g/g. h for fermentation supplemented with Tween80, MnSO₄ and MgSO₄ (Figure 5). This is logical since the maximum amount of biomass obtained for the first fermentation does not exceed 1 g/l, and to calculate this rate we divide the changes of consumed sugar by biomass variation.

![Figure 2: Evolution of optical density for both fermentations during time.](image)

![Figure 3: Evolution of residual sugars rate for both fermentations during time.](image)

![Figure 4: Evolution of specific growth rates (µ) for both fermentations during time.](image)

![Figure 5: Evolution of specific rates of sugar consumption (Qₛ) for both fermentations during time.](image)
After a time of fermentation, the specific rates of sugars consumption for both fermentations start to decline and to be cancelled after 18 hours of fermentation, which correspond to cells death described above.

Specific speeds of lactic acid production in the two fermentations give identical and maximum values of 2 g/g.h (Figure 6), they are superimposed and decline together to be cancelled after 18 hours of fermentation.

Conclusion

The bioconversion of agricultural by-products mainly the ones rich in fermentable sugars has an economic and strategic interest.

By its biochemical composition, the date's syrup is very rich in carbohydrates 56.8 g/l. which make it a substrate of choice for the development of high value substances.

The kinetics of growth under our experimental conditions showed that the date's syrup is a growing medium that induces the biosynthesis and production of lactic acid in two lactic mixed cultures of *Lactobacillus bulgaricus* and *Streptococcus thermophilus*. The results obtained from the fermentation trials show that the mixed culture can produce 26 g/l of lactic acid in medium based on date's syrup. This medium requires an addition of Magnesium, Manganese and Tween80 to allow an adequate growth and maximum production of lactic acid 49 g/l.

At the end of this study, it appears that the valuation of date's syrup by biological means (fermentation) for the production of lactic acid has certainly huge advantages.

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