

**THE SYNERGISTIC EFFECTS OF A HARVESTING AND CONSERVATION
METHOD DESIGNED FOR SMALL PRODUCERS ON THE QUALITY OF THE
PRODUCED OLIVE OIL.**

Small-scale harvesting and storage effects on oil quality (short running title)

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24 **ABSTRACT**

25 The production of ‘Premium’ olive oil depends in a large part on the quality of the fruit.
26 Small producers see themselves confronted with vast investments and logistic snags when they
27 intend to optimize the harvesting. Today, manual harvesting devices promise less damaged fruit
28 when compared to the traditional methods with nets while the use of a cooling room on the
29 farm is suggested as a solution when the harvesting needs to be stretched out over several days.
30 The use of a manual inverted umbrella during the harvest, together with a storage up to 14 days
31 at 5 °C at a family farm, was studied for three cultivars, ‘Arbequina’, ‘Picual’, and ‘Verdial’.
32 Ten quality parameters of the produced oil were examined in two consecutive years together
33 with an extended sensory analysis in the first year. The results underline the importance of the
34 used harvesting and conservation method on the quality of the extracted oil. The strength of
35 each factor varied in time and according to the cultivar. The ‘Arbequina’ c.v. showed a rapid
36 increase in the importance of the conservation factor, while ‘Picual’ c.v. was the most resistant
37 to deterioration, presenting a lower explanatory value of the conservation factor as compared to
38 the harvesting one. The results indicate that small producers with financial and logistic
39 restrictions can obtain a high-quality product. Either by combining both methods or by
40 choosing the one that guarantees the best results given the cultivar and the specific storage time
41 they need to consider.

42

43 **KEYWORDS**

44 cold storage, manual inverted umbrella, omega square, quality parameters, sensory analysis,
45 virgin olive oil

1. INTRODUCTION

The extension of the Andalusian olive fruit production is with more than 1.500.000 ha not only the core agricultural activity for that region but also the main source of income for more than 250 villages (Junta de Andalucía, 2015; Ley Del Olivar, 2011). Lesser-known characteristics are that 60 % of the 170.000 exploitations are smaller than 5 ha and 80 % less than 10 ha and that more than 50 % of the Agricultural Work Units is done in a strictly family context, in particular as non-salary-compensated work (Junta de Andalucía, 2015).

These structural factors have a direct impact on the used harvesting methods as they limit the financial possibilities of the small producers. Many of them, already of advanced age (70 % are older than 45 years) of which only 60 % define agriculture as their prime activity, are not inclined to do big investments (Serrano *et al.*, 2012; Colombo and Villanueva, 2017). There are several methods available for harvesting olives trees that target the tree in different ways (Sola-Guirado *et al.*, 2014). The impossibility to amortize sophisticated but expensive machinery on the one hand, or to contract specialized services on the other, explains why many small farmers continue to harvest their olives traditionally, beating the olive tree with sticks with nets put on the ground around it.

This traditional method implies that once the fruit is detached, the nets are dragged to the next tree, where they are spread out again until their weight is too heavy to be lugged any further. At that time the fruit is collected in containers or a truckload. The method is speedy and implies low costs but has several inconveniences that may jeopardize the intactness of the fruit. Dragging the fruit on the nets over the ground damage them inevitably, while the harvesters cannot avoid stepping on the fallen fruit while beating the branches. The relation between the quality of the fruit and the extracted oil has been the object of many studies and proven to be primordial to obtain an excellent end product (Garcia and Yousfi, 2006; Yousfi *et al.*, 2012; Rallo *et al.*, 2018; Faminiani *et al.*, 2020). However, when one decides to maximize the yield of

71 the production, the quality of the fruit becomes less important compared to the applied
72 extraction techniques.

73 During the last decade, a growing number of Spanish mills started to produce so-called
74 EVOO Premium's, instead of the common one-sided attention on maximizing the quantity. The
75 choice for quality instead of quantity goes along with the necessity to evaluate rigorously the
76 quality of the fruit in the reception yard, and an adjusted reimbursement to the producer. More
77 recently, preliminary studies to automatize this evaluation are preceding more stringent quality
78 controls soon (Puerto *et al.*, 2015; Navarro Soto *et al.* 2018; Aguilera Puerto *et al.*, 2019).

79 Meanwhile, a new kind of olive oil producer came to the foreground. Smaller-scale and
80 with a clear focus on the production of high quality. However, these producers do not only face
81 the challenge to optimize their harvesting but also confront an additional problem if they do not
82 extract the oil themselves, such as the restrictions that are imposed by the mill that processes
83 their fruit. The necessity to bring in at least several tons of olives to process them as a single
84 batch implies several days of harvesting when working on a family scale. While it is common
85 knowledge that the olives are ideally processed as soon as possible, the conservation of the
86 picked fruits becomes thus a core problem for these small producers.

87 The need to produce good quality oils is emphasized when olive production is
88 approached from a small producer's point of view, not only for them but also for the member of
89 the local cooperative who will be reimbursed not only on the yield but also on the quality of the
90 fruit. A growing amount of economic harvesting devices is coming on the market that
91 specifically addresses these small producers. One promising method for this purpose is the
92 Manual Inverted Umbrella (MIU) which consists of a foldable umbrella that is mounted on the
93 movable structure to collect the fallen fruit harvested using manual aids methods such as
94 branch shakers or shaker rakes. The use of such a MIU turned out to be competitive when

compared with the traditional method while the quality of the picked fruit was significantly better (Plasquy *et al.*, 2019, 2021).

To maintain the fruit at its best, conservation at 5°C is extensively studied and proven for more than 25 years (García and Streif, 1991; García *et al.*, 1994; Canet and García, 1999; Pereira *et al.*, 2002). These studies were mainly focused on prolonging the use of the extraction lines and thus envisioned conservation up to one month or more (García *et al.*, 1996). The benefits of adequate conservation at a shorter time have not been yet studied, especially when the aim is to produce premium quality virgin olive oil and not just avoiding a significant deterioration of its initial quality (Kalua *et al.*, 2008). Nevertheless, it might offer a solution for the individual farmer who seeks to keep his harvested olives during a limited time on the farm before their transport to the mill. It makes it possible to plan the harvesting according to the available workforce, to anticipate bad weather, and to organize in advance a convenient transport and time slot in the mill.

Knowing that both methods (harvesting and conservation) contribute in a significant way to a better result does not answer the question which one has a major impact on the quality parameters of the produced olive oil, especially when the storage time is taken into account as a complementary factor. This work aims to study the effect of MIU harvesting and cold storage on the quality of olive oils. This study becomes vital when the economic resources of the farmer are viewed as a limiting factor in the decision process.

2. MATERIAL AND METHODS

2.1. Location

The experiment took place in the olive grove of ‘Del Cetino’, situated in Bollullos par del Condado (Huelva). It covers 8 ha and includes 1700 trees, mainly of the varieties

‘Arbequina’, ‘Picual’, and ‘Verdial’, planted between 2005 and 2007 at distances of 6 x 7 m and irrigated on a deficient regimen at 50 % of the estimated crop evapotranspiration.

2.2. Harvesting method and conservation facilities

The farm disposes of a MIU and a cooling room with a storage capacity of 5000 kg. The harvesting was performed with a prototype of a MIU and with nets as the control group (Plasquy *et al.*, 2019). The device has an inverted umbrella structure with a width of 6.85 m and is formed by 14 aluminum bars covered with a resistant canvas mounted on a chassis with 4 wheels (Fig. 1). Through an opening in the canvas, the harvested fruit is collected into a plastic and perforated box of 20 kg capacity. This box is introduced and extracted by a system of pulleys and ropes. Two branch shakers (Stihl SP471) were used to detach the fruit.

Two types of harvesting were performed: one using the MIU system (R1) and another using a traditional one, collecting the fruit detached on nets on the ground (R2), both storing the caught fruit in the same perforated boxes. Then the boxes were stored in a pile in a cooling room with a size of 4.3 x 4.4 m and 2.45 m in height isolated with 10 cm thick extruded polystyrene insulating panels on the walls, floor, and roof. (Fig. 2). In the upper part the cold group was placed, formed by a compressor (EMBRACO, UNJ 9232) and an evaporator (LU-VE, SHDN 25-80). The temperature was set at 5 °C (\pm 1 °C). The boxes of the control group were stored outside under a protected roof at ambient temperature. The mean temperatures in October 2017 were 15.4 °C (min) and 29.3 °C (max) and in October 2018, 14.2 °C (min) and 24.7 °C (max). Two types of conservation were performed: one using the cooling room (C1) and another storing the boxes outside at ambient temperature (C2).

2.3. Experimental material

The trials took place between the end of September and the beginning of November during two consecutive years (2017-2018) when the majority of the olives were still green (in all cases the Maturity Index of the fruit samples was situated between 1,5 and 2,5). To assess the effects of the harvesting and conservation method on the produced oil, an equal amount of fruit from the ‘Arbequina’, ‘Picual’, and ‘Verdial’ cultivars were taken at the same moments and under the same conditions as those intended for industrial extraction. Also, it was evaluated the conservation time of 0, 4, 8, and 14 days. To evaluate each factor (harvesting method, conservation method, and conservation time), triplicates of olive samples of both harvesting methods were kept during the distinct periods each one in 6 boxes of 20 kg, which were previously distributed in the cooling room at 5 °C and outside at ambient temperature. The two harvesting methods (R1 and R2), two types of conservation (C1 and C2), and four distinct conservation periods (T0, T4, T8, and T14), gave rise to 18 different combinations of factors for each variety and each year.

2.4. Preparation of the samples

The extraction of the oil was performed in the laboratory using an “Abencor” system (Martinez *et al.*, 1975). Individual samples of 1.500 g olives were crushed in the hammer mill and the resulting paste was distributed in two subsamples of 700 g, which were weighed in two stainless steel casserole pots. Then the paste was malaxated in the thermoblender for 30 min at 30 °C. Afterward, the malaxated paste of each pot was centrifuged at 1.000 G for 1 min. The resulting solid phase of the paste was discarded and the liquid obtained was placed in a graduated 500 mL test tube for separating the aqueous phase of the lipid phase. The Virgin Olive Oil (VOO) extracted from both subsamples was taken from the lipid phases using a Pasteur pipette, filtered with filter paper, and placed in a glass bottle of 250 mL, which was

filled with nitrogen and kept at -20 °C until further examination. The experiment was carried out in triplicate.

2.5. Physico-chemical analysis

Free fatty acidity (FFA), Peroxide value, absorbency at 232 and 270 nm were determined according to the official analytical methods as described in EEC guidelines (EEC, 1991).

The oxidative stability was evaluated using the Rancimat method. The stability was expressed as the oxidation induction time (h) measured with the Rancimat apparatus (Metrohm AG, Herison, Switzerland) at a temperature of 120 °C and an airflow rate of 20 L/h.

The chlorophyll and carotenoid pigment profile was obtained by measuring the spectrophotometric absorbency in the ultraviolet, respectively at 470 nm for the carotenoids and 670 nm for the chlorophyll fraction.

The bitterness index used to estimate the presence of the attribute ‘bitterness’, was calculated using the formula: $IA = 13,33 \times K_{225} - 0,837$ (21).

An estimation of the total polyphenols was obtained by the sum of the calculated amount of polyphenols obtained by measuring the spectrophotometric absorbency at 280 and 335 nm using p-hidroxifenil acetic acid and orto-cumaric acid as calibrating patron respectively. Prior measurements revealed a calibration function for p-hidroxifenil acetic acid of $y = 0,0585x - 0,0007$ and for orto-cumaric acid of $y = 0,0218x + 0,0001$.

The content of α -tocopherols was determined through high-performance liquid chromatography (HPLC) using the IUPAC method (IUPAC, 1992).

2.6. Sensory analysis

A sensory evaluation of the oils was performed to reveal significant differences between the treatments in the first year. The analytical panel was formed by 8 approved tasters of the Instituto de la Grasa (CSIC, Sevilla) and applied the method as described in the EC regulation 640/2008. The procedure permits the classification of the EOV's according to the presence of negative attributes (Muddy, Musty, Winey, Frostbitten, Rancid, others) as well as a measurement of the intensity of positive attributes (Fruity, Bitter, Pungent). The panel members were also asked to classify the presented samples in order of preference to examine whether there were significant differences noticeable between the different treatments at each storage day. To reduce the number of unnecessary tastings the selection started with the samples of day 0 and day 14. If no significant difference between them was detected, no further analysis was performed on the oils extracted from the same cultivar on days 4 and 8.

2.7. Statistical data analysis

Statistical data analysis of the physicochemical parameters was performed using PASW Statistics 18.0 (SPSS). For each cultivar, one-way ANOVA determined the effect of the storage time, considering independently each combination of the other factors (harvesting and conservation method), as well the effect of these four combinations for each separately. Similarly, for each period, the effect of harvesting was studied independently for each conservation method and vice versa. The effect of the storage time and the treatments, defined as the four possible combinations of, on the one hand, R1, R2, and on the other S1, S2, was tested with two-way ANOVA. Finally, the effect of three factors (storage time, harvesting method, and conservation method) was studied by three-way ANOVA. If a significant effect of one of the factors was detected in a parameter, the Tuckey test was applied to discriminate mean values ($P < 0.05$) in each variable.

For each time, the effect size of the different factors and their interaction was determined by calculating the $\hat{\omega}^2$ value. The effect size is a descriptive statistic indicating the proportion of variability in the observed data that is accounted for by the treatments (Maxwell *et al.*, 2018). The effect size can be estimated in various ways, but the calculation of the Omega-Square ($\hat{\omega}^2$) was preferred because this estimation resulted to be less biased when dealing with small samples as compared to Eta and Partial Eta-Squared (Maxwell *et al.*, 2018; Yigit and Mendes, 2018).

The calculation was performed in MS Excell, using the data from the SPSS analysis in the following formula (equation 1):

$$\hat{\omega}^2 = \frac{SS_{Effect} - df_{Effect} MS_{Error}}{SS_{Total} + MS_{Error}} \quad (1)$$

with

SS_{Effect} = the Sum of Squares of each effect (R, C, or R \times C)

df_{Effect} = the degrees of freedom of each effect (R, C, or R \times C)

MS_{Error} = the mean square error

SS_{Total} = Sum of Squares total.

Negative values were set to zero. Omega squared measures become positive when the observed F value exceeds 1,0. Only in these cases, the effect accounts for variance in the population.

To estimate the tendency of the overall effect of the methods on the produced olive oil over time, the average of the $\hat{\omega}^2$ -values for all the parameters and both years for each ST were

calculated. The selection of these parameters was not predetermined because to date there is no theoretical model that integrates the various parameters and their weights.

The Friedman test was used to detect differences in treatments in the sensory evaluation (Sprent and Smeeton, 2001; Fernández, 2018). This non-parametric statistical test involves ranking each taster's judgment together, then considering the values of the ranks by columns. The null hypothesis expects that there are no differences between the treatments. When the calculated probability is significant ($P < 0.05$) it can be concluded that at least 2 of the treatments are significantly different from each other.

Once a significant difference was identified within a tasting cluster, the preference of the panel for the treatments was further deciphered through a new statistic: the Panel Preference (\bar{P}_i). \bar{P} permit a qualitative positioning of the judgments of a panel, composed of n -members who individually ranked t -different treatments. The procedure consists of two steps. In the first one, the value of each \bar{P}_i (equation 2) is calculated as a ratio with as the numerator the sum of the rank given to treatment i by each member, subtracted by the minimum sum of ranks of a treatment, which equals n , to afford comparison of \bar{P}_i -values from panels with a different size. The denominator is formed by the corrected maximum sum of ranks, being the product of t and $(n-1)$. In a second step, the \bar{P} values of the different treatments, each with a value between 0 and 1, are ranked in descending order, with the highest-ranked treatment having the highest ratio and the lowest-ranked one the lowest.

$$\bar{P}_i = 1 - \frac{\sum_{j=1}^n R_{ij} - n}{t(n-1)} \quad (2)$$

with

\bar{P}_i = Panel preference ratio of a specific treatment i ($0 < \bar{P}_i < 1$)

R_{ij} = Ranking giving by panel member j on treatment i

n = number of members in the tasting panel

t = total number of treatments to be compared in conjunction

3. RESULTS

3.1. Free Fatty Acidity

In all of the three varieties, the storage time showed to be highly significant in year 1 and 2, although the ‘Arbequina’ variety turned out to be much more vulnerable when compared to ‘Picual’ and ‘Verdial’, since in both years ‘Arbequina’ showed a highly significant effect of the factors R and S (Supl. Mat. Table 1). After 4 days there was a clear difference between the oils extracted from fruits harvested with nets and stored at room temperature (R2S2) and the other three possible combinations. In year 1, the FAA of these samples even exceeded the limit of 0,80 % of oleic acid, and as a consequence could not be classified as ‘extra’. In year 2, the effect of the cooling was more prominent when compared with year 1 and led to a clear differentiation between the oils on day 14, with the lowest values for treatment R1S1: $0,12 \pm 0,00$, and the highest for R2S2: $0,53 \pm 0,03$. The values of the $\hat{\omega}^2$ presented for both years a similar profile over 14 days. On the day of the harvest, day 0, the method of harvesting turned out to explain the variance slightly above 30 % (Table 1). From day 4 on, the effect of the conservation method was always greater than the harvesting method. However, in year 1, the interaction between both methods gained importance from day 4 on, indicating that the effects of the storage method depended in large part on the intactness of the fruit. In year 2, the role of this interaction was downplayed with an obvious effect of S from day 4, and explaining more than 80 % of the variance on day 14.

The 'Picual' variety showed to be more resistant toward an increase in the FFA, with similar tendencies on day 14, although without exceeding the official limits. In both years, the R2S2 treatment presented the highest values at day 14. In year 1, no significant effect of the harvesting method was detected (Suppl. Mat. Table 1). Measurement of the effect revealed two different profiles (Table 1). In year 1, a steep increase in the importance of the interaction between both methods became visible from day 4. The same happened for the effect of the conservation method from day 8 on. Both balanced each other in importance at day 14, explaining almost 80 % of the variance. In year 2, 76 % of the variance was explained by the storage method on day 4 which implies a downplaying of the role of the harvesting method. On day 14, the effect of the latter, as well as its interaction with the storage method, gained importance in explaining the obtained results.

Over the 2 years, the 'Verdial' variety presented a confusing image. While the storage time and the treatment were significant in both years, the conservation method showed to be highly significant and the harvesting method not in year 1, while in the following it was exactly the inverse (Supl. Mat. Table 1). This profile was reflected in the calculated effect, which showed in year 1 an increasing importance of the conservation method from day 4 up to almost 60 % on day 14. In year 2, this effect was absent while the importance of the harvesting method fluctuated between 30 and 60 % (Table 1).

3.2. Peroxides

The degree of initial oxidation of the three studied cultivars was similar two years but showed a clear difference between 'Arbequina' and 'Picual' on one side, and 'Verdial' on the other. While oils of the former presented values that were always far below the official maximum of 20 mEq O₂/kg oil, the extracted oils of the latter came close to that threshold in

the second year. The Storage time and the harvesting method stood out as a significant factor in all of the 6 different cases, while the conservation method and the interaction between the R and S-factors were only significant in three cases, namely both years in ‘Arbequina’ and the first year in ‘Verdial’. (Suppl. Mat. Table 2).

The magnitude of the effect of the different methods showed a similar profile (Table 1). In all the cases, the harvesting method stood out as the most influential factor in all cases on day 14. In ‘Arbequina’ an increase from day 4 was present in both years. The effect of the harvesting method was at the most during the first 4 days, in which it explained the variance for more than 60 % (year 1) and more than 40 % (year 2). From then on it descended below 5 % on day 14. An effect of the interaction was also involved, although not in the same matter in both years. In year 1, it becomes visible from day 8 where it attained more than 30 % on day 14, while in year 2, the effect is at its most 16 % on day 8. For the ‘Picual’ variety the effect of the conservation method was neglectable over the whole period while the effect of the recollection method differed in both years. The profile for ‘Verdial’ was consistent, although the effect of the harvesting method fluctuated between 60 and 80 % during year 1 and between 15 and 30 % in year 2.

3.3. K 232 and K 270

The calculated values of the absorbance at 232 nm were only affected by the different factors in one case out of six, namely in year 2 of the ‘Verdial’ cultivar. ‘Arbequina’ and ‘Picual’ did not showed a significant difference in the harvesting and conservation methods in neither studied years. The storage time showed to be significant in ‘Verdial’ and year 2 in ‘Arbequina’ (Suppl. Mat. Table 3). Concerning the absorbance at 270 nm the storage time was the only significant factor in all of the 6 cases. The 3 cultivars were comparable regarding the

significance of the harvesting and storage method. All showed a significant effect in the second year for the factor R, while S was not a significant factor in the Arbequina and ‘Picual’ cases, and only during year 2 in the ‘Verdial’ variety (Suppl. Mat. Table 4).

In year 2, the magnitude of the effect of the harvesting method was substantial in all the 3 varieties. In ‘Arbequina’, it was responsible for 35 % (day 8), in ‘Picual’ for 62 % (day 8), and in ‘Verdial’ for 82 % (day 4) and 72 % (day 8) of the variance. A notable effect of the conservation method in the ‘Verdial’ variety increased from day 8 (8 %) and attained 55 % on day 14, at the cost of the importance of the harvesting method, which was reduced to 21 % at that time (Table 1).

3.4. Oxidative stability

The levels of oxidative stability, expressed in hours, varied consistently in the three studied cultivars over both years. The highest values were measured in ‘Picual’, within an overall range between 90 and 145 h; ‘Verdial’ showed values between 60 and 80 h, and Arbequina between 27 and 47 h. In ‘Arbequina’, there was a clear difference in the distinct treatments and a significant effect of harvesting and storage methods. In both years there was a significant effect of the interaction between the conservation method and the storage time, however only in year 1, an interaction effect between the harvesting method and the storage time was observable. In ‘Picual’, the harvesting method turned out to be a highly significant factor, and this in both years, while this was not the case for the conservation method, except when studied in interaction with the ST. The ‘Verdial’ cultivar showed a confusing profile when comparing both years. In year 1, the storage time, harvesting method, and conservation method were very significant, however, in year 2 neither one of these factors showed a significant effect (Suppl. Mat. Table 5).

The impact of the different factors was consistent in the three varieties, characterized by a superior impact of the harvesting method up to 4 days, followed by a decrease from then on. In the course of the 14 days, the strength of the used conservation method increased, however, its maximum and velocity varied along with the cultivars: In ‘Arbequina’ it attained almost 80 % on day 14, in ‘Verdial’ 40 %, and ‘Picual’ 20 %. The results over the 2 years also indicated that the importance of the interaction varies between them. ‘Arbequina’ showed an importance of 20 % at day 4 in year 2. In year 1 of ‘Picual’, 20 % was explained by the interaction on day 8 and increased up to 40 % on day 14, while in year 2, the highest impact was on day 8 with 12 %. In the case of ‘Verdial’, an increase of up to 20 % in both years was present on day 4. However, in year 1, the impact from then on diminished, while in year 2, levels over 20 % were present up to 8 days (Table 1).

3.5. Photosynthetic pigments

The amount of carotenoids and chlorophylls were similar over the two years. In both cases, the three cultivars showed a significant effect on both the Storage time and the treatment (Suppl. Mat. Tables 6 and 7). The three varieties diverted slightly on the effect of the used recollection and harvesting methods over the two years. Overall a significant effect of these factors was present for both photosynthetic pigments over the two years. In year 2, deviant results were obtained in ‘Arbequina’ and ‘Verdial’ regarding the effect of the harvesting method on the level of carotenoids. In that same year, the ‘Verdial’ cultivar did not demonstrate an effect of the harvesting method on the amount of chlorophylls. Finally, the storage method was significant in all cases except in one case, namely in year 1, with regard to the carotenoids in the ‘Picual’ cultivar.

The measures of effect on both pigments were comparable although they varied between the cultivar (Table 1). In ‘Arbequina’, both years demonstrated the importance of the storage method, attaining from day 4 up to day 14 values situated around 80 % of the explained variance. In ‘Picual’, the importance of the conservation method was only visible in year 2, with a linear growth from day 0 (0 %) to day 14 (80 %). In year 1, the used harvesting method explained the variance with values above 50 % from day 0, only to decent at 28 % on day 14. The ‘Verdial’ cultivar, showed to be highly sensitive to the storage method. This was the clearest in year 2, where a steep increase was noticeable from day 4 to day 8, explaining more than 90 % of the variance, up to day 14 with approximately 70 %. In year 1, a linear increase started from day 0 to day 8, after which it descended beneath 10 %. This decline from day 8, went together with a remarkable increase in the importance of the interaction of the factors recollection and harvesting, explaining at day 14 more than 40 % of the variance.

3.6. Bitterness Index

The three cultivars presented distinct trends over the 14 days for the Bitterness Index. In ‘Arbequina’, the storage time and the kind of treatment-induced significant effects in both years, but the factor harvesting did not. The Factor conservation as well as its interaction with the storage time came to the fore as very significant in both years. The ‘Picual’ oils did not show a significant effect due to the storage time in year 1, however, the effect of the interactions of this factor with the factors harvesting and conservation, respectively, turned out to be very significant. Separately, the used harvesting and conservation methods exerted significant effects on this parameter. In both years, the bitterness index of ‘Verdial’ oils experimented significant effects due to the storage and treatment factor. In year 2, the effects of

the used recollection and conservation methods were significant, while in year 1 only the factor conservation (Suppl. Mat. Table 8).

The magnitude of the effects also varied according to the cultivars. While in ‘Arbequina’ the effect of the harvesting method disappeared after day 4, the storage method gained in importance from that moment on, in year 1, attaining its maximum at day 14 of 71 %, and in year 2, even 83 % at day 8 (Table 3). ‘Picual’ maintained in year 1 a high explanatory power for factor ‘harvesting’ with a value that fluctuated between 50 and 80 %. In year 2, a value around 60 % was observable up to day 4, after which it sharply descended towards a neglectable value. The factor ‘conservation’ on the other hand, presented only a slight increase in day 14 up to 30 % in year 1, while in year 2, the values did not exceed 10 %. The ‘Verdial’ cultivar expressed a steady increase in factor conservation method in year 1, attaining a maximum above 60 % on day 14. In year 2, the same maximum was reached, although interrupted with a slight decline at day 8. The difference between the two years was reflected in reverse when comparing the values of factor harvesting, characterized in year 1 with the disappearance of the effect on day 4 and rebounding at day 8 in year 2 (Table 1).

3.7. Total polyphenols

The way the amount of polyphenols was influenced by the harvesting method and storage method varied markedly between the 2 years for the ‘Arbequina’ cultivar (Suppl. Mat. Table 9). While in year 1 there was a clear effect of all the factors studied, in year 2 no effect due to the factor was detected. The ‘Picual’ and the ‘Verdial’ variety showed consistent effects due to storage time and the kind of treatment in both years but diverged in the effect of factors harvesting and conservation. In ‘Picual’ oils, the used harvesting method induced significant effects on polyphenol content in both years, while the conservation method only exerted a

significant effect during year 1. In year 2, the used harvesting and conservation methods induced significant effects on the polyphenol content of ‘Verdial’ oil, however, the effect of the factor harvesting was absent in year 1.

The three varieties expressed the strength of the various factors in different ways (Table 1). In year 1, the profile of the ‘Arbequina’ cultivar was marked by a steep increase of the importance of the used conservation method from day 8, attaining a maximum of 40 % up to day 14. Meanwhile, the impact of the used harvesting method decreased rapidly from day 4 to 20 % on day 8. In ‘Picual’, harvesting came to the fore as the main factor, responsible for explaining between 60 and 80 % during the 14 days, while the impact of the conservation method was with 7 % far less important in year 1. In year 2, the strength of the used harvesting method stayed below 60 % (day 4) and even disappeared from day 8 on. The ‘Verdial’ variety also presented a confusing result. In year 1, the magnitude of strength of the factor ‘conservation’ was characterized by an increase of almost 60 % at day 8 and followed by a subsequent decrease, while at year 2, the effect at that moment was absent whilst the impact was reduced to less than 40 % and brought forward to day 4. The factor ‘harvesting’, with no significant effect in year 1, presented in year 2 a steep increase in day 8 (60 %), only to descent rapidly to a neglectable level on day 14.

3.8. α -tocopherols

During the two assay years, the storage time and the used harvesting method induced a significant effect on the amount of α -Tocopherols in the oils of the three varieties, as well as the interaction of these factors (Suppl. Mat. Table 10). In contrast, the effect of the used conservation method was absent in both years for the ‘Arbequina’ and ‘Verdial’ oils. In the

‘Picual’ cultivar the effect of this factor was only significant in year 1, while the second year was detected a significant effect due to the interaction of this factor and the storage time.

The magnitude of the strength of the factor ‘harvesting’ varied clearly among the two years and the distinct varieties (Table 1). In ‘Arbequina’ there was a clear difference at day 0 (both years 60 %), followed by a descent to 0 % at day 8 and a renewed increase up to almost 80 % in year 1 but only 14 % in year 2. In ‘Picual’, the same high values during day 0 were present, although the profile was different when compared with ‘Arbequina’. On day 4 the effect faded away, only to rise to 75 % on day 8 after which it once again descended towards a neglectable value on day 14. During both years, the ‘Verdial’ variety showed no effect of the factor ‘harvesting’ at day 0 but expressed from day 4 a profile that was comparable with ‘Picual’ at year 1, with a steady increase in day 8 (60-80%) after which descent was set in towards values below 10 %.

3.9. Overall effect of the factors

The means of the obtained $\hat{\omega}^2$ -values for both years and all parameters, were used to express the tendency of the magnitude of strength that characterized the different factors along the storage time of the olives (Figure 3). In ‘Arbequina’ the profile of the three factors under study, namely the harvesting and conservation methods, and the interaction between both, was characterized by a rapid decrease of the initial importance of the factor ‘harvesting’ towards day 4 (60 %) after which its explanatory share settled around 15 % for the rest of the studied period. On day 4, the factor ‘conservation’ became responsible for more than 30 % and further increased to almost 40 % on day 14. The interaction of both factors was situated at 15 % on day 4 after which it slightly fell around 10 %. The descending importance of the used harvesting method and the increasing one of the used conservation methods became equal around day 3

after which the factor ‘conservation’ started to exert a major effect on the final result (Figure 1a). In ‘Picual’, the recollection method was the most important factor in explaining the variance. Despite a moderate decrease from day 1 (55%) to day 4, the value on this day and day 8, were situated around 30 %. From then on the value descended further to less than 15 %. The factor ‘conservation’ on the contrary did not attained a level above 15 % until day 14 where it surpassed the impact of the factor ‘harvesting’ and attained about 25 %. The effect of the interaction came only into play on day 8 with values slightly above 10 % (Figure 1b). The ‘Verdial’ cultivar presented a similar profile as ‘Picual’ concerning the used harvesting method, although with lesser present importance at day 0 (25 %) and overall lower values when compared to the latter. The same can be observed for the factor ‘conservation’, although its effect gained more importance from day 8 (25 %) up to day 14 (30%). The steeper inclination of both curves advanced the crossing point to an earlier moment in time (Figure 1c). When in ‘Picual’ this took place around day 13, it occurred in ‘Verdial’ around day 10. In a similar way as in ‘Arbequina’, the interaction factor exerted his influence at his maximum (15 %) around day 4, after which it decreased to values below 10 %.

3.10. Sensory Analysis

The research design foresaw that, if no significant differences were found between the samples of D0 and D14 or between those of D14, the samples of D4 and D8 would not be tasted. It turned out that only in the 'Arbequina' variety, the panel was able to significantly distinguish differences between the treatments after 14 days of storage. Therefore, only D4 and D8 oils of this variety were further examined. The official limit that disallows the use of the quality label of ‘Extra Virgin’ was only exceeded once: In the R2C2 treatment of the

492 'Arbequina' sample on day 14, the 'extra' category was lost due to a median of the 'Mold' defect
493 above 0, namely 1.2.

494 The evolution of the positive attributes disclosed a clear difference between the
495 'Arbequina' cultivar on the one hand and the 'Picual' and 'Verdial' varieties on the other
496 (Table 4). The bitterness and pungency levels of the 'Arbequina' oils were significantly
497 affected by the storage time and the conservation method, while no significant effect was found
498 due to the interaction between the factors 'harvesting' and 'conservation'. The fruity attribute
499 was not affected by the storage time, nor by the type of treatment. The 'Picual' variety presented
500 no effect of the studied factors on the different attributes. In the 'Verdial' cultivar, the attributes
501 'fruitiness' and 'bitterness' were only significantly influenced by the type of harvesting, showing
502 that the oils from olives collected with the MIU presented values significantly higher, while the
503 storage time decreased the intensity of the attribute 'pungency'.

504 According to the official regulations, most of the samples analyzed did not receive the
505 minimum amount of negative evaluations necessary to lower their quality. Nevertheless, these
506 negative evaluations can be taken into account to detail the applied treatments that showed to
507 be different, as was the case with the 'Arbequina' variety. Calculating the median of the
508 maximum grouped defects of each sample revealed that the deterioration became first visible in
509 the treatments that were kept at room temperature. On day 14, fruit harvested with the
510 traditional method (R2) and cold stored (S1) presented the onset of deterioration. The oils from
511 fruit picked with the MIU (R1) and kept at 5 °C (C1), did not show a median above 0 during
512 the time under investigation (Table 3).

513 For each cultivar and each storage time, the tasters ranked the oils in order of
514 preference. The two from day 0 and the four from day 14 were evaluated together. The
515 Friedman's test detected significant differences, with a $\chi^2 = 11.07$ ($p = 0.05$), between the six

presented samples in the three cultivars, with χ^2_R values of 13,43 for the ‘Arbequina’ c.v., 14,34 for the ‘Picual’ c.v. and 11,35 for the ‘Verdial’ c.v.. However, when the ranking was restricted to the four samples on day 14, no significant difference could be detected by the panel between the treatments of the ‘Picual’ and ‘Verdial’ varieties. As a consequence, no further sensory analysis was performed on the samples of day 4 and day 8 of these varieties. The oil samples of the ‘Arbequina’ cultivar not only showed a significant difference between the treatments at day 14 ($\chi^2_R = 12.45^{**}$), but also on day 4 and day 8, with respectively a χ^2_R value of 12.64* and 13.07*.

The Panel Preference (\bar{P}_i), as calculated with the formula (2) used the obtained ranking results for the Arbequina oil samples from storage days 4, 8, and 14 (Fig. 4). The calculation and the subsequent ranking of the obtained \bar{P}_i values revealed a clear preference for the cold stored olives from day 4 up to day 14. The impact of the harvesting method does not come to the fore as a major factor. On day 0, the tasters almost split in giving preference over one of the two methods (3 preferred R1 against 5 in favor of R2), while only on day 14 a consistent pattern was observed in giving preference over a treatment that included the R1.

4. DISCUSSION

The positive correlation of the conservation temperature on the level of FAA as well as the combined effect of a mechanized harvest and the conservation method was confirmed in the three varieties, as well as the fact that the effect of these factors varied between the cultivars tested.

The observed degree of oxidation (Peroxides, K_{232} , and K_{270}) demonstrated no consistent tendencies for the factors ‘harvesting’ and ‘conservation’ over the studied years. It is only in

year 2 that a significant effect was observed for the Peroxides and the K_{270} in all of the three varieties, while only in the ‘Arbequina’ and ‘Verdial’ cultivar concerning the K_{232} . Yousfi *et al.* (2012) mention values of Peroxides, K_{232} , and K_{270} that were significantly higher when ‘Arbequina’ is recollected mechanically and relate this to the internal ruptures as a consequence of the received blows during the harvesting. The results in year 2 supported this hypothesis, indicating a slight difference between the oils from ‘Arbequina’ and ‘Verdial’ olives in front of ‘Picual’ oils.

For the oxidative stability, the total amount of polyphenols, and the Bitterness Index, a clear distinction was present between the varieties. The results support the hypothesis that the reduction of the oxidation time is not only related to the progress of ripening but also to the aggressiveness of the harvesting method, especially when harvesting the ‘Arbequina’ variety. In the same way, the presence and the evolution of the α -tocopherols are genetically related. The hypothesis of Yousfi *et al.* (2012) that mechanized harvesting and conservation at 18 °C favors the degradation of these compounds is not confirmed as the ‘Arbequina’ cultivar showed an inverse relationship in both years.

The photosynthetic pigments (K_{470} and K_{670}) evolved consistently in the three varieties. However, the calculated strength of the present factors underlines the increasing importance of the factor ‘conservation’ in explaining the observed differences, especially for the ‘Arbequina’ and ‘Verdial’ oils.

While various parameters indicated the importance of the genetic factor when evaluating the effect of the harvesting and conservation method on the various parameters, it is only when the magnitude of the strength of these factors are taken together and compared that their full impact comes to the fore. The obtained results pointed to the critical interrelation that exists between the two factors and their interaction, and demonstrate the differences between

the 3 cultivars as the storage time increases. The vulnerability of the ‘Arbequina’ towards deterioration as compared to the ‘Picual’, and a lesser degree to the ‘Verdial’, is obvious when taking the crossing point of the two curves (harvesting- and conservation-strength) as a point of reference. The curious rebounding of the harvesting-strength in the ‘Picual’ and ‘Verdial’-cultivars at day 8, may indicate that the effects of a produced damage due to a more detrimental harvesting method, can be constrained during the first week due to the cooling of the fruit. However, this initial compensating effect loses power over the following week.

The results of the tasting panel did follow the results based on the physicochemical analysis. The outcome of the ‘Arbequina’ oil judging allowed to specify in detail how the effects of the different treatments were reflected in the quality levels of the samples at each storage time. The fine-tuning on the negative attributes revealed a striking parallel with the measured levels of free fatty acidity and peroxide levels while the presence of positive attributes was mirrored in the raking scores and the panel preference. The different strength profiles were closely related to the results of the tasting panel, especially in the case of the ‘Arbequina’ cultivar where the increase of the storage factor from day 4 is matched with a clear panel preference for cool stored oils.

5. CONCLUSIONS

The study underlines the importance of both harvesting method and conservation in the quality of the oils extracted before their processing and confirms the presence of a genetic predisposition of the different varieties studied. The use of the MIU and the consequent storage of the picked olives at 5 °C does affect the majority of the parameters in a significant way when compared with traditional harvesting and storage at ambient temperature. Especially in a more sensible variety as ‘Arbequina’ as compared with ‘Picual’ or ‘Verdial’. The calculation of the

magnitude of strengths and the calculated panel preference made it possible to discern the explanatory weight of each of the factors, to understand the differences between the varieties to the factors, and to emphasize the need to take into account the days of storage when evaluating their importance. This information is crucial whether to decide which solution fits the farmer best given the specific constraints he has to deal with.

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669 Table 1. Values of $\hat{\omega}^2$ (omega square), as a measure of the effect of the factor harvesting (R), conservation (C), and the interaction between
670 both (R \times C), for each of the 4 storage times during year 1 and 2. Negative values are set to .00.

| | | Storage Time (days) | | | | 0 | | | | 4 | | | | 8 | | | | 14 | | | | | | | | | |
|-----|---|---------------------|------|------|-----|-----|-----|-------|-----|-----|-----|-----|-----|-------|-----|-----|-----|-----|-----|-------|-----|-----|-----|-----|-----|-----|---|
| | | Factor | | R | | C | | R x C | | R | | C | | R x C | | R | | C | | R x C | | | | | | | |
| | | Year | 1 | 2 | 1 | 2 | 1 | 2 | 1 | 2 | 1 | 2 | 1 | 2 | 1 | 2 | 1 | 2 | 1 | 2 | 1 | | | | | 2 | |
| | | ARBEQUINA | | | | | | | | | | | | | | | | | | | | | | | | | |
| 671 | Table tens- three ive at- utes iness, ness, | FFA | .26 | .40 | .00 | .00 | .00 | .00 | .17 | .17 | .33 | .64 | .03 | .08 | .27 | .17 | .40 | .66 | .22 | .10 | .35 | .14 | .37 | .82 | .28 | .02 | 2. In- ity of posit- trib- (Fruit- Bitter- Pun- |
| 672 | | PV | .00 | .61 | .00 | .00 | .00 | .00 | .06 | .08 | .71 | .42 | .02 | .14 | .31 | .34 | .46 | .15 | .00 | .16 | .44 | .60 | .05 | .09 | .32 | .05 | |
| | | K232 | .38 | .59 | .00 | .00 | .00 | .00 | .00 | .00 | .02 | .09 | .30 | .42 | .00 | .24 | .00 | .11 | .00 | .23 | .00 | .09 | .00 | .37 | .27 | .00 | |
| | | K270 | .62 | .09 | .00 | .00 | .00 | .00 | .07 | .28 | .00 | .02 | .00 | .41 | .00 | .35 | .14 | .00 | .00 | .06 | .00 | .24 | .26 | .00 | .57 | .00 | |
| | | Oxidative Stability | .03 | .25 | .00 | .00 | .00 | .00 | .34 | .13 | .00 | .41 | .00 | .30 | .34 | .01 | .07 | .76 | .00 | .07 | .18 | .00 | .70 | .69 | .04 | .00 | |
| | | K470 | .75 | .49 | .00 | .00 | .00 | .00 | .05 | .02 | .88 | .83 | .00 | .03 | .04 | .00 | .86 | .91 | .02 | .00 | .00 | .03 | .89 | .71 | .01 | .00 | |
| | | K670 | .75 | .56 | .00 | .00 | .00 | .00 | .09 | .15 | .65 | .49 | .00 | .18 | .15 | .00 | .57 | .89 | .09 | .00 | .03 | .20 | .52 | .57 | .13 | .00 | |
| | | Bitterness Index | 1.00 | .38 | .00 | .00 | .00 | .00 | .00 | .01 | .00 | .57 | .00 | .22 | .14 | .00 | .21 | .83 | .00 | .00 | .04 | .00 | .71 | .30 | .00 | .11 | |
| | | Total polyphenols | 1.00 | .42 | .00 | .00 | .00 | .00 | .59 | .36 | .00 | .00 | .18 | .34 | .17 | .00 | .42 | .00 | .01 | .00 | .21 | .00 | .30 | .04 | .03 | .00 | |
| | | α-tocopherols | .55 | .55 | .00 | .00 | .00 | .00 | .41 | .30 | .00 | .16 | .00 | .02 | .00 | .02 | .32 | .00 | .08 | .00 | .77 | .14 | .00 | .01 | .00 | .24 | |
| | | PICUAL | | | | | | | | | | | | | | | | | | | | | | | | | |
| | | FFA | .00 | 1.00 | .00 | .00 | .00 | .00 | .00 | .00 | .76 | .00 | .00 | .07 | .00 | .00 | .72 | .60 | .00 | .05 | .27 | .44 | .54 | .34 | .00 | .09 | |
| | | PV | .51 | .27 | .00 | .00 | .00 | .00 | .48 | .00 | .06 | .00 | .13 | .59 | .05 | .08 | .00 | .06 | .18 | .28 | .00 | .00 | .00 | .05 | .11 | .00 | |
| | | K232 | .13 | .08 | .00 | .00 | .00 | .00 | .11 | .21 | .37 | .09 | .00 | .00 | .09 | .00 | .15 | .00 | .04 | .12 | .01 | .00 | .00 | .00 | .00 | .13 | |
| | | K270 | .00 | .93 | .00 | .00 | .00 | .00 | .80 | .29 | .03 | .00 | .09 | .00 | .73 | .62 | .14 | .13 | .05 | .00 | .15 | .07 | .00 | .33 | .07 | .00 | |
| | | Oxidative Stability | .00 | .87 | .00 | .00 | .00 | .00 | .74 | .00 | .04 | .00 | .00 | .00 | .58 | .43 | .03 | .00 | .22 | .12 | .35 | .05 | .21 | .26 | .35 | .00 | |
| | | K470 | .79 | .84 | .00 | .00 | .00 | .00 | .48 | .39 | .00 | .29 | .07 | .04 | .87 | .27 | .08 | .57 | .03 | .08 | .28 | .07 | .00 | .81 | .00 | .00 | |
| | | K670 | .73 | .83 | .00 | .00 | .00 | .00 | .55 | .44 | .21 | .27 | .00 | .01 | .81 | .22 | .03 | .54 | .08 | .06 | .00 | .01 | .25 | .78 | .00 | .00 | |
| | | Bitterness Index | .64 | .62 | .00 | .00 | .00 | .00 | .77 | .59 | .01 | .08 | .00 | .05 | .69 | .00 | .00 | .13 | .00 | .08 | .47 | .06 | .31 | .15 | .10 | .33 | |
| | | Total polyphenols | .74 | .29 | .00 | .00 | .00 | .00 | .82 | .60 | .01 | .00 | .03 | .20 | .52 | .00 | .07 | .15 | .28 | .17 | .83 | .56 | .03 | .00 | .00 | .11 | |
| | | α-tocopherols | .96 | .83 | .00 | .00 | .00 | .00 | .05 | .00 | .00 | .27 | .00 | .11 | .75 | .17 | .05 | .02 | .00 | .08 | .01 | .19 | .38 | .15 | .24 | .21 | |
| | | VERDIAL | | | | | | | | | | | | | | | | | | | | | | | | | |
| | | FFA | .17 | .55 | .00 | .00 | .00 | .00 | .43 | .35 | .00 | .00 | .51 | .00 | .07 | .31 | .22 | .12 | .43 | .03 | .06 | .58 | .54 | .00 | .00 | .00 | |
| | | PV | .61 | .33 | .00 | .00 | .00 | .00 | .71 | .10 | .01 | .00 | .23 | .11 | .83 | .29 | .00 | .00 | .09 | .00 | .65 | .00 | .11 | .00 | .00 | .00 | |

gency) as noted by a sensory analysis panel in the ‘Arbequina’, ‘Picual’ and ‘Verdial’ olive oils extracted from fruit, picked with a Manual Inverted Umbrella (R1) and in a traditional way (R2) and stored during 0, 4, 8 and 14 days at 5 °C (C1) and at ambient temperature (C2)^a. The panel was formed by 8 approved tasters.

| Storage Time (days) | Treatment R (1,2); C (1,2) | Arbequina | | | Picual ^b | | | Verdial ^b | | |
|---------------------|----------------------------|------------|-------------------------------------|--------------------------------------|---------------------|------------|-----------|----------------------|------------|--------------------|
| | | Fruitiness | Bitterness | Pungency | Fruitiness | Bitterness | Pungency | Fruitiness | Bitterness | Pungency |
| 0 | 1,1 | 2.4 ± 1.0 | 3.9 ± 1.5 | 6.1 ± 1.1 | 4.3 ± 1.1 | 4.6 ± 0.7 | 5.3 ± 0.8 | 3.5 ± 0.6 | 4.1 ± 1.0 | 5.2 ± 0.8 |
| | 1,2 | 2.4 ± 1.0 | 3.9 ± 1.5 | 6.1 ± 1.1 A | 4.3 ± 1.1 | 4.6 ± 0.7 | 5.3 ± 0.8 | 3.5 ± 0.6 | 4.1 ± 1.0 | 5.2 ± 0.8 A |
| | 2,1 | 3.0 ± 1.3 | 4.5 ± 1.2 | 6.1 ± 1.4 | 4.6 ± 1.3 | 4.8 ± 1.1 | 5.2 ± 0.7 | 3.4 ± 0.6 | 3.7 ± 1.1 | 4.5 ± 0.6 A |
| | 2,2 | 3.0 ± 1.3 | 4.5 ± 1.2 A^a | 6.1 ± 1.4 A | 4.6 ± 1.3 | 4.8 ± 1.1 | 5.2 ± 0.7 | 3.4 ± 0.6 | 4.1 ± 1.0 | 4.5 ± 0.6 |
| 4 | 1,1 | 1.9 ± 1.2 | 3.6 ± 1.1 | 5.2 ± 1.1 α | - | - | - | - | - | - |
| | 1,2 | 2.2 ± 1.2 | 3.6 ± 0.9 | 4.4 ± 1.5 B β | - | - | - | - | - | - |
| | 2,1 | 2.4 ± 1.4 | 3.3 ± 1.6 | 5.5 ± 0.9 α | - | - | - | - | - | - |
| | 2,2 | 1.9 ± 0.8 | 3.0 ± 1.1 AB | 4.0 ± 1.5 B β | - | - | - | - | - | - |
| 8 | 1,1 | 2.4 ± 1.3 | 3.3 ± 1.1 α | 4.6 ± 1.7 α | - | - | - | - | - | - |
| | 1,2 | 1.8 ± 1.0 | 2.2 ± 0.9 β | 2.7 ± 0.7 C β β | - | - | - | - | - | - |
| | 2,1 | 2.6 ± 1.5 | 3.1 ± 1.3 α | 5.0 ± 1.1 α | - | - | - | - | - | - |
| | 2,2 | 1.7 ± 1.1 | 2.1 ± 1.3 B β | 3.3 ± 1.4 B α β | - | - | - | - | - | - |
| 14 | 1,1 | 2.8 ± 1.6 | 3.9 ± 1.5 α | 5.5 ± 1.2 α | 4.5 ± 1.2 | 4.3 ± 0.8 | 5.1 ± 0.8 | 3.0 ± 0.9 | 4.7 ± 0.7 | 5.2 ± 0.5 |
| | 1,2 | 2.4 ± 1.2 | 3.5 ± 1.4 α β | 4.2 ± 1.2 BC α β | 4.3 ± 1.2 | 4.1 ± 1.2 | 4.4 ± 1.3 | 2.9 ± 1.1 | 3.8 ± 1.0 | 4.7 ± 1.2 B |
| | 2,1 | 2.7 ± 1.8 | 4.8 ± 1.1 α | 4.9 ± 1.1 α | 4.2 ± 1.1 | 4.3 ± 1.0 | 4.9 ± 0.9 | 3.8 ± 0.8 | 3.1 ± 0.7 | 4.1 ± 0.8 B |
| | 2,2 | 1.7 ± 0.8 | 2.0 ± 1.1 B α β | 3.4 ± 1.4 B α β | 4.4 ± 0.8 | 4.1 ± 1.0 | 4.7 ± 1.1 | 2.2 ± 0.8 | 3.6 ± 1.2 | 4.8 ± 1.0 |
| Storage Time (ST) | | 0.217 | 0.000 | 0.000 | 0.059 | 0.055 | 0.900 | 0.718 | 0.568 | 0.000 |
| Treatment (T) | | 0.203 | 0.010 | 0.000 | 0.952 | 0.749 | 0.965 | 0.031 | 0.150 | 0.340 |
| ST × T | | 0.752 | 0.057 | 0.199 | 0.960 | 0.724 | 0.962 | 0.383 | 0.486 | 0.348 |
| R | | 0.700 | 0.774 | 0.780 | 0.682 | 0.924 | 0.749 | 0.010 | 0.014 | 0.767 |
| R × ST | | 0.421 | 0.311 | 0.366 | 0.720 | 0.733 | 0.728 | 0.677 | 0.814 | 0.324 |
| C | | 0.062 | 0.002 | 0.000 | 0.980 | 0.613 | 0.707 | 0.171 | 0.126 | 0.130 |
| C × ST | | 0.326 | 0.166 | 0.738 | 0.682 | 0.334 | 0.878 | 0.632 | 0.775 | 0.035 |

682

^a In each variable the values of different treatments followed by different letters are significantly different according to the Tukey test (P < 0.05). Absence of letters means no significant effect due to treatment according to one-way ANOVA (P < 0.05). In each column, values at different storage times (ST) and the same harvesting method (R) and conservation method (C), followed by different upper bold case letters are significantly different; four values at each ST, followed by different lower case letters (a, b, c, d) are different; two values at the same ST and same harvesting method, but different conservation method, followed by different Greek letters are significantly different. Each value is the mean ± SD of 3 replicates.

^b When the Friedman test detected that the panel was not able to differentiate significantly the overall quality of the olive oils at day 0 and day 14, the samples at day 4 and day 8 were not sensory evaluated.

688
689

Table 3. Number of defects and the median of their intensity as reported by the sensory panel, in the oil extracted from the ‘Arbequina’ cultivar, harvested with a Manual Inverted Umbrella (R1) or in a traditional way (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) or at ambient temperature (C2). The panel was formed by 8 approved tasters.

| Treatment | D0 | | D4 | | D8 | | D14 | |
|-----------|--------|-----------|--------|-----------|--------|-----------|--------|-----------|
| | median | # defects | median | # defects | median | # defects | median | # defects |
| R1 C1 | 0.00 | 1 | 0.00 | 3 | 0.00 | 1 | 0.00 | 2 |
| R1 C2 | 0.00 | 1 | 0.00 | 5 | 0.00 | 1 | 1.60 | 6 |
| R2 C1 | 0.00 | 1 | 0.80 | 10 | 1.95 | 8 | 2.25 | 11 |
| R2 C2 | 0.00 | 1 | 2.65 | 13 | 1.55 | 8 | 2.70* | 11 |

*median for the attribute ‘musty’=1.2

707 Figure legends

708

709 Figure 1. Recollecting olive fruit with the Manual Inverted Umbrella (MUI), 4
710 operators and the use of 2 branch shakers (R1)

711

712 Figure 2. Cooling room situated in a barn on the farm with a capacity of 5.000 kg,
713 and where the fruit can be kept at 5 °C. (C1)

714

715 Figure 3. Magnitude of strength of the factors recollection and conservation and their
716 interaction for three different varieties (Arbequina, Picual, and Verdial), based on the mean of
717 the calculated omega squared ($\hat{\omega}^2$) values of 10 parameters (FFA, peroxides, K232, K270,
718 oxidative stability, photosynthetic pigments, bitterness index, total polyphenols, α -
719 tocopherols) over 14 days.

720

721 Figure 4. The Panel Preference (\bar{P}_i) for the oil extracted from the ‘Arbequina’ cultivar,
722 recollected with a Manual Inverted Umbrella (R1) or in a traditional way (R2) and stored
723 during 4, 8, and 14 days at 5 °C (C1) or at ambient temperature (C2). The panel was formed
724 by 8 approved tasters. Data at day 0 are not showed as the Friedman test can not be performed
725 on two treatments.

726

727 Supplementary material Table legend

728 Table 1. Free Fatty Acid (% oleic acid) noted in the ‘Arbequina’, ‘Picual’, and ‘Verdial’ olive
729 oils extracted from fruit, picked with a Manual Inverted Umbrella (R1) and in a traditional
730 way (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient temperature (C2).

731

732 Table 2. Peroxide Value (mEq O₂/ kg oil) noted in the ‘Arbequina’, ‘Picual’, and ‘Verdial’
733 olive oils extracted from fruit, picked with a Manual Inverted Umbrella (R1) and in a
734 traditional way (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient
735 temperature (C2).

736

737 Table 3. Absorbance at 232 nm (K232) noted in the ‘Arbequina’, ‘Picual’, and ‘Verdial’
738 olive oils extracted from fruit. picked with a Manual Inverted Umbrella (R1) and in a
739 traditional way (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient
740 temperature (C2).

741

742 Table 4. Absorbance at 270 nm (K270) noted in the ‘Arbequina’. ‘Picual’ and ‘Verdial’ olive
743 oils extracted from fruit, picked with a Manual Inverted Umbrella (R1) and in a traditional
744 way (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient temperature (C2).

745

746 Table 5. Oxidative Stability (h) noted in the ‘Arbequina’. ‘Picual’ and ‘Verdial’ olive oils
747 extracted from fruit, picked with a Manual Inverted Umbrella (R1) and in a traditional way
748 (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient temperature (C2).

749 Table 6. Absorbance at 470 nm (Carotenoids) noted in the ‘Arbequina’. ‘Picual’ and
750 ‘Verdial’ olive oils extracted from fruit picked with a Manual Inverted Umbrella (R1) and in a
751 traditional way (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient
752 temperature (C2).

753

754 Table 7. Absorbance at 670 nm (Chlorophyll) noted in the ‘Arbequina’. ‘Picual’ and
755 ‘Verdial’ olive oils extracted from fruit, picked with a Manual Inverted Umbrella (R1) and in
756 a traditional way (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient
757 temperature (C2).

758

759 Table 8. Bitterness Index noted in the ‘Arbequina’. ‘Picual’ and ‘Verdial’ olive oils extracted
760 from fruit, picked with a Manual Inverted Umbrella (R1) and in a traditional way (R2) and
761 stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient temperature (C2).

762

763 Table 9. Total Polyphenols (mg/kg) noted in the ‘Arbequina’, ‘Picual’, and ‘Verdial’ olive
764 oils extracted from fruit, picked with a Manual Inverted Umbrella (R1) and in a traditional
765 way (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient temperature (C2).

766

767 Table 10. α -Tocopherols (mg/kg) noted in the ‘Arbequina’, ‘Picual’, and ‘Verdial’ olive oils
768 extracted from fruit, picked with a Manual Inverted Umbrella (R1) and in a traditional way
769 (R2) and stored during 0, 4, 8, and 14 days at 5 °C (C1) and ambient temperature (C2).