



HAL
open science

Using $\delta^{13}\text{C}$ and hydroscares for discriminating cultivar specific drought responses

Marc Plantevin, Mark Gowdy, Agnès Destrac-Irvine, Elisa Marguerit, Gregory Gambetta, Cornelis van Leeuwen

► To cite this version:

Marc Plantevin, Mark Gowdy, Agnès Destrac-Irvine, Elisa Marguerit, Gregory Gambetta, et al.. Using $\delta^{13}\text{C}$ and hydroscares for discriminating cultivar specific drought responses. *OENO One*, 2022, 56 (2), pp.239 - 250. 10.20870/oeno-one.2022.56.2.5434 . hal-03745484

HAL Id: hal-03745484

<https://hal.inrae.fr/hal-03745484>

Submitted on 4 Aug 2022

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License



ORIGINAL RESEARCH ARTICLE

Using $\delta^{13}\text{C}$ and hydroscares for discriminating cultivar specific drought responses

Marc Plantevin^{1*}, Mark Gowdy¹, Agnès Destrac-Irvine¹, Elisa Marguerit¹, Gregory A. Gambetta¹ and Cornelis van Leeuwen¹

¹ EGFV, Univ. Bordeaux, Bordeaux Sciences Agro, INRAE, ISVV, F-33882 Villenave d'Ornon, France

Marc Plantevin and Mark Gowdy contributed equally to this article and should both be considered as first authors.

► This article is published in cooperation with Terclim 2022 (XIVth International Terroir Congress and 2nd ClimWine Symposium), 3-8 July 2022, Bordeaux, France.



*correspondence:
marc.plantevin@inrae.fr

Associate editor:
Cassandra Collins



Received:
27 February 2022

Accepted:
19 April 2022

Published:
24 June 2022



This article is published under the Creative Commons licence (CC BY 4.0).

Use of all or part of the content of this article must mention the authors, the year of publication, the title, the name of the journal, the volume, the pages and the DOI in compliance with the information given above.

ABSTRACT

Measurement of carbon isotope discrimination in berry juice at maturity ($\delta^{13}\text{C}$) provides an integrated assessment of vine water status and water use efficiency (WUE) during the period of berry ripening, and when collected over multiple seasons, can provide an indication of drought stress responses. Berry juice $\delta^{13}\text{C}$ measurements were carried out on 48 different varieties planted in a common garden experiment in Bordeaux, France from 2014 through 2020 and found important differences across this large panel of varieties. Cluster analysis showed that $\delta^{13}\text{C}$ values are likely affected by the differing phenology of each variety, resulting in berry ripening of different varieties taking place under different conditions of soil water availability within the same year. Accounting for these phenological differences, the cluster analysis created a classification of varieties that corresponds well to our current empirical understanding of their relative drought tolerance. In addition, using measurements of predawn and midday leaf water potential measurements collected over four seasons on a subset of six varieties, a hydroscape approach was used to develop a list of metrics indicative of the sensitivity of stomatal regulation to water stress (i.e. an/isohydric behaviour). Key hydroscape metrics were also found to be well correlated with some $\delta^{13}\text{C}$ metrics. A variety's water potential regulation as characterized by a minimum critical leaf water potential as determined from hydroscares was strongly correlated to $\delta^{13}\text{C}$ values under well-watered conditions, suggesting that the latter may be a useful indicator of drought stress response.

KEYWORDS: water use efficiency, carbon isotopic discrimination, water potential, drought tolerance, VitAdapt, hydroscares, grapevine, *Vitis vinifera*

INTRODUCTION

Vines are cultivated in a wide range of climates, from very dry (such as Cyprus or Aragon, Spain) to extremely wet (such as the Hunter Valley, Australia). The reason for the commercial success of vineyards across such a large range of environments can be explained by the winegrowers' ability to adapt through viticultural practices or by choosing plant material adapted to the local climatic conditions. These adaptations are increasingly important as local environmental conditions are changing under the effect of climate change (IPCC, 2021). Rainfall patterns are changing, with precipitation increasing, or decreasing depending on the region. An increase in temperature and thus evaporative demand will induce drier conditions in most winegrowing regions, even in those with unchanged precipitation regimes. As a result, adaptations are needed to maintain commercially viable viticulture in current winegrowing areas (Fraga *et al.*, 2012; Ollat *et al.*, 2016; van Leeuwen and Darriet, 2016).

Vines react to water deficits by closing their stomata (Winkel and Rambal, 1990; Williams and Araujo, 2002), which reduces carbon assimilation (Flexas *et al.*, 1998). Despite this reduction in photosynthesis, mild water deficit has positive effects in fine wine production: micro-climatic conditions are improved (better light penetration in the canopy due to lower vigour); more carbohydrates are available for berry ripening due to reduced competition with shoot growth; and berries become smaller with increased levels of anthocyanins in the produced wines (Roby *et al.*, 2004; Keller, 2010; Carbonneau *et al.*, 2015; Triolo *et al.*, 2019). The positive impact of mild water deficit has also been shown on most aroma compounds in grapes and wines (van Leeuwen *et al.*, 2020) as well as on the sensory properties of aged red Bordeaux wines (Le Menn *et al.*, 2019).

However, water deficits can also lead to yield reductions that may negatively impact commercial viability (van Leeuwen *et al.*, 2019a). Under severe water deficit, yield is highly impacted and grape ripening (e.g., sugar accumulation) can slow down or even stop in extreme conditions (van Leeuwen and Darriet, 2016). Although vines can survive under severe droughts and are unlikely to die (Charrier *et al.*, 2018), commercially viable fine wine production may become more difficult under increased levels of water deficit.

Many options for adapting viticulture to a warmer and drier future are available, such as changes in canopy management practices, vineyard planting densities, or plant material selection (e.g., clones, rootstocks, and/or varieties) (Fraga *et al.*, 2012; van Leeuwen *et al.*, 2019a; van Leeuwen *et al.*, 2019b). To inform future plant material selection, the drought responses of different varieties need to be better characterized. Assessing the regulation of vine water status is fundamental in making this characterization. Measurements of vine water potentials and the use of carbon isotope discrimination are two methods that have proven particularly useful in assessing water status, each with its own caveats (Choné *et al.*, 2001; Gaudillère *et al.*, 2002; Cifre *et al.*, 2005; van Leeuwen *et al.*, 2009; Santesteban *et al.*, 2012).

$\delta^{13}\text{C}$ provides an integrative measure of the water status of the plant over a given period of time and can be assessed in leaves and/or berry juice at the end of the season (Gaudillère *et al.*, 2002; Santesteban *et al.*, 2012). During the process of photosynthesis there is a discrimination against assimilation of CO_2 containing ^{13}C isotopes due to the higher reactivity of ^{12}C isotopes with the Rubisco enzyme in the photosynthesis reaction. In addition, the rate of diffusion of $^{13}\text{CO}_2$ through the stomata and leaf boundary layer is less than for $^{12}\text{CO}_2$. This results in lower $^{13}\text{C}/^{12}\text{C}$ isotope ratios in carbohydrates photosynthesized at that time when compared to atmospheric CO_2 (Farquhar 1989; Santesteban *et al.* 2012). As stomata close in response to drought stress, the $^{13}\text{C}/^{12}\text{C}$ ratio rises in the intercellular space and sugars formed and incorporated into various plant tissues at that time will contain a modified ratio of carbon isotopes (Bchir *et al.*, 2016).

When assessed on berry juice at maturity, $\delta^{13}\text{C}$ is a useful measure of the level of water deficit that existed during the time sugars in the berry juice accumulate (van Leeuwen *et al.*, 2009). $\delta^{13}\text{C}$ effectively provides a measure of the ratio between the rate of photosynthesis and the stomatal opening over a period of time (Farquhar, 1989). This is similar to the ratio representing water use efficiency (WUE), which at the leaf level is the amount of carbon assimilated by photosynthesis (A_n) per amount of water transpired (g_s), with a less negative $\delta^{13}\text{C}$ corresponding to less stomatal opening and higher WUE (Medrano *et al.*, 2010, Souza *et al.*, 2005, Bchir *et al.*, 2016). A study of eight own-rooted *Vitis vinifera* varieties found a strong positive correlation between $\delta^{13}\text{C}$ and leaf-level WUE (Tomás *et al.*, 2014).

Vine water status can also be assessed by different measures of leaf water potential. Midday leaf water potential (Ψ_L) is a measure of the immediate negative water potential (i.e., suction) in the leaf in response to many factors, such as vapour pressure deficit, stomatal closure, soil water status, and plant hydraulic conductivity (Choné *et al.*, 2001). Predawn leaf water potential (Ψ_{pp}) is experienced at night when the vines have equilibrated with the soil water potential (Williams and Araujo, 2002) and can be considered as a proxy for soil water availability (Gaudin *et al.*, 2017), or more precisely of the water availability in the most humid soil layer (Améglio *et al.*, 1999).

The amount of stomatal regulation in response to changing Ψ_L varies among genotypes, a concept which has been referred to as (an)isohydricity (Tardieu and Simonneau, 1998; Hochberg *et al.*, 2018). An anisohydric species allows transpiration and photosynthesis at more negative Ψ_L while an isohydric species will regulate stomata at less negative Ψ_L , decreasing transpiration and eventually photosynthesis (Tardieu and Simonneau, 1998). Plant species do not adhere to strict (an)isohydric definitions, but instead vary across a continuum from anisohydric to isohydric (Chaves *et al.*, 2010; Klein, 2014) and this is also true across *Vitis vinifera* varieties, for which differences across genotypes mainly occur at medium levels of water deficit (Levin *et al.*, 2019). In addition, growth conditions (e.g., greenhouse vs field grown) and the climatic conditions which impact the range of

water potentials can influence isohydric versus anisohydric behaviour even within the same genotype (Chaves *et al.*, 2010; Charrier *et al.*, 2018, Hochberg *et al.*, 2018).

Recently the “hydroscape” approach was proposed to visualize how plants regulate their water potential by means of stomatal regulation and to classify genotypes according to their behaviour under drought stress (Martínez-Vilalta *et al.*, 2014). The area of the hydroscape plot defines the range over which a plant regulates its Ψ_L as a function of Ψ_{PD} , with the larger the range, the less strict (i.e., more anisohydric) the genotype is at controlling its stomatal conductance under water deficit (Meinzer *et al.*, 2016). Conversely, a small hydroscape indicates a genotype with more strict stomatal control (i.e., more isohydric). This methodology has shown to be effective in classifying drought-responses of different species (Li *et al.*, 2019, Meinzer *et al.*, 2016, Álvarez-Maldini *et al.*, 2021). Hydroscares, however, are time-consuming to produce, requiring multiple measurements of Ψ_{PD} and Ψ_L over a wide range of water deficits.

This study evaluates the drought stress responses of several grapevine varieties by means of $\delta^{13}C$ measurements in berry juice sugars and by hydroscares. Some apparent relationships between the results obtained by both methods are also discussed.

MATERIALS AND METHODS

1. VitAdapt vineyard

Data for this study were obtained in the VitAdapt experimental vineyard located at the INRAE (Institut National de Recherche pour l’Agriculture et l’Alimentation) research center of Bordeaux, in France (44°47’23.8”N, 0°34’39.3”W) (Destrac-Irvine and van Leeuwen, 2017). The soil is composed mainly of gravel and sand, representative of the Pessac-Leognan AOP (Bordeaux, France). The vineyard was planted in 2009 with 52 different varieties, including 4 hybrids and 48 *Vitis vinifera* L. cultivars on a 0.7 ha parcel with 46 rows of 75 vines each on a density of 5,555 vines/ha (row spacing of 1.8 m and inter vine spacing of 1.0 m). The trunk height is 0.5 m and grapevines are pruned to a double guyot and topped during the growing season at 1.5 m. Each variety was planted on Selection Oppenheim 4 (SO4) rootstock, clone 761. Each variety was tested for major virus diseases and only clean material was planted. There is a mowed cover crop in between each vine row with mechanical tillage under the vine row. The vineyard is dry-farmed and fertilisation is adjusted based on petiole analyses carried out at mid-veraison. To account for soil variability, the VitAdapt experimental design was laid out in randomized blocks with each block containing two rows of five vines per variety. For this study, four blocks were sampled.

2. Carbon isotope discrimination ($\delta^{13}C$)

Measurements of $\delta^{13}C$ were taken from berry juice sugar collected each year at maturity from 10 vines of each variety in each of the four blocks. Measurements of grape sugars and

acids were carried out weekly starting at mid-veraison and maturity was considered when sugars started to plateau-off and acidity was low. Sampling was started in 2014, five years after initial planting in 2009, and continued each year through 2020. Berries were harvested from the 10 vines in each block and juice was extracted and centrifuged at 13,500 rpm for 10 minutes (Sigma 13 6K15, SIGMA Laborzentrifugen GmbH, Osterode am Harz, Germany). The juice was then analyzed on a Vario Micro Cube elemental analyzer coupled in a continuous flow mode to an isotopic ratio mass spectrometer (IsoPrime, Elementar). Glutamic acid USGS40 and Caffeine IAEA-600 were used as international standards during the analyses. All results are expressed in delta notation ($\delta^{13}C$) and reported relative to the Vienna Pee Dee Belemnite (VPDB) international reference. Over the seven years a total of 17 to 28 $\delta^{13}C$ measurements were obtained for each variety.

From 2017 through 2020 $\delta^{13}C$ was also measured on eight individual vines of six varieties (Cabernet-Sauvignon, Merlot, Grenache, Tempranillo, Semillon, Ugni blanc), two of each from the four blocks, upon which water potential measurements were also taken periodically over the season. Of these individual vines, some started to lose too many leaves due to disease pressure or multiple water potential measurements, in which case water potential was taken on an adjacent vine, upon which $\delta^{13}C$ was also measured with the values being averaged.

3. Water Potential

Water potential measurements were taken periodically over the seasons 2017 through 2020 on two vines each in the four blocks of Cabernet-Sauvignon, Grenache, Merlot, Semillon, Tempranillo and Ugni blanc. These varieties are some of the most widely planted varieties worldwide (Anderson and Aryal, 2013). Merlot, Tempranillo and Semillon (Rogiers *et al.*, 2009) are known for being sensitive to drought stress, while Cabernet-Sauvignon and Grenache are renowned for being less sensitive (Santesteban *et al.*, 2009). No data were obtained on Grenache in 2018 because of downy mildew damage and measurements on Ugni blanc did not start until 2018. Measurements were collected at seven timepoints in 2017 and 2018 and six timepoints in 2019 and 2020.

From each vine on each sampling date, three types of water potential were measured with leaf water potential (Ψ_L) and stem water potential (Ψ_S) measurements beginning at 2:00 p.m. on a given day and predawn water potential (Ψ_{PD}) started around 3 a.m. the morning after, with all measurements completed within two hours. For midday Ψ_S measurements the leaves were sealed in foil covered bags while on the plant one hour prior (Choné *et al.*, 2001). Measurements were taken by the method of Scholander *et al.* (1965) using a pressure chamber with digital manometer (DG MECA, 33175 Gradignan, France).

4. Hydroscares

The method for preparing hydroscares was adapted from the concepts presented by Meinzer *et al.* (2016), Charrier *et al.* (2018), and Hochberg *et al.* (2018). First, following the procedure of Meinzer *et al.*, (2016), a scatter plot is

made of Ψ_L on the vertical axis and corresponding Ψ_{PD} on the horizontal axis using only the minimum values of Ψ_{PD} and Ψ_L measured from replicate vines on each date. By using minimum Ψ_L values, data that was more influenced by non-limiting conditions for other variables, such as vapour pressure deficit, were removed. Then a 1:1 line is added representing the point at which $\Psi_{PD} = \Psi_L$, a theoretical condition when stomata are completely closed and the whole vine is in equilibrium with soil matric potential.

Then following the method of Charrier *et al.* (2018), bins of Ψ_{PD} for every 0.05 MPa were made and the average of all the minimum Ψ_L measurements within each bin was calculated. Then starting with the bin-averaged Ψ_L at zero Ψ_{PD} (the intercept), the bin averages for progressively more negative Ψ_{PD} are sequentially compiled into a linear regression using

bin averages at all previous, less negative Ψ_{PD} . This is done iteratively until the coefficient of determination (r^2) starts to decrease, suggestive of a breakpoint (called critical leaf water potential, Ψ_{Lcrit}) in the linear relationship between Ψ_L and Ψ_{PD} (Hochberg *et al.*, 2018). From this point, the data suggests the bin-average Ψ_L at more negative Ψ_{PD} will have similar values that form a horizontal line of slope zero at Ψ_{Lcrit} . This represents the lowest Ψ_L against which the plant will regulate its stomatal conductance at increasingly negative Ψ_{PD} . In uncontrolled field conditions, the condition of $\Psi_{PD} = \Psi_L$ is rarely achieved, so the horizontal line at Ψ_{Lcrit} must be extended to meet the 1:1 line in order to complete the enclosure of the hydroscape. The area of the hydroscape is then bounded by the horizontal line at Ψ_{Lcrit} (a in Figure 1), the 1:1 line (b in Figure 1), the vertical line at $\Psi_{PD} = 0$ (c in Figure 1), and the regressed line between the

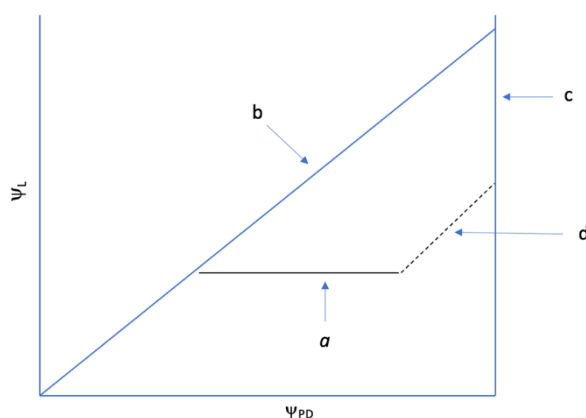


FIGURE 1. A conceptual hydroscape bordered by the line representing the Ψ_{Lcrit} (a), the 1:1 line (b), the vertical line at $\Psi_{PD} = 0$ (c) and the regressed line between the intercept at $\Psi_{PD} = 0$ and Ψ_{Lcrit} (d).

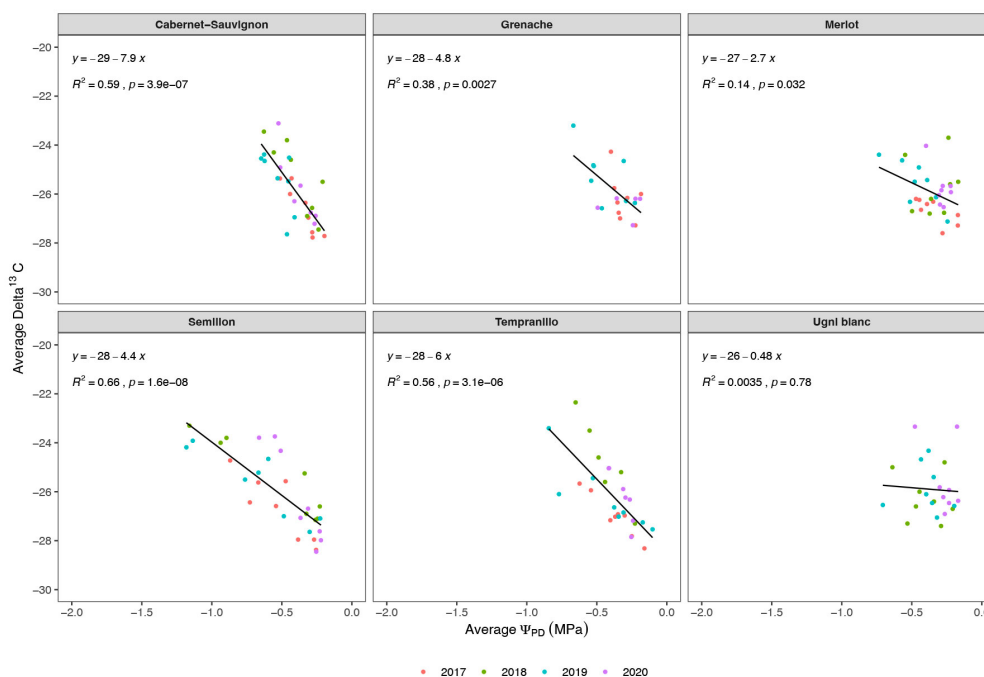


FIGURE 2. Individual vine $\delta^{13}C$ as a function of Ψ_{PD} , averaged over the data collected between 20 days before until 60 days after veraison, for six varieties in 2017 (except for Ugni blanc), 2018 (except for Grenache), 2019 and 2020 in Villenave d’Ornon, France.

intercept at $\Psi_{PD} = 0$ and Ψ_{Lcrit} (d in Figure 1). In theory, the greater the area enclosed by the hydroscape, the more anisohydric the variety (Meinzer *et al.*, 2016).

Filtering, graphing, and linear regression analysis was performed in the *R* software environment using several functions from the *dplyr* package (Wickham *et al.*, 2021a) and the *ggplot2* package (Wickham *et al.*, 2021b) with hierarchical clustering analyses and dendrograms being done using the *cluster* package.

RESULTS AND DISCUSSION

1. Climate conditions

The two years with the greatest water deficits were 2016 followed by 2018. Some years showed particular climatic conditions. In 2015 (globally a dry year), August was very rainy (109.5 mm). In 2017, evaporative demand was low in May and rainfall was high in September (82 mm), which explains why water deficit was low in this year despite a dry July and August. 2020 was one of the earliest vintages ever in Bordeaux, with high rainfall in May and June, while July was very dry (2.5 mm) and rainfall in August was close to average (46.5 mm). More information regarding vine water status during these years is presented in Figures S1 and S2.

2. Carbon isotope discrimination ($\delta^{13}\text{C}$)

2.1 Relationship of berry juice $\delta^{13}\text{C}$ to leaf water status

Figure 2 presents plots of $\delta^{13}\text{C}$ on an individual vine basis for six varieties from 2017 through 2020 versus the corresponding average Ψ_{PD} measured during berry ripening (i.e., from 20 days before mid-veraison until 60 days after veraison). The $\delta^{13}\text{C}$ and average Ψ_{PD} were well correlated for Cabernet-Sauvignon, Grenache, Semillon and Tempranillo, while not as well correlated for Merlot and Ugni blanc (Note, only three years of data were collected for Ugni blanc and Grenache). Plots of $\delta^{13}\text{C}$ versus Ψ_s (not shown) showed similar relationships. These results are consistent with previous studies showing $\delta^{13}\text{C}$ as being well correlated to Ψ_{PD} (Gaudillère *et al.*, 2002) and to midday Ψ_s (Santesteban *et al.*, 2012) suggesting increased water use efficiency in response to increasing vine water deficit.

2.2 Varietal differences in $\delta^{13}\text{C}$

Figure 3 presents the range of $\delta^{13}\text{C}$ values for each of the 48 varieties across the seven years of measurement, with the thickness of the violin plot indicative of the distribution of values and the black dot representing the mean of all measurements. As discussed in the next section, the average $\delta^{13}\text{C}$ measured in the year with the most negative values is an important metric of drought responses for a given variety. The red dots on the violin plots represent this minimum mean value for each variety and are the values upon which the varieties are ordered along the vertical axis in Figure 3.

$\delta^{13}\text{C}$ is variety dependent, with Tinto Cão showing the most negative mean values across all years (black dots) and Colombard showing the least negative mean values. The difference in these mean $\delta^{13}\text{C}$ values across the 48

varieties is up to 2.8 ‰, representing a 44 % variation across the full range of varieties relative to Tinto Cão. Some varieties (e.g., Lillorila and Ugni blanc) appear to operate in a narrower range of $\delta^{13}\text{C}$ values across the years, while others (e.g., Touriga nacional and Saperavi) operate across a wider range of $\delta^{13}\text{C}$. As discussed in the next section, the range of $\delta^{13}\text{C}$ provides an indication of the plasticity of a variety to adapt its WUE to dry versus wet conditions.

An effect is also observed across the years with the lowest and highest mean $\delta^{13}\text{C}$ values for each variety. For years with the most negative $\delta^{13}\text{C}$ values, as associated with wetter conditions and lower WUE during ripening, a difference of up to 3.1 ‰, or a 50 % variation was observed across all varieties relative to the variety with the most negative corresponding $\delta^{13}\text{C}$ value (Saperavi). For years with the least negative $\delta^{13}\text{C}$ values, as associated with drier conditions and higher WUE during ripening, a difference of up to 4.0 ‰, or a 63 % variation was observed across all varieties relative to the variety with the least negative corresponding $\delta^{13}\text{C}$ values (Xynomavro)

In addition to apparent varietal differences, measured $\delta^{13}\text{C}$ may also be impacted by the different phenology of each variety, with later ripening varieties being more likely to experience water deficits during the sugar loading period than earlier varieties. Thus, in attempting to characterize genetically determined varietal differences in $\delta^{13}\text{C}$, differences in the phenology of the berry ripening period in relationship to the driest times of the season must also be considered.

2.3 Categorization based on key traits

Several combinations of key $\delta^{13}\text{C}$ and other physiological traits were used in a hierarchical clustering analysis (HCA) to identify groupings of varieties with similar characteristics. The most consistent groupings were obtained when HCA was applied using the following three traits: “veraison date”, “minimum $\delta^{13}\text{C}$ ” (i.e., $\delta^{13}\text{C}$ in non-limiting conditions) and “range of $\delta^{13}\text{C}$ ” (i.e., the difference between $\delta^{13}\text{C}$ in a wet vintage compared to a dry vintage). The HCA in Figure 4 suggests a range and groupings of varieties based on the above three key traits. All varieties defined traditionally as drought-tolerant (blue arrows) and all varieties defined traditionally as non-drought-tolerant (green arrows) were separated between the two distinct groups. Moreover, varieties generally grown in dry climates were separated from varieties generally grown in more humid climates. While it is very complex to define drought tolerance quantitatively, a consensus could possibly be reached by considering the yield and quality data of the different varieties in the scientific literature and from the wine industry.

Interestingly, the classification based on HCA provided more consistent results with the minimum value of $\delta^{13}\text{C}$ (i.e., non-limiting conditions) compared to the maximum value of $\delta^{13}\text{C}$ (i.e., limiting conditions). Varieties with a more negative $\delta^{13}\text{C}$ in non-limiting conditions may have a greater stomatal conductance and thus greater transpiration and water use early in the season which could more quickly deplete available

soil water (Lebon *et al.*, 2003) and increase the likelihood of water deficits during berry ripening. This hypothesis is supported by the Ψ_{PD} data (Figure 2) where Tempranillo and Semillon, varieties with very relatively negative $\delta^{13}C$ in non-limiting conditions are the two varieties that reached the most negative Ψ_{PD} . The other metric of interest in this HCA classification is the difference between the minimum value of $\delta^{13}C$ (i.e., the $\delta^{13}C$ under non-limiting conditions)

and the maximum value of $\delta^{13}C$ (i.e., the $\delta^{13}C$ under limiting conditions). This trait would characterize the plasticity of the variety to adapt to the drought conditions of the vintage.

One confounding factor in assessing the $\delta^{13}C$ characteristics of different varieties is the timing of phenological stages. Of particular concern with $\delta^{13}C$ in berry juice sugars is the vine water status during the period of sugar loading in the berries, roughly starting one week to ten days before

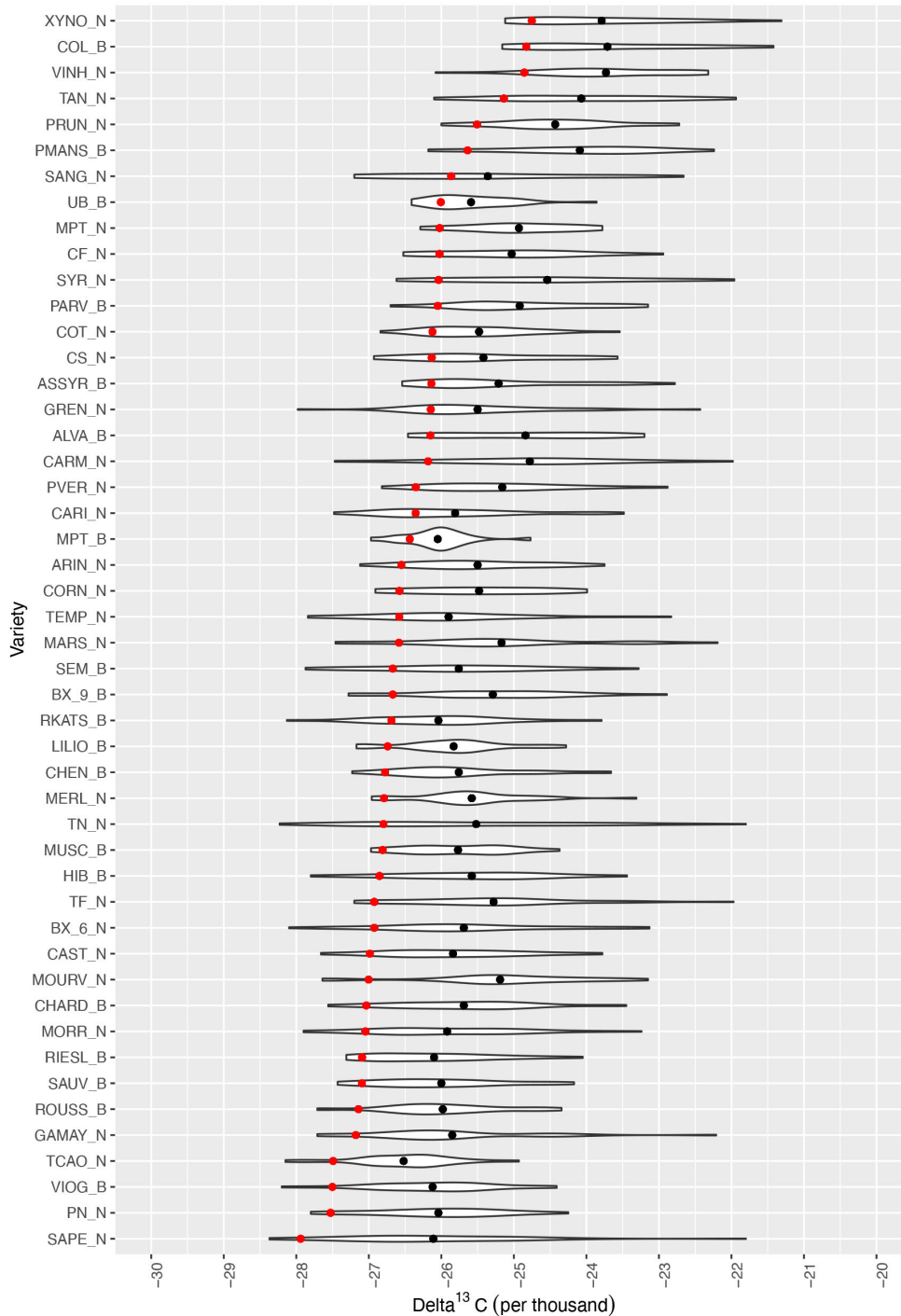


FIGURE 3. Violin plot of the $\delta^{13}C$ data for 48 varieties from 2014 to 2020 in Villenave d’Ornon, France. The black dots represent the average value of $\delta^{13}C$ and the red dots the average value of $\delta^{13}C$ in the wettest year (classified from the less negative to the most negative). The full names of the varieties are presented in Supplementary Materials, Table S1.

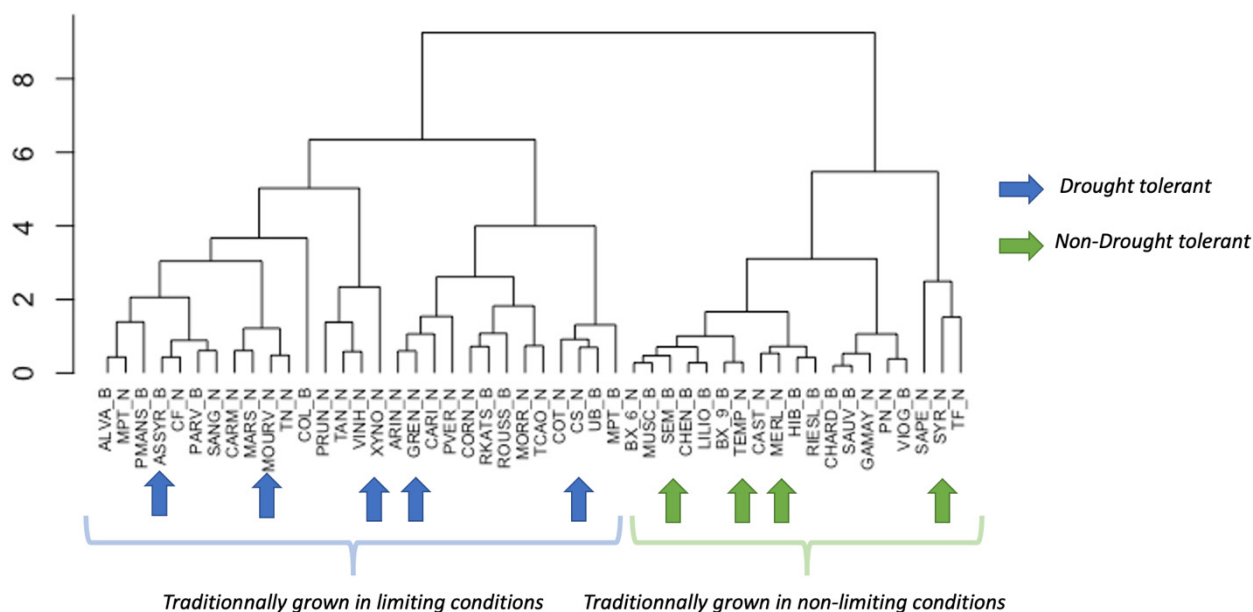


FIGURE 4. Hierarchical Clustering Analyses (HCA) performed with the Ward method of the 48 varieties as a function of “Average Veraison (DOY)”, “Minimum $\delta^{13}\text{C}$ ” and “Range of $\delta^{13}\text{C}$ ”.

mid-veraison and ending three or four weeks after mid-veraison (Suter *et al.*, 2021). In most years, water deficit increases as the season progresses, which means that an early variety faces less water deficit during berry ripening compared to late varieties. Interestingly, adding average mid-veraison dates in the HCA classification improves the separation of drought sensitive versus drought tolerant varieties, as defined by hydroscares in section 3 below. The relationship between the characteristics “late ripening” and “drought tolerant” may be due to the fact that late ripening varieties were selected to grow in warm climates, which are also most often dry climates. In the HCA presented in Figure 4, most varieties grown around Mediterranean basin are classified in the left panel, such as Assyrτικο, Sangiovese, Mourvèdre, Grenache, Carignan, and Rkatsiteli.

3. Hydroscares

3.1. Constructed hydroscares

Hydroscares are presented in Figure 5 for the six varieties using individual vine data for which there was both $\delta^{13}\text{C}$ and corresponding water potential data. Grenache has the smallest hydroscape surface and is known for being very strict in its stomatal regulation (Schultz, 2003). Semillon, the variety with the largest hydroscape, is known for its anisohydric behaviour, due to its poor control over stomatal conductance, leading to substantial nighttime transpiration (Rogiers *et al.*, 2009; Dayer *et al.*, 2020; Dayer *et al.*, 2021). In the same study of Rogiers *et al.*, (2009), Cabernet-Sauvignon and Merlot were not significantly different in terms of stomatal conductance which corresponds with our finding of the two varieties having a similar hydroscape surface (10.7 MPa^2 and 10.9 MPa^2 , respectively).

These results are also consistent with empirical knowledge gathered in production areas. In La Rioja (Spain), Grenache

is planted in the driest parts of the production area around the town of Logroño, while Tempranillo dominates in the cooler locations at higher altitude, where drought- and heat-stress are less frequent. Semillon thrives in the Atlantic climate of Bordeaux (France) and is the major variety in the Hunter Valley (Australia), one of the wettest production areas in the world (Johnson, 2013). The drought responses of Ugni blanc (Trebiano) is less well documented, but these results suggest that it may be well adapted to dry conditions. The hydroscape approach, however, only refers to stomatal behaviour of plants under various levels of water deficit. Drought responses involve other mechanisms like hydraulic conductance, osmotic adjustment, leaf versus root development, duration of the crop cycle, and timing of phenology to ensure development in the wettest part of the season (Tardieu *et al.*, 2018).

3.2. The metrics of the hydroscares

The areas of the hydroscares presented in Figure 5 provide an insight into the stomatal behaviour of plants under drought conditions, but other metrics can also be extracted (see also Figure S3 in supplementary materials). For example, the intercept of the hydroscape represents the least negative Ψ_L observed under non-limiting water availability (least negative Ψ_{pd}). Interestingly, this intercept is very different for Grenache compared to Semillon (-0.96 MPa of Ψ_L and -1.4 MPa of Ψ_L , respectively). This may be explained by differences in stomatal conductance, which was observed as being higher in Semillon than Grenache (Rogiers *et al.*, 2009). The slope of the hydroscape also differs among varieties, with Grenache having the steepest slope compared to the other varieties.

A distinct critical midday leaf water potential (Ψ_{Lcrit}) is more evident for Grenache, Ugni blanc, Cabernet-Sauvignon and Merlot, but less so for Tempranillo and Semillon.

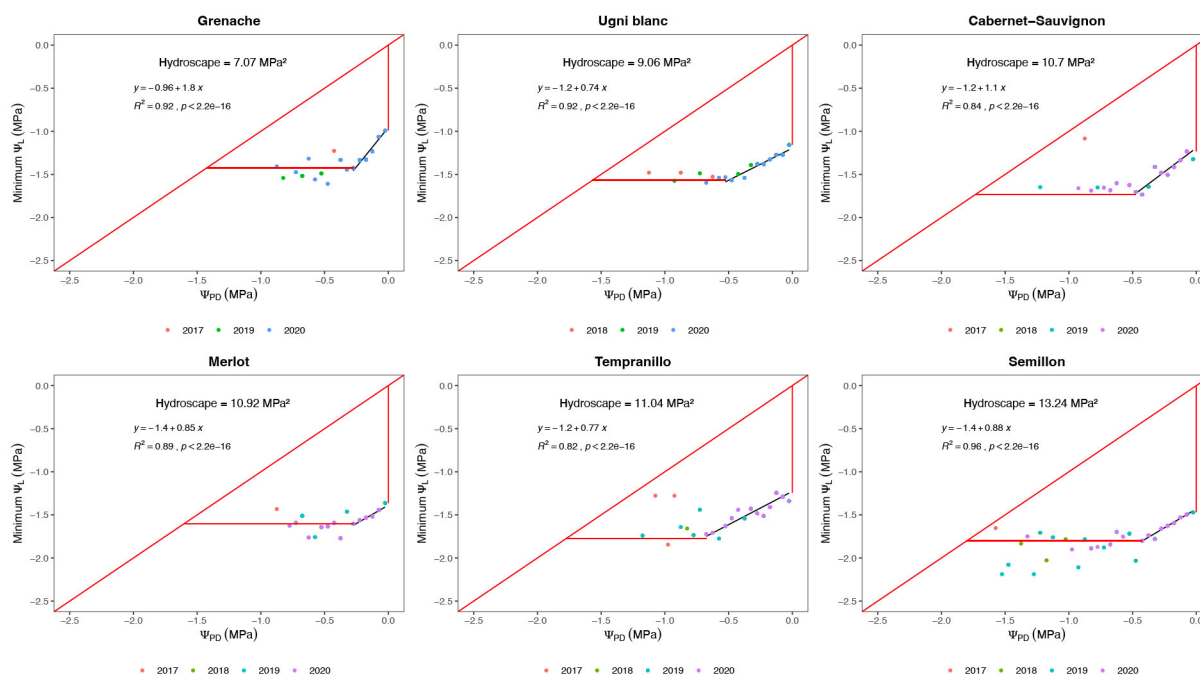


FIGURE 5. Modified hydroscape representation for field data, with their corresponding areas in MPa². Varieties are classified from the smallest to the largest hydroscape. The study was carried out in Villenave d’Ornon in 2017 (except for Ugni blanc), 2018 (except for Grenache), 2019 and 2020.

The variation observed for Tempranillo and Semillon could be due to a variety of factors, including differences in vapour pressure deficit during the precise time of measure, and/or measurement artifacts (e.g., severe levels of stress causing petiole embolisms, hydraulically disconnecting the measured leaf from the vine). There are also differences in the Ψ_{PD} at which Ψ_{Lcrit} is reached. For example, Merlot reaches Ψ_{Lcrit} at $\Psi_{PD} = -0.3$ MPa, while Tempranillo reaches Ψ_{Lcrit} at $\Psi_{PD} = -0.7$ MPa.

The Ψ_{Lcrit} is a very important metric related to the mortality thresholds of various plant species under drought (Choat *et al.*, 2018). Cultivars with high stomatal conductance in well-watered conditions will deplete soil water earlier in the season. Depending on the degree to which stomatal conductance is adjusted under decreasing Ψ_{pp} , these cultivars could face more severe water deficits. For example in this study, Tempranillo had one of the largest hydroscares and most negative critical Ψ_L . It was also observed that among the 48 varieties, Tempranillo was one of the earliest varieties to show leaf wilting and leaf abscission.

Another metric of interest is the range of Ψ_{PD} at which a variety normally operates, representing its ability to extract water at low soil water potential. As discussed above, this can be linked to less stringent stomatal control (e.g., Grenache operates over a small range of Ψ_{PD} because of a very strict stomatal control). Some varieties appear able to extract water at very low Ψ_{PD} which may be due to the root architecture associated with the cultivar x rootstock combination. It is impossible, however, to assess the rooting zone of the vines in field conditions, which may vary from one vine to the other in the VitAdapt parcel due to soil heterogeneity. It is also

possible that the variety influences the ability of the rootstock to explore the soil more or less profoundly (Tandonnet *et al.*, 2010). Vines having access to a greater volume of soil water due to a greater rooting depth are likely to experience less negative values of Ψ_{PD} .

The approach presented offers a possible way of classifying varieties according to their level of (an)isohydricity, an important criterion in drought tolerance (i.e., the larger the hydroscape, the more drought sensitive). Results are in line with the literature regarding stomatal behaviour, in particular for the well documented varieties Grenache and Semillon (Schultz, 2003; Rogiers *et al.*, 2009). Results were obtained on six varieties only and need to be confirmed on a wider range of cultivars. It would also be interesting to assess whether, for a given variety, the hydroscares have a different shape when the data are collected on different soil types, or with different rootstocks. One of the limitations of the hydroscape approach is that it requires time-consuming water potential field data collected, ideally over multiple years.

4. Comparison of $\delta^{13}C$ and hydroscares

Both $\delta^{13}C$ and hydroscares are mainly driven by the stomatal behaviour of plants (Farquhar, 1989; Meinzer *et al.*, 2016). Hydroscares were constructed as described above for six varieties for which there were also measurements of $\delta^{13}C$. Several metrics were extracted from the analysis performed using these two tools and are listed in Table 1 along with the associated values for each variety.

A correlation analysis of the metrics presented in Table 1 across the different varieties is presented in Figure 6. It is observed that the size of the hydroscares is strongly

TABLE 1. Metrics of hydroscares and $\delta^{13}\text{C}$ for six grapevine varieties. Varieties are classified from the lowest to the highest hydroscape area as representative of (an)isohydricity. For each metric, values are classified with shades of green.

Variety	Intercept (MPa)	Slope (MPa)	Ψ_{Lcrit} (MPa)	Range of Ψ_{PD} (MPa)	Hydroscape (MPa ²)	Range of Ψ_{L} (MPa)	Range of $\delta^{13}\text{C}$ (‰)	Max $\delta^{13}\text{C}$ (‰)	Min $\delta^{13}\text{C}$ (‰)
Grenache	-0.96	1.80	-1.43	-0.89	7.07	0.47	4.08	-23.20	-27.28
Ugni blanc	-1.2	0.74	-1.57	-1.12	9.06	0.37	4.06	-23.34	-27.40
Cabernet-Sauvignon	-1.2	1.10	-1.73	-1.23	10.70	0.53	4.65	-23.11	-27.76
Merlot	-1.4	0.85	-1.60	-0.89	10.92	0.20	3.90	-23.70	-27.60
Tempranillo	-1.2	0.77	-1.78	-1.18	11.04	0.58	5.96	-22.35	-28.31
Semillon	-1.4	0.88	-1.80	-1.58	13.24	0.40	5.15	-23.30	-28.45

* Difference on the yaxis between the intercept of the hydroscape and the Ψ_{Lcrit} .

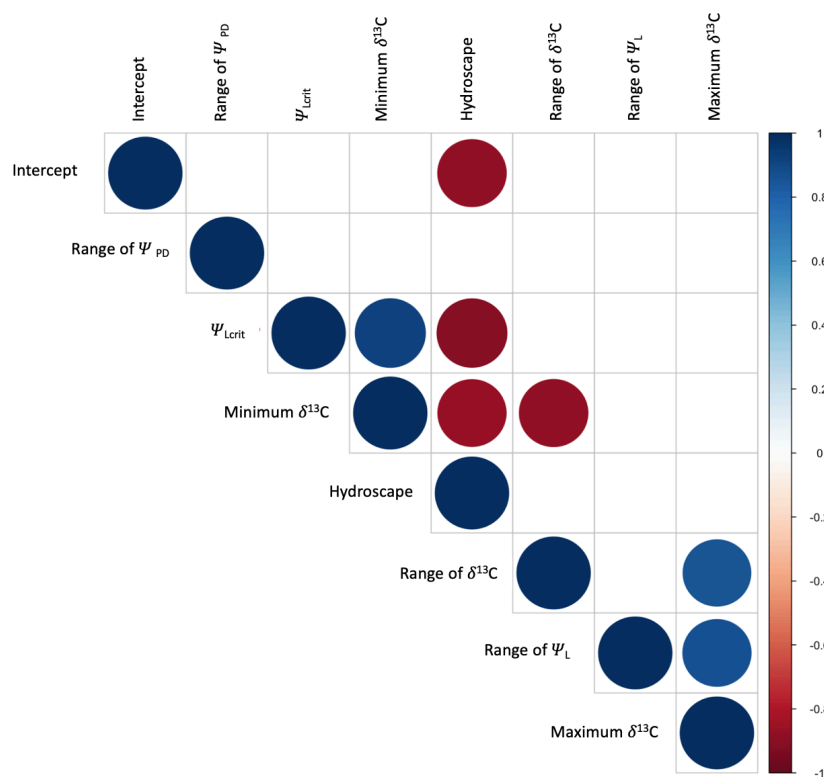


FIGURE 6. A correlation heatmap of the metrics extracted from the hydroscares and from the $\delta^{13}\text{C}$ (as shown in Table 1). Only significant correlations at $P < 0.05$ are indicated.

dependent on two variables, the intercept and the Ψ_{Lcrit} . While the intercept becomes more negative the size of the hydroscape increases, suggesting a generally anisohydric behaviour even at low Ψ_{PD} . A similar dynamic exists with a more negative Ψ_{Lcrit} also being associated with a larger hydroscape.

The minimum $\delta^{13}\text{C}$, which is the most negative $\delta^{13}\text{C}$ obtained in non-limiting conditions, appears better correlated to hydraulic traits from the hydroscares than does the maximum $\delta^{13}\text{C}$, which is the least negative $\delta^{13}\text{C}$ obtained in dry conditions. Minimum $\delta^{13}\text{C}$ is well correlated negatively with the area of the hydroscares (Hydroscape) and positively with Ψ_{Lcrit} , suggesting the anisohydric behaviour of a larger

hydroscape is somehow associated with a lower level of water use efficiency in well-watered conditions (Chaves *et al.*, 2010). Increased water use in non-limiting conditions, as suggested by more negative Ψ_{L} , may lead to greater water deficits later in the season and hence less negative $\delta^{13}\text{C}$.

By integrating the $\delta^{13}\text{C}$ metrics with the insights provided by the hydroscares metrics, minimum $\delta^{13}\text{C}$ appears to be a relatively easily accessible metric to classify grapevine varieties according to their drought responses. It should be noted that, although these metrics are easy to collect in field conditions, they need to be measured over multiple years in homogeneous soil conditions, or with enough replicates to rule-out possible variability in soil water holding

capacity. These conditions were met in the VitAdapt setting (Destrac-Irvine and van Leeuwen, 2017).

CONCLUSION

The measured $\delta^{13}\text{C}$ in berry juice sugars provides an indication of vine water status during the period of sugar accumulation from veraison through maturity, as was confirmed by comparison against corresponding measurements of predawn water potential on six varieties over four seasons. Distinct differences in $\delta^{13}\text{C}$ measured in berry juice sugar collected in a vineyard setting over seven years of differing climatic conditions found significant differences across 48 varieties. The drought responses of these varieties were then ranked using a hierarchical cluster analysis based on: i) the mean $\delta^{13}\text{C}$ in years of relatively low water stress; ii) the range of $\delta^{13}\text{C}$ for a variety as measured in wet versus dry years; and iii) the day of year of mid-veraison. This ranking corresponded well with generally understood drought tolerance characteristics of the different varieties. Of particular importance were the highly variable mid-veraison dates for the different varieties affecting the water deficits experienced by each variety, and hence the corresponding measured $\delta^{13}\text{C}$ in a given year, independent of the genotype response.

The predawn measurements on the six varieties over four years, along with corresponding midday leaf water potential measurements were used to construct hydroscales, which allowed insight into the stomatal behaviour of these varieties. Among several hydroscale metrics, minimum critical leaf water potential (Ψ_{Lcrit}), and the closely related total area of the hydroscale were considered to provide a good indication of drought stress sensitivity. Less negative values of Ψ_{Lcrit} and correspondingly smaller hydroscale areas suggest an isohydric variety, which regulates its stomata more strictly to maintain water potential. Conversely, more negative Ψ_{Lcrit} and larger hydroscales are suggestive of a more anisohydric variety, which regulates its stomata less strictly, allowing for more negative water potentials.

Comparison of $\delta^{13}\text{C}$ and hydroscale metrics found minimum $\delta^{13}\text{C}$ (i.e., from years with less limiting conditions) to be correlated negatively with the area of the hydroscales and positively correlated with Ψ_{Lcrit} , suggesting the anisohydric behaviour of a larger hydroscale is somehow associated with a lower level of water use efficiency. This may be due to increased water use in non-limiting conditions as suggested by more negative Ψ_{L} , resulting in greater water deficits and more stress later in the season and hence less negative $\delta^{13}\text{C}$. This also suggests that this minimum $\delta^{13}\text{C}$ could provide a relatively easy way to classify grapevine varieties according to their drought stress responses.

ACKNOWLEDGMENTS

This study was carried out with support from Jas. Hennessy & Co. (16100 Cognac, France). The VitAdapt Project is supported by the Conseil Interprofessionnel des Vins de Bordeaux (CIVB), the Conseil Régional d'Aquitaine and

the Institut National de Recherche pour l'Agriculture, l'Alimentation et l'Environnement (INRAE). This study has been carried out with financial support from the French National Research Agency (ANR) in the frame of the Investments for the future Programme, within the Cluster of Excellence COTE (ANR-10-LABX-45). The authors would like to thank INRAE, UEVB, F-33882, Villenave d'Ornon, France, for its contribution to the maintenance of the VitAdapt experiment.

REFERENCES

- Álvarez-Maldini, C., Acevedo, M., & Pinto, M. (2021). Hydroscales: A useful metric for distinguishing iso-/anisohydric behaviour in almond cultivars. *Plants*, *10*(6), 1249. <https://doi.org/10.3390/plants10061249>
- Améglio, T., Archer, P., Cohen, M., Valancogne, C., Daudet, F. A., Dayau, S., & Cruiziat, P. (1999). Significance and limits in the use of predawn leaf water potential for tree irrigation. *Plant and Soil*, *207*(2), 155-167. <https://doi.org/10.1023/A:1026415302759>
- Anderson, K., & Aryal, N. R. (2013). *Which winegrape varieties are grown where? A global empirical picture* (p. 700). University of Adelaide Press. <https://doi.org/10.20851/winegrapes>
- Bchir, A., Escalona, J., Galle, A., Hernández-Montes, E., Tortosa, I., Braham, M., & Medrano, H. (2016). Carbon isotope discrimination ($\delta^{13}\text{C}$) as an indicator of vine water status and water use efficiency (WUE): Looking for the most representative sample and sampling time. *Agricultural Water Management*, *167*, 11-20. <https://doi.org/10.1016/j.agwat.2015.12.018>
- Carbonneau, A., Deloire, A., Torregrosa, L., Jaillard, B., Pellegrino, A., Metay, A., Ojeda, H., Lebon, E., & Abbal, P. (2015). *Traité de la vigne: Physiologie, terroir, culture*. (Issue 2ème ed., p. 592 p.). Dunod. <https://hal.archives-ouvertes.fr/hal-02154373>
- Charrier, G., Delzon, S., Domec, J.-C., Zhang, L., Delmas, C. E. L., Merlin, I., Corso, D., King, A., Ojeda, H., Ollat, N., Prieto, J. A., Scholach, T., Skinner, P., van Leeuwen, C., & Gambetta, G. A. (2018). Drought will not leave your glass empty: Low risk of hydraulic failure revealed by long-term drought observations in world's top wine regions. *Science Advances*, *4*(1), eaao6969. <https://doi.org/10.1126/sciadv.aao6969>
- Chaves, M. M., Zarrouk, O., Francisco, R., Costa, J. M., Santos, T., Regalado, A. P., Rodrigues, M. L., & Lopes, C. M. (2010). Grapevine under deficit irrigation: Hints from physiological and molecular data. *Annals of Botany*, *105*(5), 661-676. <https://doi.org/10.1093/aob/mcq030>
- Cifre, J., Bota, J., Escalona, J. M., Medrano, H., & Flexas, J. (2005). Physiological tools for irrigation scheduling in grapevine (*Vitis vinifera* L.): An open gate to improve water-use efficiency?. *Agriculture, Ecosystems & Environment*, *106*(2-3), 159-170. <https://doi.org/10.1016/j.agee.2004.10.005>
- Choat, B., Brodrick, T. J., Brodersen, C. R., Duursma, R. A., López, R., & Medlyn, B. E. (2018). Triggers of tree mortality under drought. *Nature*, *558* (7711), 531-539. <https://doi.org/10.1038/s41586-018-0240-x>
- Choné, X., van Leeuwen, C., Dubourdieu, D., Gaudillère, J. P., (2001). Stem water potential is a sensitive indicator of grapevine water status. *Annals of Botany*, *87*(4), 477-483. <https://doi.org/10.1006/anbo.2000.1361>
- Dayer, S., Herrera, J. C., Dai, Z., Burlett, R., Lamarque, L. J., Delzon, S., Bortolami, G., Cochard, H., & Gambetta, G. A. (2020).

- The sequence and thresholds of leaf hydraulic traits underlying grapevine varietal differences in drought tolerance. *Journal of Experimental Botany*, 71(14), 4333–4344. <https://doi.org/10.1093/jxb/eraa186>
- Dayer, S., Herrera, J. C., Dai, Z., Burlett, R., Lamarque, L. J., Delzon, S., Bortolami, G., Cochard, H., & Gambetta, G. A. (2021). Nighttime transpiration represents a negligible part of water loss and does not increase the risk of water stress in grapevine. *Plant, Cell & Environment*, 44(2), 387–398. <https://doi.org/10.1111/pce.13923>
- Destrac-Irvine, A., & van Leeuwen, C. (2017). VitAdapt : an experimental program to study the behaviour of a wide range of *Vitis vinifera* varieties in a context of climate change in the Bordeaux vineyards. In: proceedings of Climwine, sustainable grape and wine production in the context of climate change, 11-13 April 2016, Bordeaux, p. 165-171.
- Farquhar, G. (1989). Carbon isotope discrimination and photosynthesis. *Annual Review of Plant Physiology and Plant Molecular Biology*, 40(1), 503–537. <https://doi.org/10.1146/annurev.arplant.40.1.503>
- Flexas, J., Escalona, J. M., & Medrano, H. (1998). Down-regulation of photosynthesis by drought under field conditions in grapevine leaves. *Functional Plant Biology*, 25(8), 893-900. <https://doi.org/10.1071/PP98054>
- Fraga, H., Malheiro, A. C., Moutinho-Pereira, J., & Santos, J. A. (2012). An overview of climate change impacts on European viticulture. *Food and Energy Security*, 1(2), 94–110. <https://doi.org/10.1002/fes3.14>
- Gaudillère, J., van Leeuwen, C., & Ollat, N. (2002). Carbon isotope composition of sugars in grapevine, an integrated indicator of vineyard water status. *Journal of Experimental Botany*, 53(369), 757–763. <https://doi.org/10.1093/jexbot/53.369.757>
- Gaudin, R., Roux, S., & Tisseyre, B. (2017). Linking the transpirable soil water content of a vineyard to predawn leaf water potential measurements. *Agricultural Water Management*, 182, 13-23. <https://doi.org/10.1016/j.agwat.2016.12.006>
- Hochberg, U., Rockwell, F. E., Holbrook, N. M., & Cochard, H. (2018). Iso/Anisohydry: A Plant–Environment Interaction Rather Than a Simple Hydraulic Trait. *Trends in Plant Science*, 23(2), 112–120. <https://doi.org/10.1016/j.tplants.2017.11.002>
- Johnson, H.J. R. (2013). *The World Atlas of Wine, 7th Edition*. Mitchell Beazley.
- Keller, M. (2010). *The Science of Grapevines—1st Edition*. Elsevier Science Publishing Co Inc, Academic Press Inc.
- Klein, T. (2014). The variability of stomatal sensitivity to leaf water potential across tree species indicates a continuum between isohydric and anisohydric behaviours. *Functional Ecology*, 28(6), 1313–1320. <https://doi.org/10.1111/1365-2435.12289>
- Lebon, E., Dumas, V., & Pieri, P. (2003). Modelling the seasonal dynamics of the soil water balance of vineyards. *Functional Plant Biology*, 30, 12. <https://doi.org/10.1071/FP02222>
- Le Menn, N., van Leeuwen, C., Picard, M., Riquier, L., Revel, G., & Marchand, S. (2019). Effect of Vine Water and Nitrogen Status, as Well as Temperature, on Some Aroma Compounds of Aged Red Bordeaux Wines. *Journal of Agricultural and Food Chemistry*, 67. <https://doi.org/10.1021/acs.jafc.9b00591>
- Levin, A. D., Williams, L. E., Matthews (2019). A continuum of stomatal responses to water deficits among 17 wine grape cultivars (*Vitis vinifera*). *Functional Plant Biology*, 47(1), 11–25. <https://doi.org/10.1071/FP19073>
- Li, X., Blackman, C. J., Peters, J. M. R., Choat, B., Rymer, P. D., Medlyn, B. E., & Tissue, D. T. (2019). More than iso/anisohydry: Hydroscares integrate plant water use and drought tolerance traits in 10 eucalypt species from contrasting climates. *Functional Ecology*, 33(6), 1035–1049. <https://doi.org/10.1111/1365-2435.13320>
- Martínez-Vilalta, J., Poyatos, R., Aguadé, D., Retana, J., & Mencuccini, M. (2014). A new look at water transport regulation in plants. *New Phytologist*, 204(1), 105–115. <https://doi.org/10.1111/nph.12912>
- Medrano, H., Flexas, J., Ribas-Carbó, M., & Gulías, J. (2010). Measuring Water Use Efficiency in Grapevines. In S. Delrot, H. Medrano, E. Or, L. Bavaresco, & S. Grando (Eds.), *Methodologies and Results in Grapevine Research* (pp. 123–134). Springer Netherlands. https://doi.org/10.1007/978-90-481-9283-0_9
- Meinzer, F. C., Woodruff, D. R., Marias, D. E., Smith, D. D., McCulloh, K. A., Howard, A. R., & Magedman, A. L. (2016). Mapping ‘hydroscares’ along the iso- to anisohydric continuum of stomatal regulation of plant water status. *Ecology Letters*, 19(11), 1343–1352. <https://doi.org/10.1111/ele.12670>
- Ollat, N., Touzard, J.-M., & van Leeuwen, C. (2016). Climate Change Impacts and Adaptations: New Challenges for the Wine Industry. *Journal of Wine Economics*, 11(01), 139–149. <https://doi.org/10.1017/jwe.2016.3>
- Roby, G., Harbertson, J. F., Adams, D. A., & Matthews, M. A. (2004). Berry size and vine water deficits as factors in winegrape composition: Anthocyanins and tannins. *Australian Journal of Grape and Wine Research*, 10(2), 100–107. <https://doi.org/10.1111/j.1755-0238.2004.tb00012.x>
- Rogiers, S. Y., Greer, D. H., Hutton, R. J., & Landsberg, J. J. (2009). Does night-time transpiration contribute to anisohydric behaviour in a *Vitis vinifera* cultivar? *Journal of Experimental Botany*, 60(13), 3751–3763. <https://doi.org/10.1093/jxb/erp217>
- Santesteban, L. G., Miranda, C., & Royo, J. B. (2009). Effect of water deficit and rewatering on leaf gas exchange and transpiration decline of excised leaves of four grapevine (*Vitis vinifera* L.) cultivars. *Scientia Horticulturae*, 121(4), 434–439. <https://doi.org/10.1016/j.scienta.2009.03.008>
- Santesteban, L. G., Miranda, C., Urretavizcaya, I., & Royo, J. B. (2012). Carbon isotope ratio of whole berries as an estimator of plant water status in grapevine (*Vitis vinifera* L.) cv. ‘Tempranillo.’ *Scientia Horticulturae*, 146, 7–13. <https://doi.org/10.1016/j.scienta.2012.08.006>
- Scholander, P. F., Bradstreet, E. D., Hemmingsen, E. A., & Hammel, H. T. (1965). Sap Pressure in Vascular Plants. *Science*, 148(3668), 339–346. <https://doi.org/10.1126/science.148.3668.339>
- Schultz, H. R. (2003). Differences in hydraulic architecture account for near-isohydric and anisohydric behaviour of two field-grown *Vitis vinifera* L. cultivars during drought. *Plant, Cell & Environment*, 26(8), 1393–1405. <https://doi.org/10.1046/j.1365-3040.2003.01064.x>
- Souza, D., R, C., Maroco, J. P., Santos, D., P, T., Rodrigues, M. L., Lopes, C. M., Pereira, J. S., & Chaves, M. M. (2005). Impact of deficit irrigation on water use efficiency and carbon isotope composition ($\delta^{13}\text{C}$) of field-grown grapevines under Mediterranean climate. *Journal of Experimental Botany*, 56(418), 2163–2172. <https://doi.org/10.1093/jxb/eri216>
- Suter, B., Destrac-Irvine, A., Gowdy, M., Dai, Z., & van Leeuwen, C. (2021). Adapting Wine Grape Ripening to Global Change Requires a Multi-Trait Approach. *Frontiers in Plant Science*, 12, 624867. <https://doi.org/10.3389/fpls.2021.624867>
- Tandonnet, J.-P., Cookson, S., Vivin, P., & Ollat, N. (2010). Scion genotype biomass allocation and root development in grafted grapevine. *Australian Journal of Grape and Wine Research*, 16, 290–300. <https://doi.org/10.1111/j.1755-0238.2009.00090.x>

- Tardieu, F., & Simonneau, T. (1998). Variability among species of stomatal control under fluctuating soil water status and evaporative demand : modelling isohydric and anisohydric behaviours. *Journal of Experimental Botany*, 49(Special), 419-432. https://doi.org/10.1093/jxb/49.special_issue.419
- Tardieu, F., Simonneau, T., & Muller, B. (2018). The physiological basis of drought tolerance in crop plants : a scenario-dependent probabilistic approach. *Annual Review of Plant Biology*, 69(1), 733-759. <https://doi.org/10.1146/annurev-arplant-042817-040218>
- Tomás, M., Medrano, H., Brugnoli, E., Escalona, J. m., Martorell, S., Pou, A., Ribas-Carbó, M., & Flexas, J. (2014). Variability of mesophyll conductance in grapevine cultivars under water stress conditions in relation to leaf anatomy and water use efficiency. *Australian Journal of Grape and Wine Research*, 20(2), 272–280. <https://doi.org/10.1111/ajgw.12069>
- Triolo, R., Roby, J. P., Pisciotto, A., Di Lorenzo, R., & van Leeuwen, C. (2019). Impact of vine water status on berry mass and berry tissue development of Cabernet franc (*Vitis vinifera* L.), assessed at berry level. *Journal of the Science of Food and Agriculture*, 99(13), 5711–5719. <https://doi.org/10.1002/jsfa.9834>
- van Leeuwen, C., & Darriet, P. (2016). The Impact of Climate Change on Viticulture and Wine Quality. *Journal of Wine Economics*, 11(01), 150–167. <https://doi.org/10.1017/jwe.2015.21>
- van Leeuwen, C., Tregoat, O., Choné, X., Bois, B., Pernet, D., Gaudillère, J.-P., & others. (2009). Vine water status is a key factor in grape ripening and vintage quality for red Bordeaux wine. How can it be assessed for vineyard management purposes. *J. Int. Sci. Vigne Vin*, 43(3), 121–134. <https://doi.org/10.20870/oeno-one.2009.43.3.798>
- van Leeuwen, C., Pieri, P., Gowdy, M., Ollat, N., & Roby, J. P. (2019a). Reduced density is an environmental friendly and cost effective solution to increase resilience to drought in vineyards in a context of climate change. *OENO One*, 53(2), 129-146. <https://doi.org/10.20870/oeno-one.2019.53.2.2420>
- van Leeuwen, C., Destrac-Irvine, A., Dubernet, M., Duchêne, E., Gowdy, M., Marguerit, E., Pieri, P., Parker, A., de Risséguier, L., & Ollat, N. (2019b). An update on the impact of climate change in viticulture and potential adaptations. *Agronomy*, 9(9), 514. <https://doi.org/10.3390/agronomy9090514>
- van Leeuwen, C., Barbe, J. C., Darriet, P., Geffroy, O., Gomès, E., Guillaumie, S., Helwi, P., Laboyrie, J., Lytra, G., Le Menn, N., Marchand, S., Picard, M., Pons, A., Schüttler A. & Thibon, C. (2020). Recent advancements in understanding the terroir effect on aromas in grapes and wines: This article is published in cooperation with the XIIIth International Terroir Congress November 17-18 2020, Adelaide, Australia. Guest editors: Cassandra Collins and Roberta De Bei. *OENO One*, 54(4), 985-1006. <https://doi.org/10.20870/oeno-one.2020.54.4.3983>
- Wickham, H., François, R., Henry, L., Müller, K., & RStudio. (2021a). *Dplyr : A Grammar of Data Manipulation* (1.0.7) [Logiciel]. <https://cran.r-project.org/web/packages/dplyr/index.html>
- Wickham, H., Chang, W., Henry, L., Pedersen, T. L., Takahashi, K., Wilke, C., Woo, K., Yutani, H., Dunnington, D., & RStudio. (2021b). *Ggplot2 : Create elegant data visualisations using the grammar of graphics* (3.3.5) [Logiciel]. <https://cran.r-project.org/web/packages/ggplot2/index.html>
- Williams, L., & Araujo, F. J. (2002). Correlations among Predawn Leaf, Midday Leaf, and Midday Stem Water Potential and their Correlations with other Measures of Soil and Plant Water Status in *Vitis vinifera*. *Journal of the American Society for Horticultural Science*, 127, 448–454. <https://doi.org/10.21273/JASHS.127.3.448>
- Winkel, T., & Rambal, S. (1990). Stomatal conductance of some grapevines growing in the field under a Mediterranean environment. *Agricultural and Forest Meteorology*, 51(2), 107-121. [https://doi.org/10.1016/0168-1923\(90\)90010-4](https://doi.org/10.1016/0168-1923(90)90010-4)