# Relationships among Ambient Temperature and Vapor Pressure Deficit and Leaf and Stem Water Potentials of Fully Irrigated, Field-Grown Grapevines

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**Abstract:** Four *Vitis vinifera* L. cultivars grown at five locations throughout California were studied to determine the relationships among temperature and vapor pressure deficit (VPD) and leaf water potential ( $\Psi_1$ ) measured under clear skies at midday (solar noon) or in some instances midmorning to midafternoon. Stem water potential ( $\Psi_{\text{stem}}$ ) was also measured on several occasions. Vines were irrigated at 100% or greater of measured or estimated vine-yard evapotranspiration, and deficit or nonirrigated vines were included for comparison. Temperature and VPD were determined at the time of measurement. The highest and lowest values of  $\Psi_1$  measured on well-watered grapevines were -0.51 and -1.15 MPa, respectively. Leaf and stem water potentials were linearly related to VPD and ambient temperature. The coefficient of determination was greater for the relationship between  $\Psi_1$  and VPD ( $R^2 = 0.74$ ) than ambient temperature ( $R^2 = 0.58$ ). Based on the regressions, estimates of  $\Psi_1$  at a VPD of 2 and 5 kPa for fully irrigated grapevines would be -0.65 and -0.89, respectively, while those of  $\Psi_{\text{stem}}$  at the same VPDs would be -0.37 and -0.57 MPa, respectively. Leaf water potential of water-stressed vines was less responsive to VPD or temperature when  $\Psi_1$  values ranged from -1.2 to -1.45 MPa. The values of  $\Psi_1$  and  $\Psi_{\text{stem}}$  as a function of VPD or temperature could serve as baselines indicating whether grapevines are fully irrigated or not water stressed under the environmental conditions found in semiarid grapegrowing regions.

**Key words:** leaf water potential, stem water potential, grapevines, vapor pressure deficit, nonstressed baseline

Plant-based measurements as a tool in irrigation management are growing in popularity in the California grape industry. The measurement of vine water status is used to determine when to initiate irrigations early in the growing season and the frequency of water applications once irrigation has begun (L.E. Williams, personal observation). Grapevine water status is determined by measuring leaf  $(\Psi_1)$  or stem  $(\Psi_{\text{stem}})$  water potential at midday or a few hours before or after midday.

Leaf water potential of grapevines undergoes diurnal changes (Williams et al. 1994), with daily minimum values occurring when vine water use is greatest or shortly thereafter (Williams et al. 2003b, Williams and Ayars 2005a) because it is affected by environmental conditions,

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including light, temperature, and vapor pressure deficit (VPD) (van Zyl 1987, Smart and Barrs 1973). Environmental effects on plant water status extend to other plant species. Grimes et al. (1987) found that  $\Psi_1$  of cotton was a linear function of both VPD and ambient temperature when measured on a diurnal basis. McCutchan and Shackel (1992) found that leaf and stem water potentials of prune during midseason were more highly correlated with VPD than with relative humidity, solar radiation, ambient temperature, or wind speed at midday. Williams and Trout (2005) reported similar relationships between VPD and  $\Psi_1$ or  $\Psi_{\text{stem}}$  of grapevine using a limited data set. The value of measured Ψ<sub>1</sub> will vary depending on environmental conditions at the time of measurement. However, light could be eliminated as a major factor if  $\Psi_1$  measurements were taken on cloudless days and if photon flux density (PFD) measured perpendicular to the leaf blade was greater than 1500 µmol m<sup>-2</sup> s<sup>-1</sup>.

The relationships between either  $\Psi_1$  (Grimes et al. 1987) or  $\Psi_{\text{stem}}$  (McCutchan and Shackel 1992, Shackel et al. 1997) and VPD have been used to establish a nonstressed baseline for those two parameters in cotton and deciduous fruit trees, respectively. In the former study (Grimes et al. 1987), cotton  $\Psi_1$  was "climate-normalized" for scheduling an irrigation event. In the latter study (Shackel et al. 1997), the relationship between  $\Psi_{\text{stem}}$  and VPD was designated the fully irrigated baseline. The ability to determine if grapevines are not water stressed, similar to a fully irrigated baseline, would give growers another tool to develop an objective irrigation management program.

This study was conducted to determine the relationships among  $\Psi_{\rm l}$ , and to a lesser extent  $\Psi_{\rm stem}$ , and ambient temperature and VPD for grapevines irrigated at 100% of measured evapotranspiration (ET<sub>c</sub>) or slightly greater than estimated ET<sub>c</sub>. Four different cultivars growing in vineyards at five locations in California were studied. At one location, water applications to replenish ET<sub>c</sub> were determined with the use of a weighing lysimeter (Williams et al. 2003b). Crop coefficients and reference evapotranspiration (ET<sub>c</sub>) were used at the other four locations to estimate vineyard ET<sub>c</sub>. Vines were irrigated at 1.12, 1.2, or 1.25 times ET<sub>c</sub>. All measurements were made on cloudless days to minimize the effects of light on  $\Psi_{\rm l}$  or  $\Psi_{\rm stem}$ .

## Materials and Methods

A weighing lysimeter was installed in 1986 at the University of California Kearney Agricultural Center located in the San Joaquin Valley (lat. 36°48'N; long. 119°30'W). Two *Vitis vinifera* L. (cv. Thompson Seedless clone 2A) grapevine cuttings were planted in the lysimeter and in the vineyard surrounding the lysimeter on 9 April 1987. Vine and row spacings were 2.15 and 3.51 m, respectively (7.55 m² per vine). The length allocated to the canopies of the two vines within the lysimeter was similar to that of the vines down each row in the vineyard surrounding the lysimeter. Row direction was 6° north of the east/west axis. The vineyard was ~1.4 ha (168 x 82 m).

Vines were supported on a trellis that consisted of a 2.13-m wooden stake driven 0.45 m into the soil at each vine. A 0.6-m cross arm was placed atop the stake and wires attached at either end of the cross arm to support the fruiting canes. The trellis for the vines in the lysimeter was self-contained to ensure it was part of the lysimeter mass.

The soil container of the lysimeter was 2 m x 4 m x 2 m deep. The tank was weighed with a balance beam and load-cell configuration, with most of the weight eliminated using counter weights. A detailed description of the lysimeter is given elsewhere (Williams et al. 2003a,b, Williams and Ayars 2005a).

Vines within the lysimeter and the surrounding vineyard were irrigated with 4 L h<sup>-1</sup> in-line drip emitters, spaced every 0.30 m in the vine row. Drip tubing was attached to a wire suspended 0.4 m aboveground. The lysimeter was weighed hourly to determine ET of the two vines and was irrigated when the decrease in mass exceeded a 16 kg (a volume of 8 L vine<sup>-1</sup>) threshold value. The number of irrigations per day throughout the growing season ranged from 0 to 7.

The irrigation pump for the rest of the vineyard was controlled by the lysimeter's datalogger (21x Micrologger, Campbell Scientific, Logan, UT). When the lysimeter was irrigated, the vineyard pump was activated to irrigate the field. Vines were irrigated with the amount of water the lysimeter vines used. In-line water meters downstream from the solenoid valves in each row measured actual ap-

plied water amounts. In another treatment vines were not irrigated. Data used in this study were collected in 2005.

The second site was a Merlot vineyard in Madera County (lat. 36°55'N; long. 120°9'W). The vines were planted on their own roots with 2.13 and 3.66 m vine and row spacings, respectively. The trellis was a cordon wire at a height of 1.28 m and a foliage catch wire 0.3 m above that. Vineyard rows were approximately east/west. Vines were drip-irrigated at either 0.4 or 1.2 of estimated ET<sub>a</sub>. The two amounts of water were achieved using different numbers of emitters or emitters with different discharge rates. The seasonal crop coefficients (K<sub>s</sub>) used to schedule irrigations at this site were developed in the previous four years by measuring the shade cast on the ground beneath the canopy and then using the relationship between the percentage of shade and the crop coefficient (Williams and Ayars 2005b). Reference ET (ET<sub>o</sub>) was obtained from California Irrigation Management Information System (CIMIS) weather station 145, located ~15 km from the vineyard. Vineyard ET<sub>c</sub> was calculated as ET<sub>o</sub> x K<sub>c</sub>. Irrigation treatments did not commence until midday  $\Psi_1$  reached -1.0 MPa. Vines were irrigated once weekly, beginning on Friday and ending by Sunday, with applied water amounts equal to that required for the week. Water potential readings were generally measured on Thursday. Data from the 2002 to 2005 growing seasons were used.

The third site was a Cabernet Sauvignon vineyard in Livermore Valley (lat. 37°40'N. long. 121°46'W). Vines were grafted onto 5C rootstock. Two trellis/training treatments were used: vertical shoot-positioned (VSP) and Smart-Dyson. Vine and row spacings were 1.83 m, and row direction was approximately north/south. Irrigation amounts were 1.12 of estimated ET<sub>c</sub> and applied with a drip-irrigation system. The seasonal crop coefficients for the VSP trellis were developed in a Chardonnay vineyard in Napa Valley (L.E. Williams, unpublished data), adjusted for a row width of 1.83 m and adjusted with the shaded area technique (Williams and Ayars 2005b). The seasonal crop coefficients for the Smart-Dyson trellis/training system at a row width of 1.83 m were established using the shaded area technique in 2000. Reference ET was obtained from a CIMIS weather station located 30 km from the vineyard. Calculation of applied water amounts was similar to that described for the Merlot site, and vines were irrigated 1 to 3 times weekly, depending on the required amounts. Data used were collected in 2002 and 2003. All Ψ<sub>1</sub> readings were taken on the west side of the canopy between 1330 and 1600 hr, Pacific Daylight Time (PDT).

The fourth vineyard site, located near Paso Robles (lat. 35°41'N; 120°39'W), was planted to Cabernet Sauvignon grafted onto 5C. Vine and row spacings in the vineyard were 1.83 and 3.05 m, respectively, with a VSP trellis/training system. Row direction was approximately north/south. In one treatment, vines were irrigated at 1.12 of estimated ET<sub>c</sub>. A second treatment consisted of irrigating vines once every two weeks with 90.7 L (16.2 mm) of water per vine. The seasonal crop coefficients used were those for a VSP

trellis at a row width of 3.05 m. Reference ET was obtained from the PR1 weather station operated by the Paso Robles Wine Country Alliance (PRWCA), located ~3 km from the vineyard. Calculation of applied water amounts was similar to that described for the Merlot site. Vines were irrigated 1 to 3 times weekly, depending upon the required amounts. Data used from this site were collected in 2002 and 2005. Measurements of  $\Psi_1$  were taken on the east side of the canopy in the morning hours (no earlier than 1000 hr PDT) and on the west side in the afternoon (no later than 1600 hr PDT).

The fifth vineyard was located in the Temecula Valley (lat. 33°33'N; long. 117°02'W) of southern California. Chardonnay grapevines on their own roots were planted to vine and row spacings of 2.44 and 3.66 m, respectively. The vines were trained to quadrilateral cordons with a 1.0-m cross arm. Rows were oriented north/south. The seasonal crop coefficients used to schedule irrigations were developed at the site in 1997. The shaded area technique, as in the Merlot vineyard, was used in 1998 and 1999. Reference ET was obtained from CIMIS weather station 137 located ~5 km from the vineyard. Once irrigations began the vines received water five days a week at 1.25 of estimated ET<sub>c</sub> to ensure they were well watered. Data were collected in 1998 and 1999.

Variables measured and calculations used to determine hourly and daily ET<sub>o</sub> from CIMIS can be found in Synder and Pruitt (1992). Degree-day data were obtained from the University of California Statewide Integrated Pest Management Project website (www.ipm.ucdavis.edu). Degree days were calculated using the sine method with a lower threshold of 10°C. Temperature data used in calculating degree days were obtained from the CIMIS (or PRWCA) weather station nearest to the vineyard site.

Water potential readings at all locations were measured as described by Williams and Araujo (2002). Specifically, leaf  $(\Psi_I)$  and stem  $(\Psi_{stem})$  water potentials were measured with a pressure chamber (model 1000; PMS Instrument, Corvallis, OR). Leaf  $\Psi$  was measured on fully expanded, mature leaves exposed to direct solar radiation located on the outside of the canopy. Leaf blades for Ψ, determinations were covered with a plastic bag, quickly sealed, and petioles then cut within 1 to 2 sec. The time between leaf excision and chamber pressurization was generally less than 10 to 15 sec. Approximately 30 min before measurements, leaves for determination of  $\Psi_{\mbox{\tiny stem}}$  were enclosed in plastic bags covered with aluminum foil. Leaves chosen for  $\Psi_{\mbox{\tiny stem}}$  measurements were of similar age and type as those used for  $\Psi_1$  but were located on the shaded side of the canopy to minimize any possible heating effects. A single leaf from five to six individual vine replicates was measured and used for data analysis. Stomatal conductance (g<sub>s</sub>) was measured with a steady-state diffusion porometer (model 1600; LI-COR, Lincoln, NE) on leaves similar to those used for  $\Psi_1$  measurements. The porometer had been sent to the LI-COR factory each year for recalibration (Turner 1991).

Temperature and relative humidity were measured at all locations with two hand-held temperature/relative humidity probes (model DM-84 Multimeter with MultiMeterMate RH/T probe, A.W. Sperry Inst., Inc., Hauppauge, NY) and on occasions a Pocket Sling Psychrometer (Cole-Parmer, Vernon Hills, IL). The probes were positioned just beneath the canopy of vines trained to a VSP trellis (ensuring they were in the shade) and just below the fruiting zone of vines at the other vineyard sites. The probes were placed at two different locations within each vineyard. Measurements with the sling psychrometer were made between rows at a height of ~2 m. The probes were routinely calibrated in the laboratory and the outputs from the two were within 1°C and 2% relative humidity. Photon flux density (PFD) was measured with a quantum sensor (model LI-190SA; LI-COR) or using the quantum sensor on the diffusion porometer.

Data were analyzed via regression analysis using linear and quadratic terms. Regressions with the best fit are presented. Values of  $\Psi_1$  and  $\Psi_{\text{stem}}$  are the means of five to six individual leaf replicates. The homogeneity of linear regression slopes was tested for all individual vineyard data sets. Data were also analyzed using analysis of variance and means separated using the Tukey-Kramer test. CoStat statistics software (CoHort Software, Monterey, CA) was used for data analysis.

### Results

The vineyards used in this study spanned a wide distance from northern (Livermore) to southern (Temecula) California. The accumulation of degree days and seasonal ET differed only slightly from location to location (Table 1). Estimated or measured seasonal grapevine water use ranged from ~450 to 800 mm with vineyard water use dependent on both row spacing and trellis type and individual vine water use dependent on vine spacing within the row and trellis type. Maximum weekly irrigation requirements during the middle of the growing season ranged from 165 to 333 L vine-1 or 30 to 44 mm. In most cases the amount of water requested in this study to meet vineyard ET in the commercial vineyards was applied by the grower/cooperator. Water potential measurements spanned the period at most locations from before bloom until close to fruit harvest.

All measurements were made on cloudless days, and photon flux density (PFD) was in excess of 1500  $\mu mol\ m^{-2}\ s^{-1}$  at the time  $\Psi_l$  or  $\Psi_{stem}$  was taken. A wide range of temperatures and vapor pressure deficits (VPD) were recorded at all sites when measurements were taken. The high and low temperatures across all locations at the time of data collection were 44.6 and 20.1°C, respectively (Table 2). High and low values of VPD were 8.71 and 1.19 kPa, respectively. Using all data points, VPD was highly correlated with ambient temperature (Ta) in this study. The linear relationship between VPD and temperature was VPD = -5.69 + 0.289 \* Ta (R^2 = 0.87, p < 0.001). A second order

Table 1 Seasonal climatic variables, dates of measurements, and water requirements at the five locations of grapevines in this study.

Location	DDa	ET <sub>o</sub> ª (mm)	Date⁵ (month/day)	ET <sup>a</sup> (L vine <sup>-1</sup> /mm)	Max. applied H <sub>2</sub> O/week <sup>-1a</sup> (L vine <sup>-1</sup> /mm)
Madera	2289°	1289	4/13 to 8/19	5742 / 760	302 / 39
Livermore	2260	1084	5/27 to 9/4	1973 / 589	147 / 44
Kearney	2503	1131	5/24 to 8/24	6044 / 800	333 / 44
Paso Robles	1981	1208	5/6 to 8/21	2536 / 458	165 / 30
Temecula	2121	1072	5/20 to 9/17	5288 / 607	293 / 33

<sup>&</sup>lt;sup>a</sup>Degree days (DD), reference ET (ET<sub>o</sub>), and measured (Kerney) or estimated (other locations) vineyard evapotranspiration (ET<sub>o</sub>) are from 15 March to 31 Oct at each location. Maximum weekly, applied water amounts refer to water use at 100% of ET<sub>o</sub>.

Table 2 Low and high values of temperature, vapor pressure deficit (VPD), and leaf water potential (Ψ<sub>1</sub>) measured at each location throughout the course of the study.

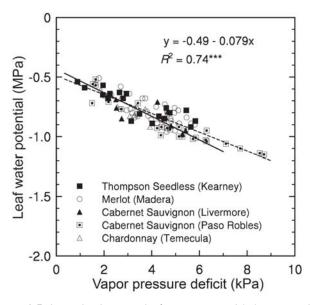
	Temp (°C)		VPD (kPa)		Ψ <sub>ι</sub> (MPa)	
Location	Low	High	Low	High	Low	High
Madera	20.1	39.9	1.78	5.65	-0.97	-0.64
Livermore	28.5	37.1	2.52	5.42	-0.98	-0.71
Kearney	22.7	39.4	1.19	5.83	-0.92	-0.54
Paso Robles	22.0	44.6	1.50	8.71	-1.14	-0.53
Temecula	22.2	35.6	1.65	3.93	-0.92	-0.61

polynomial regression between VPD and  $T_a$  (3.34 - 0.284 \*  $T_a$  + 0.00886 \*  $T_a$ <sup>2</sup>) increased the R<sup>2</sup> to 0.91.

Differences in the high and low values of  $\Psi_1$  at the various locations were determined by taking measurements at different times during the growing season or at different times during the day. The highest and lowest  $\Psi_1$  values were -0.53 and -1.14 MPa, both measured at the Paso Robles site (Table 2). The lowest values at the other locations ranged from -0.92 to -0.98 MPa.

The relationship between  $\Psi_1$  and VPD did not differ among the five locations. The best fit of the relationships between  $\Psi_1$  and VPD or temperature for the entire data set was a linear function (Figure 1, Figure 2). The coefficient of determination was greater for the regression of  $\Psi_1$  on VPD than for the regression of  $\Psi_1$  and temperature. While fewer measurements of  $\Psi_{\text{stem}}$  were taken during the course of the study, a linear relationship was detected between  $\Psi_{\text{stem}}$  and VPD (Figure 3). The relationship between  $\Psi_{\text{stem}}$  and temperature (y = -0.118 - 0.0185 \* VPD) had a  $R^2$  value of 0.49 (p < 0.01).

On several occasions  $g_s$  and leaf transpiration (E) were measured with  $\Psi_1$  (Table 3). In general,  $g_s$  increased after the first measurement of the day, was highest around solar noon, and slightly decreased thereafter. Values of  $g_s$  for the fully irrigated vines were always greater than those of vines that were deficit irrigated or not irrigated at all. Leaf E values of vines irrigated at 1.0 of  $ET_c$  or greater were highest at solar noon or later. Leaf  $\Psi$  of the deficit-irrigated vines (Cabernet Sauvignon and Merlot) tended to



**Figure 1** Relationship between leaf water potential ( $\Psi_{\rm l}$ ) measured on four grapevine cultivars and vapor pressure deficit (VPD) at the time of measurement at five locations. The solid line represents the relationship between  $\Psi_{\rm l}$  and VPD (Williams and Trout 2005). Each data point is the mean of at least five individual leaf replicates (\*\*\* indicates significance at p < 0.001; n = 90).

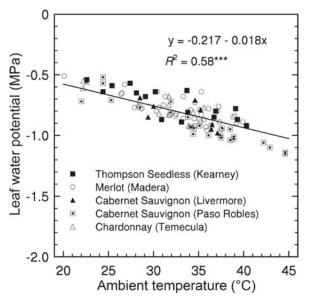
decrease during the day, while that of the nonirrigated Thompson Seedless vines remained similar from 1000 to 1600 hr. The relationship between  $\Psi_1$  values lower than -1.2 MPa and VPD was also best fit with a linear regression (Figure 4). However, the slope of that relationship differed from that between  $\Psi_1$  and VPD for vines irrigated at 1.0 ET<sub>c</sub> or greater (from Figure 1).

## **Discussion**

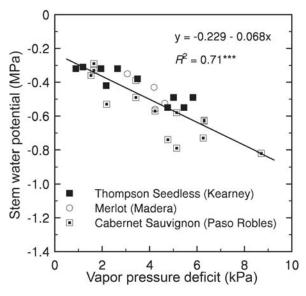
Previous studies on grapevine have examined the response of  $\Psi_1$  to various environmental parameters on a diurnal basis (Smart 1974, Smart and Barrs 1973, Stevens et al. 1995, van Zyl 1987). The environmental factor having the most influence on  $\Psi_1$  in those studies was PFD and to a lesser extent VPD and temperature. The purpose of our study was to examine the response of  $\Psi_1$  and  $\Psi_{\text{stem}}$ 

Date column represents the first and last dates at which water potential data were measured at each location for use in this study.

eValues in columns at locations in which data were collected more than a single growing season represent the means of all seasons.



**Figure 2** The relationship between leaf water potential and ambient temperature at the time of measurement. Other information is as given in Figure 1.



**Figure 3** The relationship between stem water potential ( $\Psi_{\text{stem}}$ ) measured on three grapevine cultivars and vapor pressure deficit (VPD) at the time of measurement. Other information is as given in Figure 1 (n = 28).

Table 3 Time of day, temperature, vapor pressure deficit (VPD), leaf water potential (Ψ<sub>i</sub>), stomatal conductance (g<sub>s</sub>), and leaf transpiration (E) measured at that particular time for three cultivars. Vines were irrigated at various fractions of estimated ET<sub>c</sub> (1.12, 1.2, and 0.4) or measured ET<sub>c</sub> with a weighing lysimeter.

Time (hr)	Temp (°C)	VPD (kPa)	Ψ <sub> </sub> (MPa)	<b>g</b> <sub>s</sub> (mmol m <sup>-2</sup> s <sup>-1</sup> )	E (mmol m <sup>-2</sup> s <sup>-1</sup> )	Ψ <sub>ι</sub> (MPa)	<b>g</b> <sub>s</sub> (mmol m <sup>-2</sup> s <sup>-1</sup> )	E (mmol m <sup>-2</sup> s <sup>-1</sup> )		
Paso Rob	les (24 Jul 2002	2) Cabernet Sa	uvignona							
				1.12 ET <sub>c</sub>			Dry down			
1000	22.2	1.50	-0.72 a <sup>b</sup>	437 c	8.6 c	-1.28 a	182	4.8 b		
1200	28.5	2.72	-0.77 a	575 a	17.3 b	-1.33 ab	175	6.1 b		
1400	34.4	4.38	-0.99 b	524 ab	23.1 a	-1.42 bc	146	8.5 a		
1600	36.1	5.14	-1.00 b	483 bc	22.2 a	-1.45 c	145	9.5 a		
Madera (5	Jul 2002) Merlo	ot°								
				1.20 ET <sub>c</sub>			0.4 ET <sub>c</sub>			
1100	27.9	2.67	-0.77 a	590 b	16.1 c	-1.21 a	390 a	11.6		
1300	32.5	3.85	-0.83 b	728 a	24.3 b	-1.22 a	353 a	14.1		
1500	35.6	4.82	-0.87 b	756 a	29.8 a	-1.29 b	194 b	11.1		
Kearney A	Ag Center (24 A	ug 2005) Thon	npson Seedles	<b>S</b> <sup>d</sup>						
				1.0 ET <sub>c</sub>			No applied water			
1000	29.9	2.81	-0.68 a	448 b	12.7 b	-1.32	140 a	4.5		
1300	35.5	4.30	-0.83 c	587 a	20.1 a	-1.32	116 ab	5.5		
1600	35.3	4.61	-0.76 b	473 b	19.1 a	-1.31	100 b	5.2		

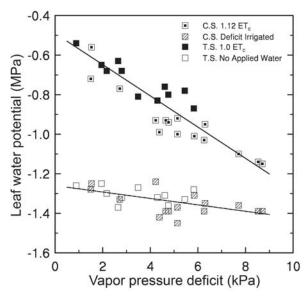
<sup>&</sup>lt;sup>a</sup>Vines in the 1.12 ET<sub>c</sub> treatment had been irrigated with 42 L (total week's irrigation requirement at 100% of ET<sub>c</sub> was 91 L vine<sup>-1</sup>) the night before measurements were taken while the dry-down treatment (irrigated at 0.56 of ET<sub>c</sub>) had not been irrigated for two weeks.

measured at midday to changes in ambient VPD and temperature at the time of measurement, as it is at midday that daily minimum values of either parameter are expected, even for well-watered vines (Grimes and Williams 1990). During the daily measurement period, between 1000 and 1600 hr (PDT) on cloudless days, it was assumed that

 $<sup>^{</sup>b}$ Values within a column for each cultivar followed by a different letter are significantly different at p < 0.05. Values within a column not followed by any letters are not significantly different.

<sup>°</sup>Vines in the 1.2 and 0.4 irrigation treatments had been irrigated with the entire week's irrigation requirement of 326 and 109 L vine<sup>-1</sup> (irrigation amount at 100% of ET<sub>c</sub> was 272 L vine<sup>-1</sup>), respectively, the weekend (30 and 31 Jun) before measurements on 5 Jul.

<sup>&</sup>lt;sup>d</sup>Water use of Thompson Seedless vines measured with the weighing lysimeter on 24 Aug 2005 was 40 L (5.3 mm) and vines were irrigated with 8 L five times that day. The no-applied water treatment had not been irrigated at any time during 2005.



**Figure 4** The relationship between  $\Psi_{\scriptscriptstyle \parallel}$  measured on deficit irrigated or nonirrigated grapevines and VPD at the time of measurement, using the equation: y = -1.24 - 0.0179 \* VPD,  $R^2$  = 0.31\*\*. Measurements were between 1000 and 1600 hr. The regression from Figure 1 and  $\Psi_{\scriptscriptstyle \parallel}$  values of the irrigated treatments for each cultivar measured at the same time are given for comparison. (T.S.: Thompson Seedless; C.S.: Cabernet Sauvignon; \*\* indicates significance at p < 0.01.) Other information is as given in Figure 1.

any changes in  $\Psi_1$  would be in response to other environmental conditions and that PFD was nonlimiting. Differences in temperature and VPD were obtained by taking measurements beginning early in the growing season through to near harvest and on some occasions from midmorning to midafternoon across a range of locations and grapevine cultivars.

It was assumed that the vines used in this study were well-watered, that is, that water was nonlimiting in the soil profile. The Thompson Seedless vines used in 2005 were grown in a weighing lysimeter and irrigated whenever they used 2 mm of water, as reported previously (Williams et al. 2003b). Vines irrigated at less than ET<sub>c</sub>, with ET<sub>c</sub> determined with a weighing lysimeter, had lower midday  $\Psi_1$ ,  $\Psi_{\text{stem}}$ ,  $g_{\text{s}}$ , and leaf net  $CO_2$  assimilation compared with those irrigated at full ET<sub>c</sub> or greater (Williams and Trout 2005). Vines grown at the other locations used in our study were irrigated based upon the calculation of ET using ET<sub>a</sub> and estimated seasonal crop coefficients (Williams and Ayars 2005b) and then using only treatments in which applied water was greater than estimated ET<sub>c</sub>. Values of g<sub>s</sub> and E measured on the Merlot and Cabernet Sauvignon vines were similar to those measured on the Thompson Seedless vines (Table 3), indicating that those vines were being irrigated at or close to full ET<sub>c</sub>.

The maximum values of  $\Psi_1$ ,  $g_s$ , and E for the fully irrigated vines in our study (Table 2) were generally greater than those found in other studies. For example, maximum  $g_s$  for field-grown vines in Spain having a "sufficient supply of water" was equivalent to 326 mmol m<sup>-2</sup> s<sup>-2</sup>, with most  $g_s$  values much less (Jacobs et al. 1996). Stomatal

conductance for vines receiving full irrigation in a field study in Portugal averaged 300 mmol m<sup>-2</sup> s<sup>-1</sup> (De Souza et al. 2003). In another study, vines "maintained in a wellwatered state" had maximum g values that approached 500 mmol m<sup>-2</sup> s<sup>-1</sup> at low ambient temperatures and VPD; afternoon (1600 hr) values of  $\Psi_1$  for the well-watered vines decreased to -1.4 MPa and g<sub>s</sub> decreased as temperature and VPD increased (Correia et al. 1995). Schultz (2003) reported a maximum  $g_s$  and a minimum  $\Psi_1$  of ~250 mmol m<sup>-2</sup> s<sup>-1</sup> and -1.4 MPa, respectively, for vines irrigated weekly with 30 L vine-1 in 1994 and 50 L vine-1 in 1995 from diurnal measurements taken in August. Maximum g of Carignane and Merlot was estimated as 560 and 440 mmol m<sup>-2</sup> s<sup>-1</sup>, respectively, by Winkel and Rambal (1990). In another study, typical values at midday for stomatal resistance and  $\Psi_1$  of a "wet" treatment were reported as ~1.0 s cm<sup>-1</sup> (equivalent to a stomatal conductance of 400 mmol m<sup>-2</sup> s<sup>-1</sup>) and -0.95 MPa, respectively (van Zyl 1987). In an irrigation study on Colombard grapevines, maximum g and E were 530 and 29 mmol m<sup>-2</sup> s<sup>-1</sup>, respectively (Stevens et al. 1995), while another study reported maximum values of 600 and >15 mmol m<sup>-2</sup> s<sup>-1</sup>, respectively (Cuevas et al. 2006). As seen from the above examples, the values of g<sub>s</sub>, E, and  $\Psi_1$  found in our study for the fully irrigated vines are equal to or greater than those that others have reported for well-watered vines.

Leaf Ψ of fully irrigated grapevines decreased as VPD and temperature increased, despite measurements being taken at various times during the growing season and different times of day (as long as light was nonlimiting) across locations and cultivars. Based on the regression equation, midday Ψ<sub>1</sub> of well-watered grapevines decreased from -0.65 to -0.89 MPa as VPD increased from 2 to 5 kPa. The slope of the equation used to describe the relationship here was not significantly different from that reported elsewhere (Williams and Trout 2005). McCutchan and Shackel (1992) also reported that prune Ψ<sub>1</sub> was linearly correlated with VPD ( $R^2 = 0.70$ ). The lowest value of Ψ, measured in this study for well-watered vines, -1.15 MPa for Cabernet Sauvignon at the Paso Robles site, occurred when VPD was 8.7 kPa at the time of measurement. This  $\Psi_1$  value is still greater than that reported by others at midday for well-watered or control vines (Correia et al. 1995, Schultz 1996).

Stem  $\Psi$  of grapevines was also linearly related to VPD in our study. Stevens et al. (1995) reported that  $\Psi_{\rm stem}$  of their well-watered vines was correlated with VPD ( $\Psi_{\rm stem}=-0.326$  -  $0.052*{\rm VPD}$ ,  $R^2=0.63$ ). McCutchan and Shackel (1992) found that  $\Psi_{\rm stem}$  of prune (*Prunus domestica* L. cv. French) was linearly related to VPD ( $\Psi_{\rm stem}=-0.41$  -  $0.12*{\rm VPD}$ ,  $R^2=0.81$ ) when some of the early and late season data were excluded from the calculation (all  $\Psi_{\rm l}$  and  $\Psi_{\rm stem}$  data collected in our study were used in the correlations with VPD). It was subsequently shown that this baseline also fit data collected on almond (*Prunus dulcis* (Mill.) Webb.) and was termed a fully irrigated (Shackel et al. 1998) or a nonstressed (Shackel et al. 1997) baseline.

The measurement of predawn leaf water potential  $(\Psi_{PD})$ is often considered a more reliable means of assessing grapevine water status than that of  $\Psi_{l}$  or  $\Psi_{stem}$  measured at midday (Gruber and Schultz, 2005). One reason for this preference is that  $\Psi_{\scriptscriptstyle 1}$  was shown to be highly correlated with daily maximum temperature—it increased or decreased at midday as ambient temperature increased or decreased, respectively, for vines irrigated with large amounts of water (H.R. Schultz and M.A. Matthews, unpublished data, cited in Gruber and Schultz 2005). It was concluded that  $\Psi_1$  measurements taken during the day may be uncoupled from the water status of the soil since  $\Psi_1$  varied with temperature changes. The observation of Schultz and Matthews was also demonstrated in our study, Ψ<sub>1</sub> was inversely related to ambient temperature at the time of measurement, and their data fit the regression we found for  $\Psi_1$  as a function of temperature (Figure 2). Therefore, the results from our study indicate that the variation of  $\Psi_1$ to changes in the environment does not mean  $\Psi_1$  is uncoupled from soil water status. It has also been found that midday  $\Psi_1$  and  $\Psi_{stem}$  were more highly correlated with soil water content and soil matric potential than was  $\Psi_{PD}$ . (Williams and Araujo 2002, Williams and Trout 2005).

Transpiration (E) of a plant can be directly related to the liquid phase soil to leaf water potential gradient and inversely related to the total resistance following the Ohm's law analog (van den Honert 1948):

$$E = (\Psi_{soil} - \Psi_{l}) / R_{sl}$$
 (1)

where  $\Psi_{soil}$  is soil water potential,  $\Psi_{l}$  is leaf water potential, and  $R_{sl}$  is the combined soil to leaf water flow resistance. Rearranging the equation:

$$\Psi_{1} = \Psi_{\text{soil}} - R_{\text{sl}} * E \tag{2}$$

it can be seen that any variation in E, brought about by changes in evaporative demand will affect  $\Psi_1$  even when soil water is not limiting. The decrease in  $\Psi_{l}$  or  $\Psi_{stem}$  because of the increase in VPD or temperature presented in this paper and elsewhere is a reflection of the increase in evaporative demand and an increase in E. Equations 1 and 2 also explain the results reported in Figure 4. As water is depleted in the soil profile, leaf transpiration decreases (because of reduced g<sub>s</sub> or increased R<sub>1</sub>) and the influence of VPD and temperature on  $\Psi_1$  is diminished and  $\Psi_{soil}$  becomes a more dominant factor (Table 2). Therefore, the slope of the relationship between  $\Psi_1$  and VDP is less for water-stressed vines than for well-watered grapevines. These differences in the response to VPD between fully irrigated and water-stressed vines are supported by a report that diurnal  $\Psi_1$  of irrigated grapevines correlated with VPD, but no such relationship was found in vines that had not been irrigated (Smart and Barrs 1973). An extreme example of soil moisture availability having a stronger effect on  $\Psi_1$  than does VPD has been reported; midday  $\Psi_1$ of nonirrigated Chardonnay grapevines in September was -1.81 MPa with a low VPD of 1.9 kPa at the time of measurement (Williams and Araujo 2002).

The relationship between  $\Psi_1$  and/or  $\Psi_{\text{stem}}$  and VPD would be useful as a baseline to determine whether grapevines are well watered. It is often assumed that the application of water, regardless of the amount, results in vines that are not stressed for water. An example is found in a study where  $\Psi_1$  was -1.5 MPa for both irrigated and waterstressed vines by midafternoon on 7 January, which had a maximum temperature of 34.9°C and a noon humidity of 36% (Smart 1974). Based on the regression in Figure 2, the  $\Psi_1$  of well-watered vines at that temperature would be expected to be no lower than -0.85 MPa. The lowest  $\Psi_1$ value obtained in our study at that temperature was greater than -1.0 MPa. The vines in the Smart (1974) study were furrow irrigated approximately every 10 days, but it was not reported when the last irrigation event occurred before January 7 or the amount of water applied each time. Even when vines are irrigated numerous times daily, values of net CO, assimilation,  $g_s$ , and  $\Psi_1$  can be significantly lower for deficit-irrigated vines (water applied at 0.6 or less of measured ET<sub>c</sub>) compared to those irrigated at ET<sub>c</sub> or greater (Williams and Trout 2005). Therefore, it appears that vines receiving irrigation are not necessarily well-watered with maintained high values of A,  $g_s$ , and  $\Psi_l$ , as noted elsewhere (Schultz 2003).

Plant species have been classified as having either isohydric or anisohydric stomatal behavior, which affects the differences in the diurnal timecourse of E and  $\Psi_1$  for plants that are fully irrigated compared with those that are waterstressed (Tardieu and Simonneau 1998). It has been assumed that grapevines exhibit isohydric stomatal behavior because  $\Psi_1$  of water-stressed grapevines does not drop significantly below that of watered vines during the day (Medrano et al. 2003). An alternative explanation would be that the  $\Psi_1$  of grapevines at midday is similar across a range of soil water availabilities (Cifre et al. 2005). However, Schultz (2003) concluded that the V. vinifera cultivar Grenache exhibited near-isohydric stomatal behavior while Syrah exhibited anisohydric behavior (i.e.,  $\Psi_1$  of waterstressed vines was significantly lower than that of the watered control throughout most of the day). Based on the data presented here, Thompson Seedless, Cabernet Sauvignon, and Merlot could be considered cultivars with anisohydric stomatal behavior. For all three cultivars,  $\Psi_1$  of fully irrigated vines was always significantly greater than that of the deficit irrigated or nonirrigated vines, regardless time of day (Table 3) or VPD at the time of measurement (Figure 4). Under reexamination, the findings of Medrano et al. (2003) indicate that when  $\Psi_1$  of waterstressed vines of Tempranillo and Manto Negro are compared with those of well-irrigated vines (irrigation at 100% of potential evapotranspiration) both cultivars exhibit anisohydric stomatal behavior.

The data presented in this paper has practical applications. The relationship between  $\Psi_1$  or  $\Psi_{\text{stem}}$  and VPD (or temperature) would be useful in determining whether the amount of water in the soil profile is limiting growth early in the season, as the baseline derived here indicates a

fully irrigated grapevine. In addition, the time span within which  $\Psi_{\rm l}$  or  $\Psi_{\rm stem}$  is measured could be expanded since changes in environmental conditions are taken into account (as long as PFD is nonlimiting). The senior author has found that hourly values of ambient temperature and relative humidity measured in a vineyard were similar to those obtained from CIMIS and PRWCA weather stations. Thus, environmental data are readily available to growers in California. Lastly, the lessened response of  $\Psi_{\rm l}$  of waterstressed vines to changes in VPD would indicate that once  $\Psi_{\rm l}$  is less than -1.2 MPa, soil moisture availability could be assumed to be the primary factor affecting vine water status.

A nonstress baseline has been used in the calculation of the Crop Water Stress Index (CWSI) (Idso et al. 1981, Jackson et al. 1988). The nonstress baseline compares the relationship between differences in the canopy temperature of plants minus that of ambient temperature at a specific VPD for plants transpiring at their full potential (fully irrigated). Such a baseline has been developed for grapevines growing in the San Joaquin Valley for use in calculating a CWSI (Grimes and Williams 1990). A fully irrigated baseline, similar to one previously proposed (Shackel et al. 2000), could also used on grapevines with the relationship between either  $\Psi_1$  or  $\Psi_{\text{stem}}$  and VPD found in this study. In Shackel et al. (2000), the water potential deficit equaled a fully irrigated  $\Psi_{\text{stem}}$  value (derived from the fully irrigated or nonstress baseline of  $\Psi_{\text{stem}}$  as a function of VPD, [McCutchan and Shackel 1992]) minus an observed or measured  $\Psi_{\text{stem}}$ . This water potential deficit was highly correlated with the degree of irrigation deficit across several years of study in numerous orchards. The usefulness of this type of calculation currently is being examined by the senior author under different irrigation regimes.

## **Conclusions**

Both  $\Psi_{\rm l}$  and  $\Psi_{\rm stem}$  of well-watered grapevines varied as a function of ambient temperature and VPD at the time measurements were made. Approximately 75% of the variation in  $\Psi_{\rm l}$  of well-watered grapevines—using four cultivars, grown at five locations, measured on different dates and times of day throughout the growing season, across years, and irrigated at different frequencies—was explained by VPD at the time of measurement. Based on the regression here, estimates of  $\Psi_{\rm l}$  at a VPD of 2 and 5 kPa for fully irrigated grapevines would be -0.65 and -0.89 MPa, respectively. Leaf  $\Psi$  of deficit or nonirrigated grapevines was less responsive to VPD than the fully irrigated vines when  $\Psi_{\rm l}$  was <-1.2 MPa. Such information would be useful in determining whether vines were stressed for water in any vineyard irrigation management program.

### Literature Cited

Cifre, J., J. Bota, J.M. Secalona, H. Medrano, and J. Flexas. 2005. Physiological tools for irrigation scheduling in grapevine (*Vitis vin*-

- *ifera* L.). An open gate to improve water-use efficiency. Agric. Ecosyst. Environ. 106:159-170.
- Correia, J.J., J.S. Pereira, M.M. Chaves, M.L. Rodriques, and C.A. Pacheco. 1995. ABA xylem concentrations determine maximum daily leaf conductance of field-grown *Vitis vinifera* L. plants. Plant Cell Environ. 18:511-521.
- Cuevas, E., P. Baeza, and J.R. Lissarrague. 2006. Variation in stomatal behaviour and gas exchange between mid-morning and midafternoon of north-south oriented grapevines (*Vitis vinifera* L. cv. Tempranillo) at different levels of soil water availability. Sci. Hortic. 108:173-180.
- De Souza, C.R., J.P. Maroco, T.P. dos Santos, M.L. Rodrigues, C.M. Lopes, J.S. Pereira, and M.M. Chaves. 2003. Partial rootzone drying: Regulation of stomatal aperature and carbon assimilation in field-grown grapevines (*Vitis vinifera* cv. Moscatel). Funct. Plant Biol. 30:653-662.
- Grimes, D.W., and L.E. Williams. 1990. Irrigation effects on plant water relations and productivity of 'Thompson Seedless' grapevines. Crop Sci. 30:255-260.
- Grimes, D.W., H. Yamada, and S.W. Hughes. 1987. Climate-normalized cotton leaf water potentials for irrigation scheduling. Agric. Water Man. 12:293-304.
- Gruber, B.R, and H.R. Schultz. 2005. Coupling of plant to soil water status at different vineyard sites. Acta Hortic. 689:381-389.
- Idso, S.B., R.D. Jackson, P.R. Pinter, R.J. Reginato, and J.L. Hatfield. 1981. Normalizing the stress-degree-day parameter for environmental variability. Agric. Meteorol. 24:45-55.
- Jackson, R.D., W.P. Kustas, and B.J. Choudbury. 1988. A reexamination of the crop water stress index. Irr. Sci. 9:309-317.
- Jacobs, C.M.J., B.J.J.M. van den Hurk, and H.A.R. de Bruin. 1996. Stomatal behaviours and photosynthetic rate of unstressed grapevines in semi-arid conditions. Agric. For. Meteor. 80:111-134.
- McCutchan, H., and K.A. Shackel. 1992. Stem-water potential as a sensitive indicator of water stress in prune trees (*Prunus domestica* L. cv. French). J. Am. Soc. Hortic. Sci. 117:607-611.
- Medrano, H., J.M. Escalona, J. Cifre, J. Bota, and J. Flexas. 2003. A ten-year study on the physiology of two Spanish grapevine cultivars under field conditions: Effects of water availability from leaf photosynthesis to grape yield and quality. Funct. Plant Biol. 30:607-619.
- Schultz, H.R. 1996. Water relations and photosynthetic responses of two grapevine cultivar of different geographical origin during water stress. Acta Hortic. 427:251-266.
- 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 Environ. 26:1393-1405.
- Shackel, K.A., et al. 1997. Plant water status as an index of irrigation need in deciduous fruit trees. HortTechnology 7:23-29.
- Shackel, K., S. Gurusinghe, D. Kester, and W. Micke. 1998. Water stress responses of almond [*Prunus dulcis* (Mill.) Webb.] trees under field conditions. Acta Hortic. 470:309-314.
- Shackel, K.A., B. Lampinen, S. Southwick, W. Olson, S. Sibbett, W. Kreger, J. Yeager, and D. Boldhamer. 2000. Deficit irrigation in prunes: Maintaining productivity with less water. HortScience 35:1063-1066.
- Smart, R.E. 1974. Aspects of water relations of the grapevine (*Vitis vinifera*). Am. J. Enol. Vitic. 25:84-91.

- Smart, R.E., and H.D. Barrs. 1973. The effect of environment and irrigation interval on leaf water potential of four horticultural species. Agric. Meteor. 12:337-346.
- Stevens, R.M., G. Harvey, and D. Aspinall. 1995. Grapevine growth of shoots and fruit linearly correlated with water stress indices based on root-weighted soil matric potential. Aust. J. Grape Wine Res. 1:58-66.
- Synder, R.L., and W.O. Pruitt. 1992. Evapotranspiration data management in California. *In* Proceedings of the ASCE National Conference, Water Forum 1992, pp. 128-133. Am. Society of Civil Engineers, Baltimore.
- Tardieu, F., and T. Simonneau. 1998. Variability among species of stomatal control under fluctuating soil water status and evaporative demand: Modelling isohydric and anisohydric behaviours. J. Exp. Bot. 49:419-432.
- Turner, N.C. 1991. Measurement and influence of environmental and plant factors on stomata conductance in the field. Agric. For. Meteor. 54:137-154.
- van den Honert, T.H. 1948. Water transport in plants as a catenary process. Disc. Faraday Soc. 3:146-153.
- van Zyl, J.L. 1987. Diurnal variation in grapevine water stress as a function of changing soil water status and meteorological conditions. S. Afr. J. Enol.Vitic. 8:45-52.
- Williams, L.E., and F. Araujo. 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* L. J. Am. Soc. Hortic. Sci. 127:448-454.

- Williams, L.E., and J.E. Ayars. 2005a. Water use of Thompson Seedless grapevines as affected by the application of gibberellic acid (GA<sub>3</sub>) and trunk girdling: Practices to increase berry size. Agric. For. Meteor. 129:85-94.
- Williams, L.E., and J.E. Ayars. 2005b. Grapevine water use and the crop coefficient are linear functions of the shaded area measured beneath the canopy. Agric. For. Meteor. 132:201-211.
- Williams, L.E., and T.J. Trout. 2005. Relationships among vine- and soil-based measures of water status in a Thompson Seedless vine-yard in response to high-frequency drip irrigation. Am. J. Enol. Vitic. 56:357-366.
- Williams, L.E., N.K. Dokoozlian, and R.L. Wample. 1994. Grape. *In* Handbook of Environmental Physiology of Fruit Crops. Vol. 1. Temperate Crops. B. Shaffer and P.C. Anderson (Eds.), pp. 83-133. CRC Press, Orlando, Florida.
- Williams, L.E., C.J. Phene, D.W. Grimes, and T.J. Trout. 2003a. Water use of young Thompson Seedless grapevines in California. Irrig. Sci. 22:1-9.
- Williams, L.E., C.J. Phene, D.W. Grimes, and T.J. Trout. 2003b. Water use of mature Thompson Seedless grapevines in California. Irrig. Sci. 22:11-18.
- Winkel, T., and S. Rambal. 1990. Stomatal conductance of some grapevines growing in the field under a Mediterranean environment. Agric. For. Meteor. 51:107-121.