

Cold Hardiness of Dormant Buds of Grape Cultivars: Comparison of Thermal Analysis and Field Survival

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Abstract. Thermal analysis (TA) was used to evaluate dormant bud cold hardiness of nine *Vitis* cultivars weekly during the 1993-94 dormant period. TA hardiness estimates were expressed as either mean low-temperature exotherm temperature (MLTE) or temperatures lethal to 10% (LT₁₀), 50% (LT₅₀), or 90% (LT₉₀) of dormant bud sample. A destructive freeze on 19 Jan. 1994 presented an opportunity to compare dormant bud field survival with laboratory estimates of bud hardiness that had been derived from TA. Vineyard air temperatures of -24C caused primary bud kill that ranged from a mean of 15% with 'Concord' to 100% with 'Viognier'. With the exception of 'Viognier' and one of two 'Cabernet Sauvignon' clones, field mortality levels were accurately bracketed by TA estimates of LT₁₀, MLTE, and LT₉₀ values, which had been obtained in the week preceding the freeze. 'Viognier' bud hardiness was overestimated by ≈1.5C, and the hardiness of 'Cabernet Sauvignon clone UCD#6' was underestimated by <1C. The discrepancy with 'Viognier' may have been related to prior destruction of primary buds by bud necrosis and the misinterpretation of secondary bud exotherms as due to primary buds.

Thermal analysis (TA), including differential thermal analysis (DTA), is commonly used to measure cold hardiness of tissues and organs that avoid freezing by supercooling, such as floral buds of cherry (*Prunus cerasus* L.) (Callan, 1990) and peach [*Prunus persica* (L.) Batsch] (Durner and Gianfagna, 1991; Quamme, 1978) and the mixed buds of grape (Andrews et al., 1984; Quamme, 1986; Wolf and Pool, 1987). In practice, TA uses sensitive thermocouples or thermoelectric (TE) modules to detect the latent heat of fusion released when supercooled tissues freeze. Grape bud TA was originally explored by Pierquet et al. (1977) and Pierquet and Stushnoff (1980), who established a relationship between bud exothermic events and specific primordia destruction. Grape bud TA was modified by Andrews et al. (1984) to take advantage of the greater sensitivity and sample size afforded by TE modules. Refinements subsequently were made to freezing protocol (Quamme, 1986; Wolf and Pool, 1987). The advent of low-cost data-acquisition hardware (Wample et al., 1990; Wolf and Pool, 1986) has facilitated the

application of grape bud TA to studies of genotypic differences (Pool et al., 1990) and response to the environment (Pool et al., 1992; Wolf and Cook, 1992) and cultural practices (Hamman et al., 1990; Wample et al., 1993).

The conviction that laboratory hardiness tests faithfully represent actual field hardiness is central to studies that seek to predict plant survival or damage in response to field conditions. Using TA as a measure of grape bud hardiness was examined in early TA research and found valid (Andrews et al., 1984; Quamme, 1986; Wolf and Pool, 1987). Typically, comparisons were made of TA and other laboratory techniques, such as browning of specific tissues following sample exposure to a range of test temperatures. Differences in hardiness estimates measured by these techniques were ≈1 to 3C. The ability of TA and DTA to accurately estimate bud cold hardiness can be affected by bud excision, the cooling rate, and other protocol; the technique also may be inappropriate for some systems (Flinn and Ashworth, 1994). Cooling rates of ≤10C/h did not affect the low-temperature exotherm (LTE) temperature of grape bud TA (Quamme, 1986; Wolf and Pool, 1987), but LTE temperatures can be increased if buds are excised without attached nodal tissue (Quamme, 1986; Wolf and Pool, 1987). Based on work with peaches (Ashworth and Davis, 1987), the attached nodal tissue may serve as an anatomical barrier to ice nucleation of the bud primordia.

Despite more than 10 years of use, there have been few opportunities to closely compare TA hardiness estimates with actual field survival following cold stress episodes. This report resulted from the occurrence of a dam-

aging low-temperature episode on 19 Jan. 1994 that coincided with weekly TA estimates of grape bud cold hardiness. The data provided assurance that TA can accurately represent field hardiness, but some caution must be used interpreting exothermic events from samples of vines that have a significant proportion of previously destroyed primary buds.

Materials and Methods

Plant material. In this study, vines were grown at Virginia Tech's Winchester Agricultural Expt. Station, Winchester. Dormant bud cold hardiness of nine cultivars was evaluated weekly using TA from 29 Sept. 1993. *Vitis vinifera* L. cultivars were 'Cabernet Sauvignon clone UCD #6', 'Cabernet Sauvignon clone UCD #8', 'Fer Servadau', 'Petit Verdot', 'Riesling', and 'Viognier'. All vinifera cultivars were grafted to rootstock C-3309. Other cultivars were 'Concord' (*V. labrusca* L.), 'Seyval' (*Vitis* hybrid), and 'Norton' (*V. aestivalis* Michx.). All cultivars, except 'Concord', 'Seyval', and 'Riesling', were part of an evaluation study established in 1989 of 22 cultivars. Each cultivar was planted in three-vine plots, replicated five times, in a completely randomized design. 'Concord', 'Seyval', and 'Riesling' were planted in nonreplicated, independent rows in 1990 ('Seyval') and 1991 ('Concord' and 'Riesling'). All vines were spaced 2.1 m apart in north-south-oriented rows that were 3.7 m wide. Except for 'Concord' and 'Riesling', all vines were trained to bilateral cordons (1.1 m above ground), spur-pruned, and shoots positioned vertically. 'Concord' and 'Riesling' were trained to Geneva Double Curtain and shoots positioned downwards. Shoot density ranged from 15 to 20 per meter of canopy, except with 'Seyval', which had a density of 12 to 15 shoots/meter. Two recording thermographs, with a verification thermometer, were located in the vineyard, each 1.5 m above ground in National Weather Service-approved instrument shelters. Thermograph calibration was checked on 19 Jan. 1994 at -23C and -26C in a freezer (model T20-S with Watlow 942 programmer; Tenney Environmental, Philadelphia) that held a constant temperature with <1C variation.

Dormant bud TA. Weekly TA measures of dormant bud cold hardiness were made starting 29 Sept. 1993 and continuing through late Jan. 1994. For each cultivar, at least two representative canes were collected from different vines at each test date. Buds at node positions three to 15 were used for TA; they were excised in the laboratory and mounted on TE modules (MELCOR, Trenton, N.J.) with a strip of water-moistened filter paper to promote ice nucleation of moisture at the bud-TE module interface (Wolf and Pool, 1987). Buds were excised so that 0.5 to 1.0 mm of subtending nodal tissue remained attached to the bud. For each hardiness test, four or five buds were mounted on each of four or five TE modules, for a total of 20 buds per cultivar. Loaded modules were cooled from 0 to -35C, at 4C/h in the previously described freezer. TE volt-

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age and separate thermocouple temperature data were recorded every 20 sec with the data-acquisition hardware (Wolf and Pool, 1986). Data were subsequently plotted to determine the LTE temperature. The median primary bud LTE temperatures for each of the four or five TE modules were averaged to derive a single mean LTE (MLTE) value for each cultivar. In addition to MLTE temperatures, lethal temperature values were obtained for primary buds by ranking all primary bud LTEs for a given cultivar and determining the temperature lethal to 10%, 50%, and 90% (LT₁₀, LT₅₀, and LT₉₀, respectively) of the sample for each test date. The yield of LTEs judged to originate from primary buds was 0.9 to 1.0. LTE "yield" represents the ratio of exotherms to the number of buds examined. Each compound grape bud should produce one primary bud exotherm; however, the actual exotherms : number of buds ratio in a sample usually averages less than one. Reductions in LTE

yield can result from prior bud destruction, failure to register exotherms due to extended signal recording frequency, simultaneous bud freezing that superimposes two or more exotherms, and perhaps other reasons.

Assessment of bud kill from field exposure. Primary bud mortality was evaluated following the 19 Jan. freeze by collecting one representative cane per vine on 21 Jan. (≈15 canes per cultivar). The canes were held at 22C for 48 to 72 h, after which buds at nodes three through 15 were sectioned with a razor and examined visually. The primary bud was judged dead if brown and alive if green. Previous to this exercise, a similar survey had been conducted 18 to 21 Nov. 1993 to determine the primary bud necrosis level (Morrison and Iodi, 1990) among all cultivars. The bud necrosis survey used three canes per vine, whereby buds at nodes three through 15 were sectioned with a razor and evaluated in a manner similar to that described for assessing freeze injury.

Results and Discussion

Vineyard daily high and low temperatures from 1 Sept. 1993 through 1 Feb. 1994 are shown in Fig. 1. The first killing frost occurred 11 Nov., and temperatures throughout November and December were considered conducive to good cold hardiness acquisition. MLTE temperatures for the two 'Cabernet Sauvignon' clones decreased linearly through mid-December and attained levels by 1 Jan. that were lower than any previously recorded for this cultivar in Virginia (Wolf and Cook, 1992); acclimation conditions were excellent, at least for 'Cabernet Sauvignon' (Fig. 1). MLTE patterns for the other cultivars were similar in slope and stability to patterns shown for 'Cabernet Sauvignon'.

The period from 15 to 20 Jan. 1994 was characterized by unusually low temperatures. Both vineyard thermographs recorded a low of -24C from 0600 to 0800 HR on the morning of 19 Jan. Cooling rate in the vineyard was 0.7C/h from 1800 to 2400 HR on 18 Jan. and 0.6C/h from 0000 to 0800 HR on 19 Jan. Records from the National Weather Service station at Dulles Airport, Chantilly, Va. (75 km away) indicated that sustained winds from 0000 to 0600 HR on 19 Jan. were no less than 3.6 m·s⁻¹ and averaged 4.5 m·s⁻¹. Given the wind, the vineyard's small size (≈1.0 ha), and the uniform temperatures recorded in the vineyard, it is reasonable to presume that all plots experienced essentially the same temperature.

For seven of the nine cultivars evaluated during Winter 1993-94, TA provided a close approximation of actual bud hardiness as evaluated by bud kill following 19 Jan. (Table 1). MLTE temperatures and LT₅₀ values (not shown) differed by no more than 0.7C. Either estimate of hardiness could be used; however, MLTE temperature has more general use in TA studies and was the estimate of choice in Table 1. Thermal analysis overestimated cold hardiness of 'Viognier' buds by ≥1.5C. The Nov. 1993 survey indicated that 'Viognier' suffered 32% primary bud necrosis (Table 2). Causes of grape bud necrosis are unknown, but the condition observed in Virginia appeared similar to that described for table grapes in California (Morrison and Iodi, 1990) and Chile (Pérez and Kliewer, 1990). Bud necrosis also was observed among other cultivars but at

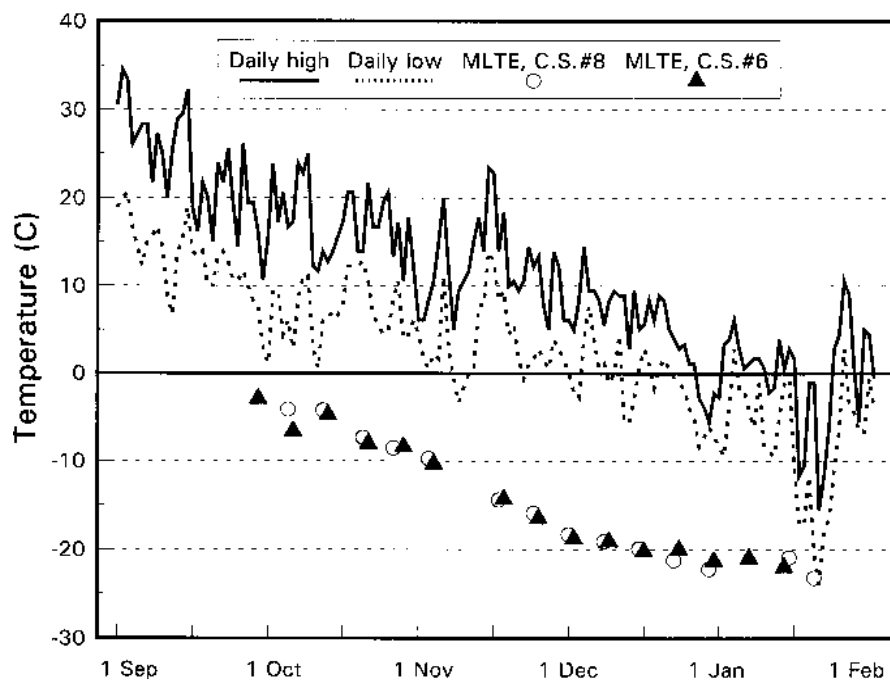


Fig. 1. Daily high and low air temperature at vineyard and mean low-temperature exotherm (MLTE) temperatures for 'Cabernet Sauvignon clone UCD#8' and 'Cabernet Sauvignon clone UCD#6' from 1 Sept. 1993 through 1 Feb. 1994.

Table 1. Comparison of dormant bud cold hardiness measured by thermal analysis and field mortality sustained by nine *Vitis* cultivars before and as a result of exposure to -24C on 19 Jan. 1994.

Hardiness test date	Cabernet Sauvignon #8 ^a			Concord			Riesling		
	LT ₁₀	MLTE	LT ₉₀	LT ₁₀	MLTE	LT ₉₀	LT ₁₀	MLTE	LT ₉₀
13 Jan.	-18.3	-21.1	-22.8	-20.0	-25.6	-27.2	-21.1	-23.3	-25.0
18 Jan.	-20.5	-23.3	-24.4	-20.5	-26.1	-30.6	-20.0	-25.0	-26.1
Dead primary buds (%) ^y		90			15			37	
		Norton				Viognier			
10 Jan.	-25.6	-28.3	-29.4	-21.1	-23.9	-25.0	-23.3	-25.0	-27.8
17 Jan.	-23.3	-28.9	-30.6	-22.9	-24.2	-25.5	-17.8	-25.6	-28.3
Dead primary buds (%) ^y		22			100			32	
		Cabernet Sauvignon #6				Petit Verdot			
5 Jan.	-19.4	-21.1	-22.2	-18.3	-22.2	-23.9	-18.9	-20.5	-22.8
12 Jan.	-20.5	-22.2	-23.3	-17.8	-22.2	-23.9	-18.3	-21.1	-23.9
Dead primary buds (%) ^y		76			95			99	

^aLT₁₀ = temperature lethal to 10% of a bud sample; MLTE = mean low-temperature exotherm temperature; LT₉₀ = temperature lethal to 90% of a bud sample.

^yDetermined on 21 Jan.

Table 2. Cultivar differences in primary bud destruction resulting from bud necrosis (assessed Nov. 1993) and field exposure to -24C on 19 Jan. 1994. Canes from nonreplicated cultivars were randomly collected from at least five vines. Percentage data were square-root-transformed before analysis of variance but are shown as nontransformed values.

Cultivar	Bud necrosis		Cold-injured	
	n ^z	% ^y	n ^z	% ^y
Viognier	18	31.5 A	10	100 A
Fer Servadau	22	5.3 B	13	99 A
Petit Verdot	12	0.0 B	8	95 AB
Cabernet Sauvignon Clone #8	30	5.7 B	15	90 AB
Cabernet Sauvignon Clone #6	30	3.2 B	15	76 B
Riesling	10	0.6 ^x	10	37 ^x
Seyval	20	1.3 ^x	12	32 ^x
Norton	12	4.8 B	13	22 C
Concord	---	--- ^x	10	15 ^x

^zn = number of canes examined.

^yMean separation within rows by Duncan's multiple range test at $P \leq 0.05$; nonreplicated vines were not included in mean separation.

^xNot replicated.

<6%. Interpretation of grape bud LTEs is complicated by the presence of secondary buds, usually two, in the compound structure (Pratt, 1974). Primary bud exotherms are judged on the basis of their greater size and higher temperature of occurrence than those of secondary buds; however, some subjectivity is inherent with interpretation. We suspect that the hardness discrepancy with 'Viognier' was produced by including some secondary bud LTEs in the data interpretation, even though 'Viognier' LTE yield was expected to be lower (≈ 0.7) due to 32% primary bud necrosis. Although the cold hardness of 'Cabernet Sauvignon clone UCD #6' was slightly underestimated, the discrepancy was reasonable, particularly because it had been >6 days since the last MLTE determination. No attempt was made to estimate the cold hardness of secondary buds or to methodically tally secondary bud mortality in the field, although this exer-

cise would have been possible. Due to the significant cold injury on 19 Jan. (Table 2), further hardness tests during Winter 1993-94 were suspended.

We demonstrated that TA can accurately estimate dormant bud cold hardness of the *Vitis* species and an interspecific hybrid that represent a wide range of inherent cold hardness. In addition to relative differences in cold hardness between population samples, TA can be confidently used to estimate the absolute cold hardness of buds. Some caution must be used in estimating the cold hardness of a population that has a significant incidence of nonfreeze injury, and some judgment is required to distinguish the LTEs of primary and secondary buds if this is of interest to the researcher.

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