

Heat tolerance of winegrapes: Experimental design

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1 Methods so far...

Cuttings taken from RMI vineyard at UC-Davis in December 2015. Growing started in January 2016. They are currently planted in 26 cm diameter pots.

Grapevines were dried down and placed into chambers with moist soils on 27 May 2016. Chambers are set (program called `VITIS.DORM`) to 6°C and 400 ppm light for 8 hours (day) then 4°C and no light for the other 16 hours. Humidity is set at 80%.

They were then watered again on 13-14 June 2016 (still greenish then) but Kea (greenhouse manager) said they appeared dormant about a week late (should check with her on this).

Plants were removed on 15 August 2016 and put into cool greenhouses (15.5-18°C during the night, 18-21°C during the day). They were pruned down to 2-bud spurs and measurements of the buds were taken. Plants were kept moist and monitored 2X/week for possible growth. Greenhouse temperatures after the first week were slowly ramped up to normal operating temperatures (I need to check but I think 10°C at night 23-28°C during the day).

2 What we measure in the greenhouse

For baseline data we measured the spur size—the size of the two buds and the distance between them also.

We monitor the plants every 2-3 days recording Eichorn Lorenz stage (for each bud) and soil moisture (for each plant).

3 Training and managing growth

We'll aim to manage plants for relatively equal growth and for one strong cluster per stem following the below:

- Make sure all the plants are pruned back to 2 bud spurs. (Done.)
- Train the whole shoot, and plan to train the two shoots (with two stakes). Sucker all the other shoots.
- Aim for one cluster per shoot; be careful removing the other clusters ... wait until they are separated from the leaves completely then *carefully* pinch them out.
- We'll aim to train to some number of leaves perhaps (around 10 or so). Try to keep the number similar across varieties. Remove the leaf and remove the apex (pinch it off once

the plant hits desired leaf number) BUT, we'll also keep an eye on shoot length for really short plants we should allow more leaves probably.

Remember the goal is to manage growth so that the plants—across varieties and across replicates—are as similar as possible. So we may need to adjust the above guidelines to make sure we reach this goal.

4 Sulphur & water

Plant receive nightly puffs of sulphur in the greenhouse. So far we have not seen mildew so we'll plan to monitor plants closely in the chamber but will only treat if we see mildew. If we see mildew we'll treat with a sulphur spray.

Plants will be kept moist, we'll record soil moisture.



5 Proposed design

We've designed the experiment to test how high temperatures alter both the *timing* and *duration* of flowering. Ideally with photos we'll also try see how the total flowering is affected (and thus possibly fruitset, as we follow the plants in the greenhouse when they come out of the chambers).

Plants will go into chambers when they hit EL stage 12 (5 leaves stage). We'll randomly assign which order they go in (e.g., if an individual of one variety hits EL stage 12 then we'll use a randomizer to decide if it goes into treatment 1, 2, 3, 4, or 5 until all the plants for that variety are in a treatment).



Plant will stay in chambers until the end of flowering. That is, they will each be removed once they reach 100% flower development (I realize they may not all flower but I think they will either flower or abort so 100% would be 100% flowering + flower aborting).



We'll rotate the treatment of each chamber once per week to remove (as possible) chamber effects.

Chamber setup:

Photoperiod: Based on a literature review of 13 studies, of which 5 were using ambient conditions (i.e., no control): 5 used 12 hours, 2 used 15 hours and 1 used 18 hours. So we'll do 12 hour days. It makes most sense for me given that we're replicating the earlier part of the season and it makes the math easier too.



Based on the literature review we found studies where temperatures ranged from 10-50°C (I know!) but for controlled studies the common range was 20-40°C with many experiments exposing the plants to these temperatures for 7-18 days (often at each stage). So I think we are covering the general range. Also—considering other crops, Porter & Gawith (1999, *European Journal of Agronomy*) found that 31°C was the max for wheat and Parent & Tardieu (2012, *New Phytologist*, this is a super cool paper by the way) found a range of optima temperatures from 21.6°C-32.7°C depending on the species.

Based on the same literature review we found the difference in daily amplitude of temperature (i.e., night versus day temperatures) is usually about 10°C. It varies though from a high of 20-



35°C in some studies to 5 or 6 in others. These lower amplitudes generally come from later in the season (for those studies using ambient conditions) or when people have higher temperature treatments. Since our chambers max out at 40°C for us to do a 10°C amplitude means the highest temperature we'll get is 35°C. Thus I think a 6°C amplitude would be better. My suggested temperatures are below (note I previously considered 4°C but now I think that seems too low):

- Mean of 20°C – 17/23°C
- Mean of 26°C – 23/29°C
- Mean of 30°C – 27/33°C
- Mean of 34°C – 31/37°C
- Mean of 37°C – 34/40°C






What we'll measure in the chambers

Here's the hoped for plan to measure every 2-3 days ...

- Shoot length
- Leaf number
- % flowering via photos perhaps and also possibly we'll count caps via bags
- Soil moisture for each pot

Possibly measure photosynthesis once a week? That's a big maybe.

6 Questions

1. Comments on the temperature ranges. 
2. Decide on which vars and how many exactly once we know what grows! 
3. What level to set the chambers' CO_2 ? 300ppm or 400ppm? 
4. Keep photoperiod in greenhouse at 12 hours once we remove grapes? 