Additional considerations

Introduction

Conditions in controlled environment plant growth rooms and chambers (CE units) should be reported in detail for comparison of results and duplication of experiments. The minimum guidelines table, along with these notes, should help meet these aims, indicating a required, minimum amount of information that should be reported. They may also highlight parameters that could be important, but that may not have been considered for measurement.

Average measurements should be reported, including their temporal standard deviation (s.d.).

All sensors should be calibrated regularly according to manufacturer's procedures and suggested frequency.

Radiation

- Output of all electric radiation sources decreases with hours of operation e.g. for some fluorescent lamps output may drop 20% after the first five months of use.
- Irradiance varies significantly across the growing area in many CE units.
- Vertical radiation gradients occur in all CE units, depending on chamber size, lamp type, lamp distribution, and luminaire shape.
- Spectra from electric lamps generally differ from that of the sun. Unnatural red to far-red light ratios may affect morphogenesis in some plants and photomorphogenic effects should be considered when interpreting results.

Temperature

- Differences may exist between the temperatures of the air and plant, especially under high radiation loads.
- Older on-off control systems can result in as much as ±5°C variation from the set point temperature.
- A vertical temperature gradient occurs in most CE units, depending on airflow rates and other factors.

Atmospheric moisture

- Air humidity affects plants in CE units directly (via transpiration and gas exchange) and indirectly (via the plant's energy balance and physical and biological environment).
- Heating and cooling cycles lasting only 1 to 3 minutes can change absolute humidity by 1 to 2%, altering relative humidity by 20 to 40%.
- Air humidity is a challenging parameter to monitor, but is critical to plant water relations and infection by foliar pathogens. Relative humidity (RH) is acceptable for

reporting humidity until CE units can control vapour pressure deficit (VPD), or portable instruments are available to measure and display VPD.

Carbon dioxide

- Carbon dioxide (CO_2) is probably the least controlled environmental parameter in CE studies. Unfortunately, too little or too much CO_2 is hard to detect until plants start to show specific symptoms.
- Small variations in CO₂ can affect plant growth and development significantly. People in or around CE units, and even greenhouses, can increase CO₂, as may motor vehicles, heating systems, and other nearby sources that produce CO₂.
- Few CE units manufactured today have CO₂ control or monitoring equipment installed as a standard feature. However, most do have some degree of ventilation or air exchange, and good air exchange can moderate CO₂ build-up or depletion.
- Even if a CE unit is well ventilated, it is important to remember that the surrounding area with which it exchanges air should also be well ventilated.

Experimental design issues

- Ideally, a single CE unit should be treated as a single replicate. True replication requires using multiple CE units, or repeating treatments in each unit with time, both expensive and time consuming options.
- Regular transfer of plants between CE units may be an alternative to avoid direct confounding of effects of an imposed environment with that of a CE unit.
- Repeating experiments in a CE unit with poorly controlled or monitored environmental parameters may lead to erroneous assumptions about treatment conditions and resulting data.

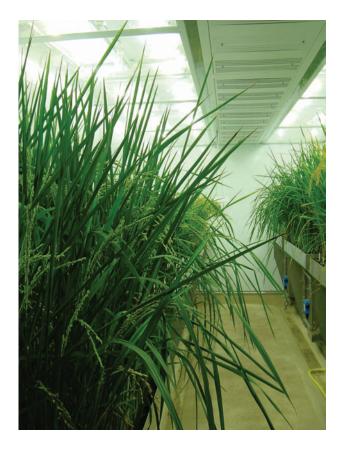
Example of a report suitable for publication

The experiment was conducted in a 3 m by 4 m growth room equipped with cool white fluorescent lamps (Model 830, Philips) mounted above a clear glass barrier, and an upward airflow distribution system using sufficient outdoor make-up air to provide ambient CO_2 conditions inside the room. The room air temperature was maintained at 25/20°C (s.d. ±2/1°C) during the light/dark period. The photosynthetically active radiation (PAR) at the top of the canopy was maintained at 400 µmol m⁻² s⁻¹ (s.d. ±10 µmol m⁻² s⁻¹) during the 12-hour photoperiod. The relative humidity in the room was maintained at 70% (s.d. ±10%). The plants were grown in 1 L pots filled with a peat-vermiculite (2:1 volume ratio) mixture. The plants were hand watered daily with a freshly prepared nutrient solution (full strength Hoagland, pH 6).

International Committee for Controlled Environment Guidelines

Minimum Guidelines for Measuring and Reporting Environmental Parameters for Experiments on Plants in Growth Rooms and Chambers

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Parameter to measure		Units ¹	Where to measure	When to measure	What to report
<i>Radiation</i> and	Photosynthetically active radiation (PAR) ²	µmol m ⁻² s ⁻¹	Top of plant canopy in centre of growing area	At start and end, and every 2 weeks of the experiment	Average and standard deviation. Radiation source (type, model, and manufacturer)
	Photoperiod	h			Duration of light and dark periods
Temperature	Air	°C	Top of plant canopy in centre of growing area	Daily during each light and dark period, at least 1 hour after light/dark change	Average and standard deviation
	Liquid culture	°C	Within solution under plants	As above for air temperature	Average and standard deviation
Atmospheric moisture or	Water vapour pressure deficit (VPD)	kPa	Top of plant canopy in centre of growing area	Daily during each light and dark period, at least 1 hour after light/dark change	Average and standard deviation
	Relative humidity (RH)	%	As above for VPD	As above fpr VPD	Average and standard deviation
<i>Carbon dioxide</i> ³		µmol mol-1	Top of plant canopy	At least hourly	Average and standard deviation
<i>Air velocity</i> ³		m s ⁻¹	At one or more representative canopy locations	At least once during the experiment	Average and standard deviation
Watering		litre (L)		Daily	Frequency, amount and type of water added
pН	Liquid culture	рН	In the bulk solution	Before and after pH correction	Average and standard deviation
Electrical conductivity (EC) ³	Liquid culture	S m ⁻¹	In the bulk solution	Before and after EC correction	Average and standard deviation
Substrate				At start of the experiment	Type and volume per container, components of soil-less substrate, container dimensions
Nutrition	Solid media	mol kg ⁻¹ (dry)		When added or replenished	Nutrients and their form added to soil media
	Liquid culture	mmol L ⁻¹		Daily, or when replenished	Ionic concentration in initial and added solution. Aeration if any. Volume of initial solution
Room or chamber properties	Specifications				Floor area. Manufacturer and model if available
	Barrier beneath lamps				Indicate if present and its composition
	Air flow				Indicate whether up, down or horizontal

1 Report in other multiples or sub-multiples of indicated units if more convenient.

² Referred to as photosynthetically active radiation (PAR: 400-700 nm) for general usage and described as photosynthetic photon flux density (PPFD) by many journals, professional societies, and manufacturers of quantum sensors. When diurnal PAR is ramped, integrals should be reported, e.g. in mol m² d⁻¹.

³ This parameter should be reported if records are available and always when it is a variable under investigation. For more information, consult the detailed guidelines published as ANSI/ASAE Engineering Practice EP411.4 (2002) 'Guidelines for measuring and reporting environmental parameters for plant experiments in growth chambers'. ASAE, 2950 Niles Road, St. Joseph, MI 49085-9659, USA.