



## Mar 18,

# Measuring PPFD on Algal Shaker 🖨

## Jakub Nedbal<sup>1</sup>

<sup>1</sup>King's College London

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#### ABSTRACT

The illuminated orbital shaker features a glowing growing area. The illumination is not entirely homogeneous. It is brightest near the centers of the LEDs and becomes dimmer closer to the edges. The light brightness of the light in the glowing area is dependent on the LEDs, their spacing, the distance between the LEDs and the transparent platform, the material of the anti-slip mat, among other parameters. To ensure consistency and reproducibility between experiments, the brightness can be measured and the light output of the illuminator calibrated. Quantum detector is used to measure the photosynthetic photon flux density (PPFD). The measurement described in this protocol is conducted for all current settings on the LED controller and in 12 different locations across the glowing area. It is done both in air and inside an Erlenmeyer flask containing water, to best capture the actual conditions experienced by the microalgae cultures.

#### **EXTERNAL LINK**

https://app.labstep.com/sharelink/fcc4e7da-8aff-4a1f-b611-3005066f8851

#### **GUIDELINES**

The goal is to calibrate the irradiance on the surface of the microalgae shaker in air and also inside a glass flask with deionized water in it against the setting of the LEDs.

For each LED current setting, the measurement must be taken in many locations due to the inhomogeneity of the illumination pattern. The resulting value is then the average of all these measurements for each current setting.

Acknowledgement: The measurements are done using professional equipment borrowed from Giorgio Perin (Patrick Jones group) at Imperial College London.

MATERIALS TEXT





Borosilicate Glass Narrow Neck
Erlenmeyer Flasks (100 ml)
by Fisher Scientific
Catalog #: 15409103

Water refers to sterilized deionized water

#### SAFETY WARNINGS

The LEDs are bright at the higher current settings. The brightness of the light right at the orbital shaker platform is comparable to the Sun on a bright day. Do not stare into the light. Keep a distance between your eyes and the glowing LEDs. Use sunglasses or other protective equipment if you find your eyes becoming sensitive due to the bright light.

### BEFORE STARTING

This protocol has been designed for a particular type of quantum light sensor (US-SQS/L, Walz) and light detector (LI-250A, Li-Cor Biosciences). It may need to be adjusted for different equipment.

1 Switch off the algal shaker

Clear out any culture flasks from the algal shaker.

Move it onto a bench.

- 2 Work in a dark room. Turn all lights in the room off. Keep them off during the measurements.
- 3 Connect the Walz probe to the Li-COR Light Meter.

Set the Li-COR Light Meter calibration to measure in air (-274.3). This value is specific to the particular probe and will be different for other probes. The value in the calibration data for specifically for the utilized probe must be used.

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4 Place the probe on a random spot on top of the illuminated shaker platform and record the value.

Hold the probe vertically upright and keep your hand as far away from the illuminator as possible to minimize effect of light scattered from the hand and sleeve.

Repeat this in twelve different random places. Find a random location by doing it with eyes closed or looking away to avoid personal bias.

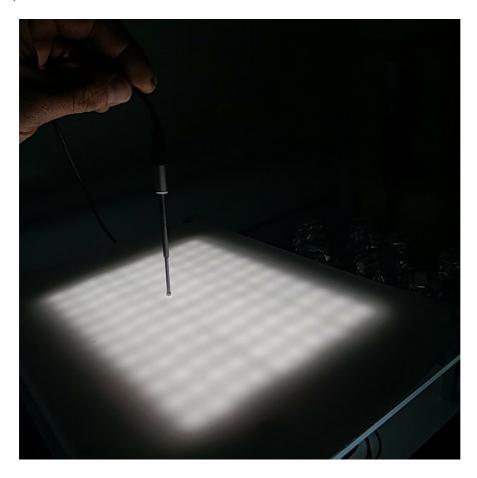


Photo showing how the proble is held perpendicularly to the surface of the illuminated orbital shaker platform with the hand as far as possible to minimize the contribution of the light scattering from the hand.

5 Repeat the above step for all eight LED current settings.

6 Record all PPFD measurements in this table:

Repeat		LED Current Setting									
	Tric kle		1	2	3	4	5	6			
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12											

Photosynthetic Photon Flux Density in Air. Table to record twelve repeats of PPFD measurements for eight LED current settings done in air on the surface of the illuminated orbital shaker platform.

- Set the Li-COR Light Meter calibration to measure in water (-471.9). This value is specific to the particular probe and will be different for other probes. The value in the calibration data for specifically for the utilized probe must be used.
- 8 Place a 100 ml Erlenmeyer flask onto the illuminated shaker platform.
- 9 Place the probe into the flask fix the flask mouth with some cotton wool, replicating the plug normally used in the culture. Try to keep the probe vertical.

Record the value.

Move the flask to another random spot, repeating the measurement in twelve different places. Find a random location by doing it with eyes closed or looking away to avoid personal bias.



Photo showing how the proble is held perpendicularly to the surface of the illuminated orbital shaker platform inside an Erlenmeyer flask with deionized water. Cotton wool plug in the neck of the flasks holds the probe and also reproduces the scattering from the cotton wool used to plug the flasks with the microalgae cultures.

10 Repeat the above step for all eight LED current settings.

5

#### 11 Record all PPFD measurements in this table:

Repeat		LED Current Setting									
	Tric kle	•	1	2	3	4	5	6			
1											
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Photosynthetic Photon Flux Density in Water. Table to record twelve repeats of PPFD measurements for eight LED current settings done in water inside an Erlenmeyer flask on the surface of the illuminated orbital shaker platform.

#### 12 References

- LI-250A Instruction Manual
- US-SQS/L Instruction Manual

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