

# **AER<sup>®</sup> Bioreactor Designs**

## **I. The f<sub>i</sub> AER Bioreactor:**

- a. Two 1-W LED (one red , one blue) illuminating**
- b. A 10 gallon tank with lid-portal, reflective lid, reflective base and walls**
- c. 0.40 sq in fresh spinach leaf on elevated wire mesh**
- d. 0.15 mole CO<sub>2</sub> released from 9" OD collection sac (30% CO<sub>2</sub> in the 10 gal tank)**
- e. Average temperature of 25° C (78° F)**
- f. Iodine to stain glucose in spinach cells**
- g. Compound light microscope with image retention device**

**Hypothesis: The AER Bioreactor will demonstrate synthesis and retention of excess glucose in spinach cells exposed to a high CO<sub>2</sub> concentration**

### **Method:**

- 1. Divide fresh spinach leaf into control and CO<sub>2</sub>-exposed samples**
- 2. Collect CO<sub>2</sub> (sterno emission) into a 9" OD sac**
- 3. Place demonstration samples on elevated wire mesh in the 10 gal tank**
- 4. Place CO<sub>2</sub> sac in tank. Withdraw sac through**

- lid-port and close port: tank's ~ 30% CO<sub>2</sub> <sup>(1)</sup>.*
- 6. Illuminate reflective chamber with red & blue LED for \_\_\_\_\_ minutes**
  - 7. Iodine-stain control and exposed samples**
  - 8. Retain 40X, 100X images of control and exposed samples**
  - 9. Compare stained-cell counts of control / exposed samples ( $\chi^2$ )**

<sup>(1)</sup> *Nishimura & Asakawa, Plant Physiol. 1978.*

## **II. The f<sub>ii</sub> AER Bioreactor:**

***This reactor design is for extraction of excess glucose from high CO<sub>2</sub>-exposed spinach cells.***

***The design's a modification of standard sugar beet glucose-extraction procedure:***

### **Method:**

- 1. Crush and float CO<sub>2</sub>-exposed leaf in water.**
- 2. Filter and compost leaf debris**
- 3. Process / concentrate sucrose solute**

**Philip Nolan**  
**[waupacacpr@gmail.com](mailto:waupacacpr@gmail.com)**

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