

CALIFORNIA STATE SCIENCE FAIR 2017 PROJECT SUMMARY

Name(s)	Project Number
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	31900
Project Title Cloning African Violets through Autotrophic Tissue Culture	
Objectives/Goals Abstract	
Traditionally, tissue cultured plantlets use sugar provided in the mediu	Im for energy because the jars in
which they are housed are sealed. My objective was to test if sections of African violet tissue would grow	
at an accelerated pace and have a decreased chance of contamination when provided a sugariess	
Methods/Materials	
Materials include CO(2) testing solution to determine gas permeability of various plastics, heat sealer to	
make small plastic bags, 60 glass vials, pressure cooker to sterilize, 4 g Murashige and Skoog medium,	
1-Naphthaleneacetic acid (synthetic auxin hormone), 6-Benzylaminopurine (synthetic cytokinin), and	
coconut water (natural cytokinin). The $CO(2)$ system is constructed from a bottle with yeast sugar, and water attached to a bubble counter to	
gauge gas production connected by airline tubing to the growth chamber. Ninety sterilized plant sections	
were housed individually in glass yials covered with gas-permeable Ziploc bag plastic (Autotrophic 1, or	
A1), small bags made from Ziploc plastic (Autotrophic 2, or A2), and vials sealed by the original screw	
cap with added sugar (Mixotrophic, or M).	
Results	
bigher than those of M. However, the explants in the M experimental group haddeveloped calli	
(undifferentiated tissue, the precursor to shoots and roots) with larger biomasses. On the other hand,	
regardless of the original hormone supplement provided, 83% of the calli in autotrophic conditions	
differentiated into green shoots with the potential of maturing into adult plants, while 67% of the M group	
produced roots, which are more difficult to work with and have less p	otential.
Conclusions/Discussion I developed a new method of plant propagation through the use of Ziploc bags and $CO(2)$ generated from	
materials adapted from the fishkeeping hobby. My original hypothesis was partially supported. A1 was	
the most successful because of its structural stability, while A2 was similar but secondary in success and	
the M group was least productive. This unique, cost-effective technique may be applied to the cultivation	
of plant medicines, production of economically significant crops, prop	bagation of fragile or sterile plants,
and conservation of endangered species.	
Summary Statement	
This project investigates the effect of varying levels of gas permeabili	ty and carbon sources on the growth
and differentiation of cloned African violet plantlets.	
Help Received	
I designed and performed the experiment at home by adapting information from previous publications	
focused on different plants and procedures.	