



**WORLD POTATO  
CONGRESS**

# Speaker Disclosure

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# Unveiling the Potential of 2*E*-Hexenal: A Breakthrough in Potato Storage Disease Management



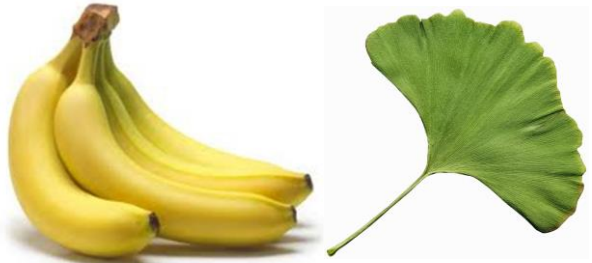
**Phillip Wharton**

**Associate Professor of Potato Pathology, University of Idaho**

**Katie Malek & Alan Malek, University of Idaho; John Immaraju, AMVAC**

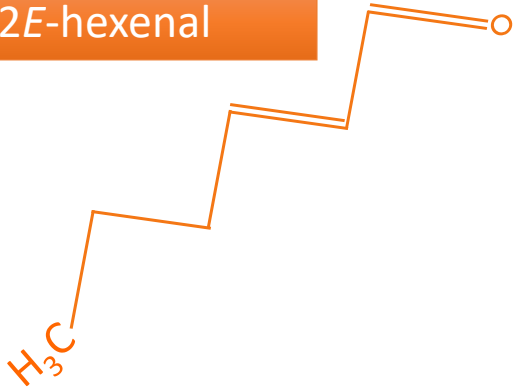


# Volatile compounds to control disease



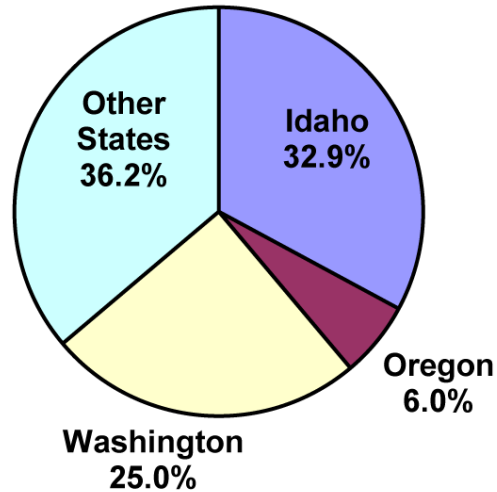
- 2*E*-hexenal is a naturally produced volatile compound (lipoxygenase pathway)
- Previous studies have shown anti-fungal and anti-bacterial properties
- Approved by the FDA as a food/flavor additive
- Volatile nature

2*E*-hexenal



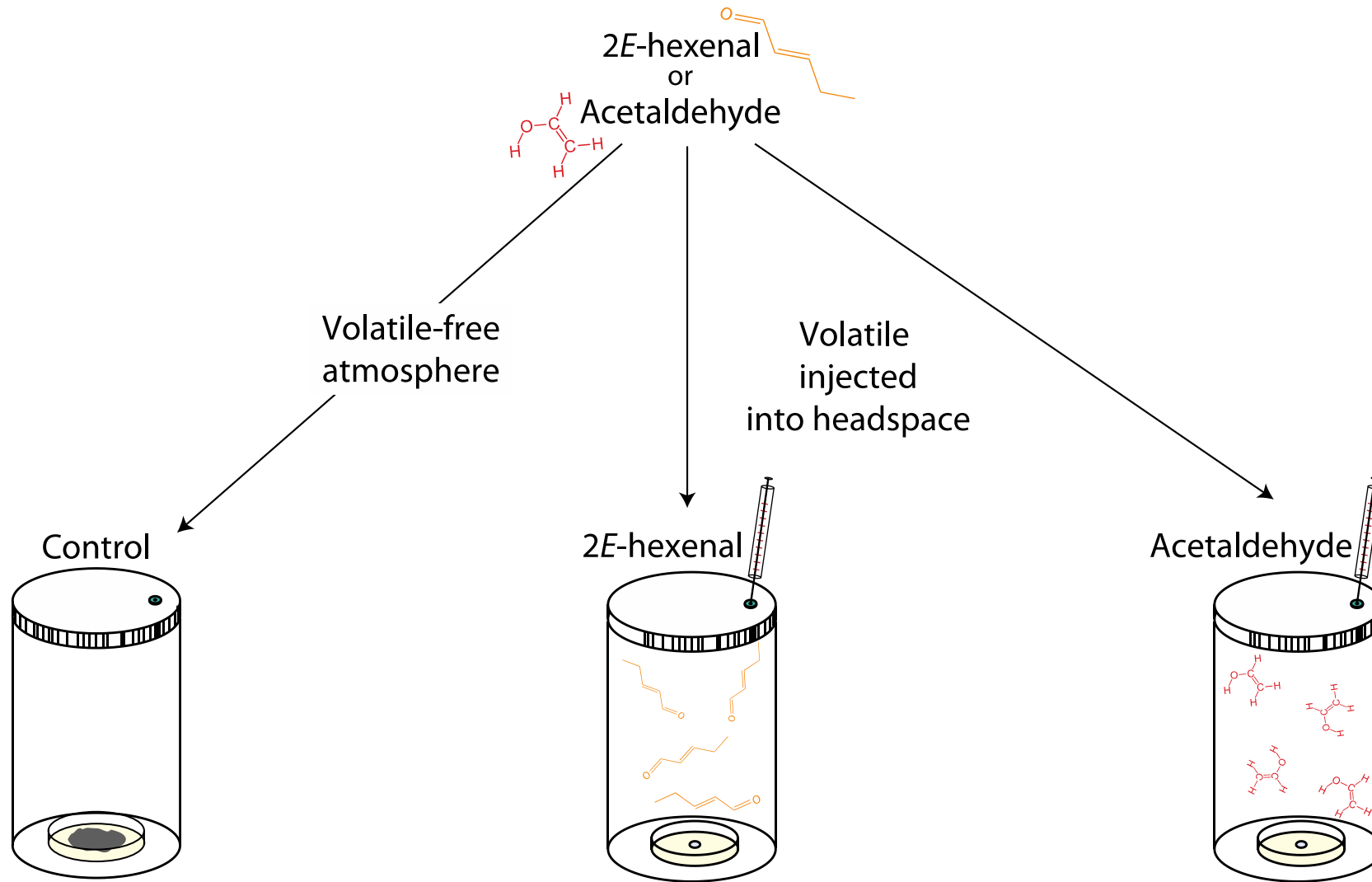
# Market overview

Percent of Fall Potato Production  
(Percents do not add to 100 due to rounding.)

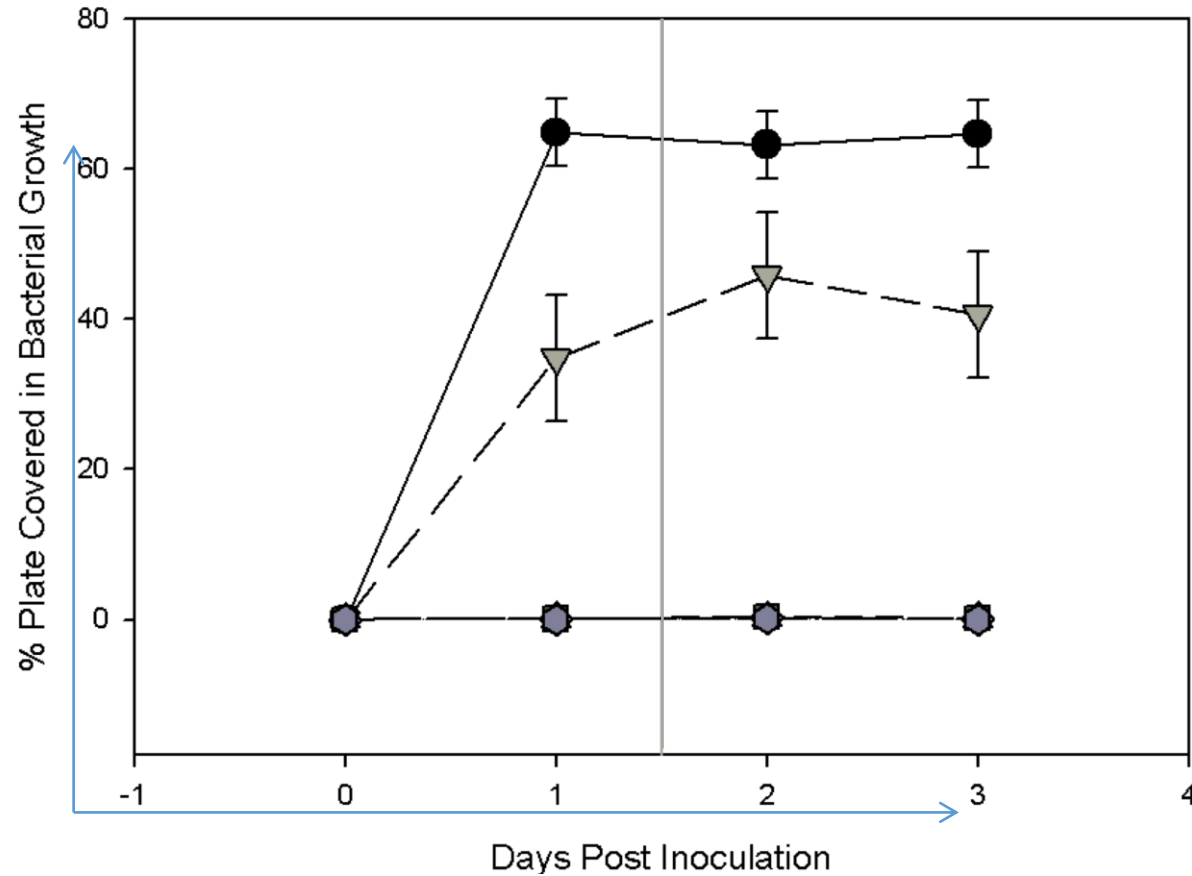
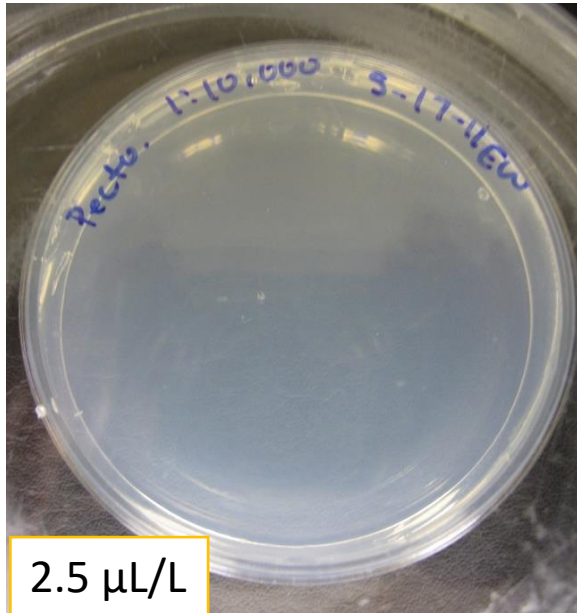


- On average, Idaho growers produced 7 million metric tonnes of potatoes valued at \$1.05 billion or approx. 5% of the Idaho economy.
- On average, 500,000 tonnes (\$73 million) of the potatoes placed in storage are lost to diseases and shrinkage.
- Growers have few effective options for the control of potato diseases once potatoes have been placed in storage.
- At the University of Idaho, our research showed that the anti-fungal plant volatile 2*E*-hexenal is highly effective in controlling potato diseases in storage.

# *In vitro* methods



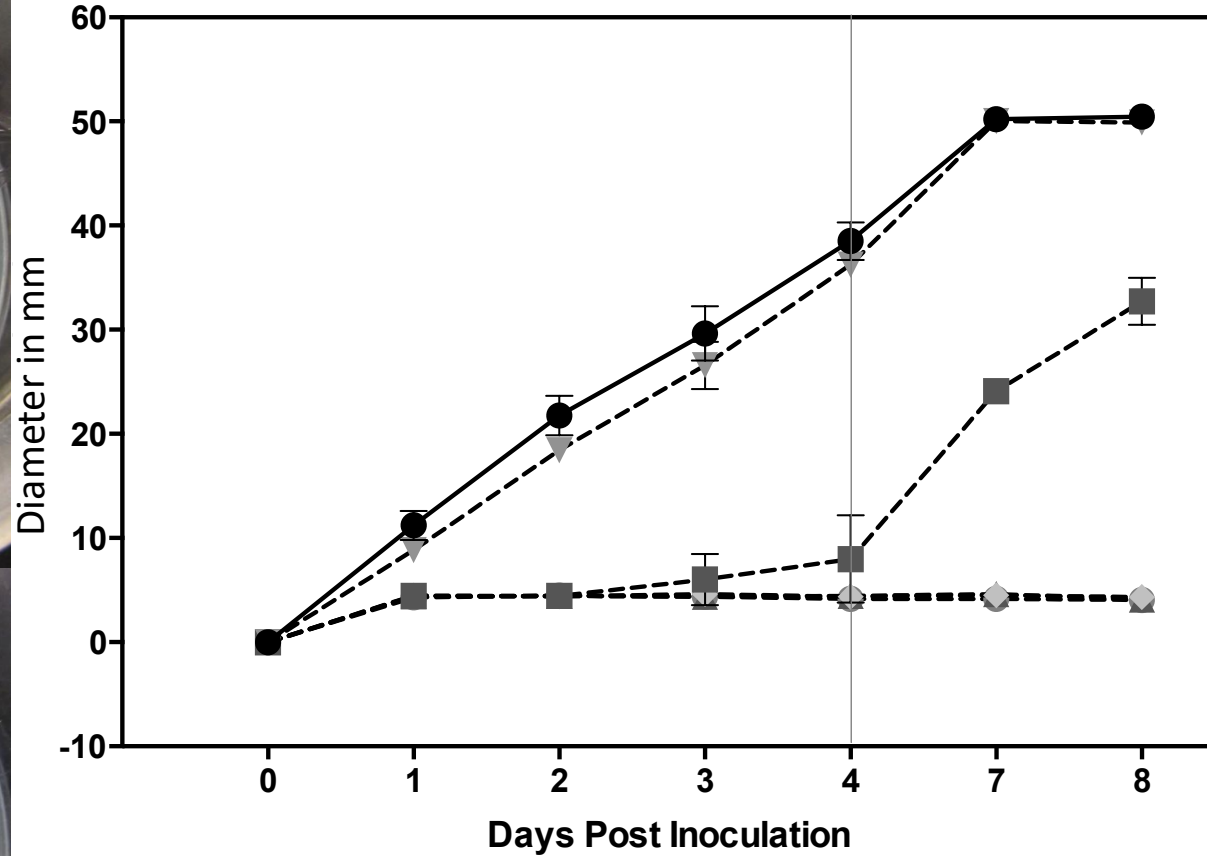
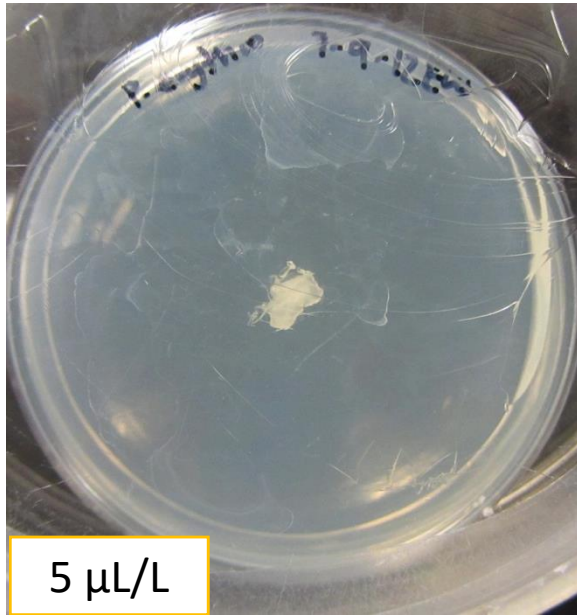
(●) UTC (▼) IS Check (■) 2.5  $\mu\text{L/L}$  (◆) 5  $\mu\text{L/L}$  (▲) 7.5  $\mu\text{L/L}$  (◆) 10  $\mu\text{L/L}$



Results: *Soft rot*

2.5  $\mu\text{L/L}$  of 2E-hexenal was capable of inhibiting growth of *P. atrosepticum* completely *in vitro*.

(●) UTC (▼) IS Check (■) 2.5  $\mu\text{L/L}$  (◆) 5  $\mu\text{L/L}$  (▲) 7.5  $\mu\text{L/L}$  (●) 10  $\mu\text{L/L}$

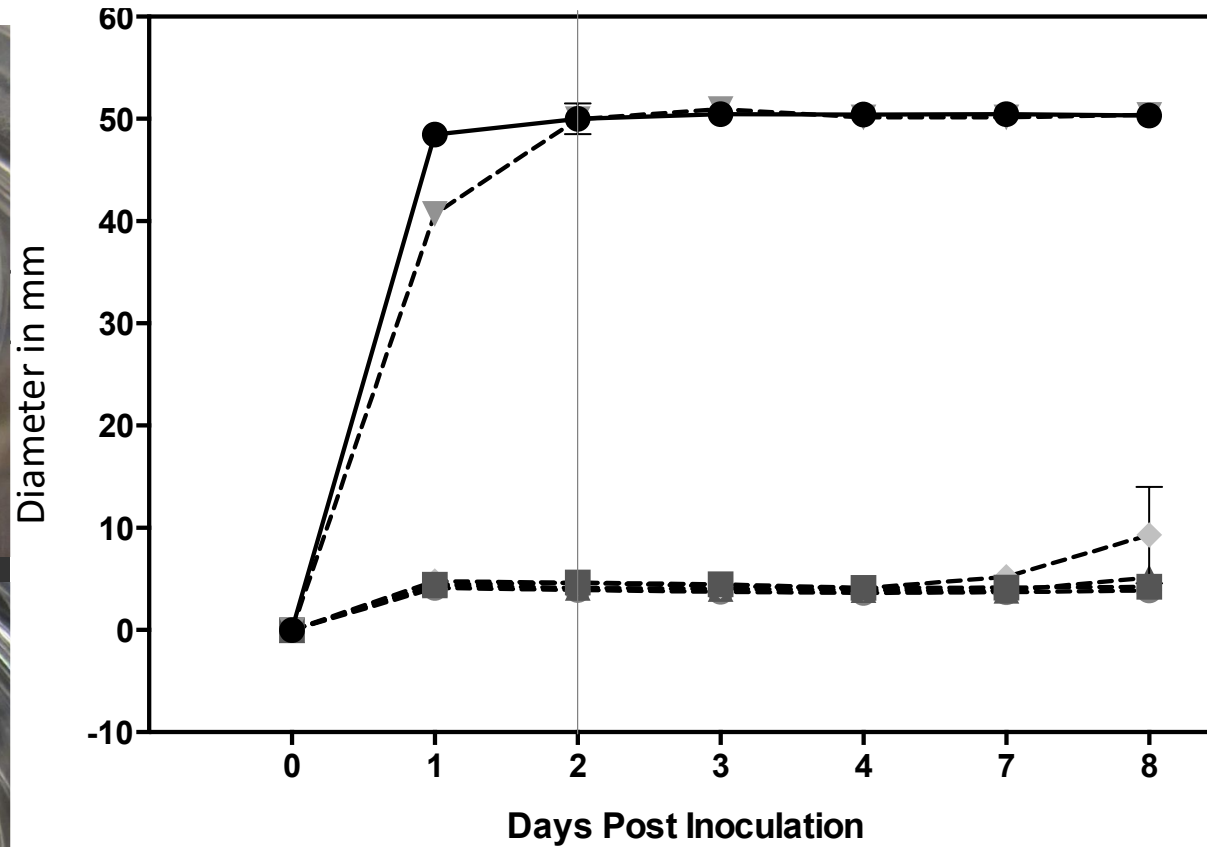
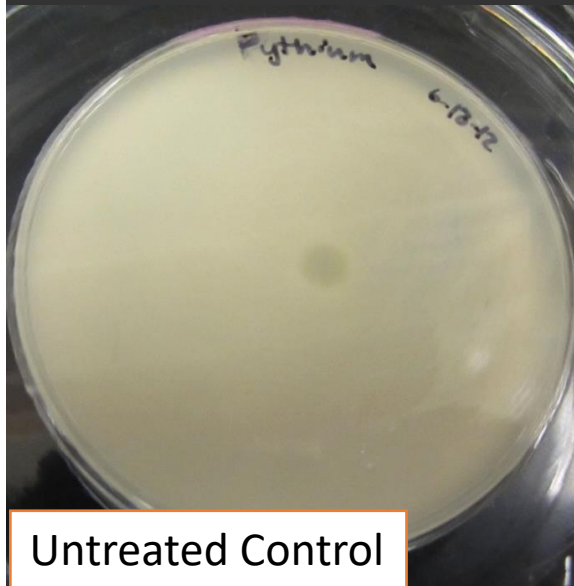
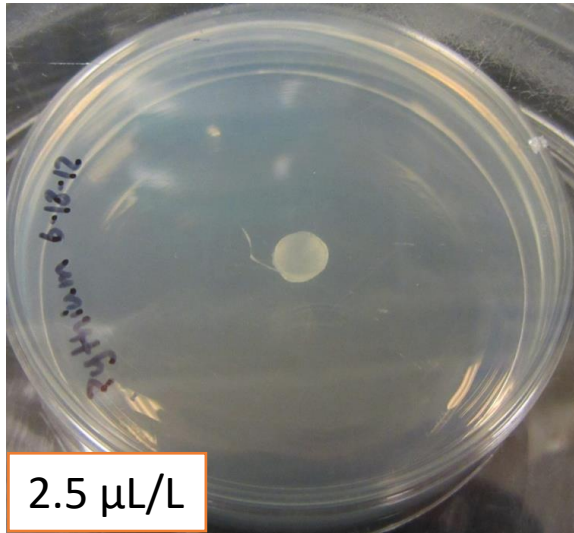


## Results: *Pink Rot*

2.5  $\mu\text{L/L}$  was capable of slowing *P. erythroseptica* growth, 5  $\mu\text{L/L}$  capable of inhibiting growth completely *in vitro*.



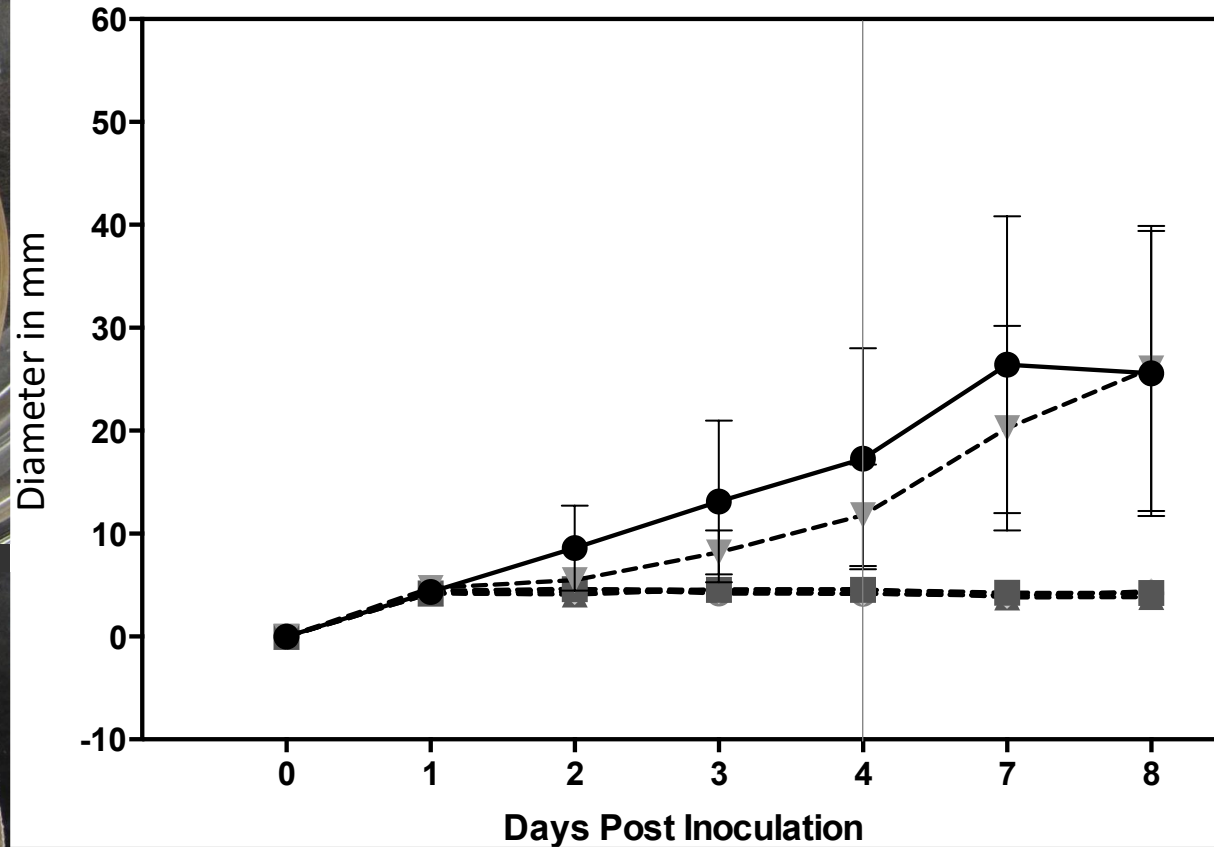
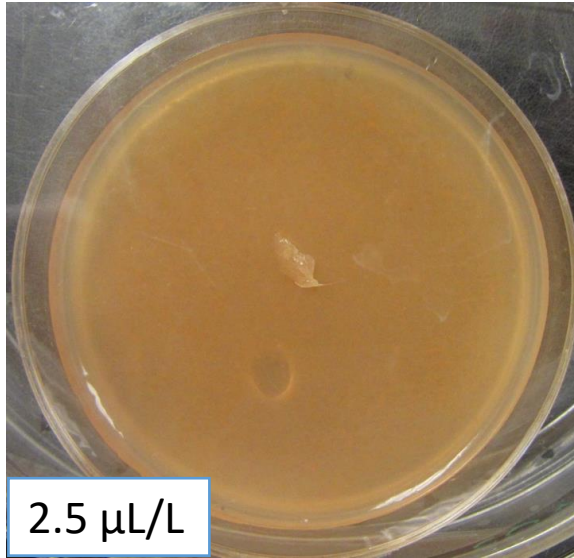
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## Results: *Pythium Leak*

2.5  $\mu\text{L/L}$  was capable of capable of inhibiting *P. ultimum* growth completely *in vitro*.

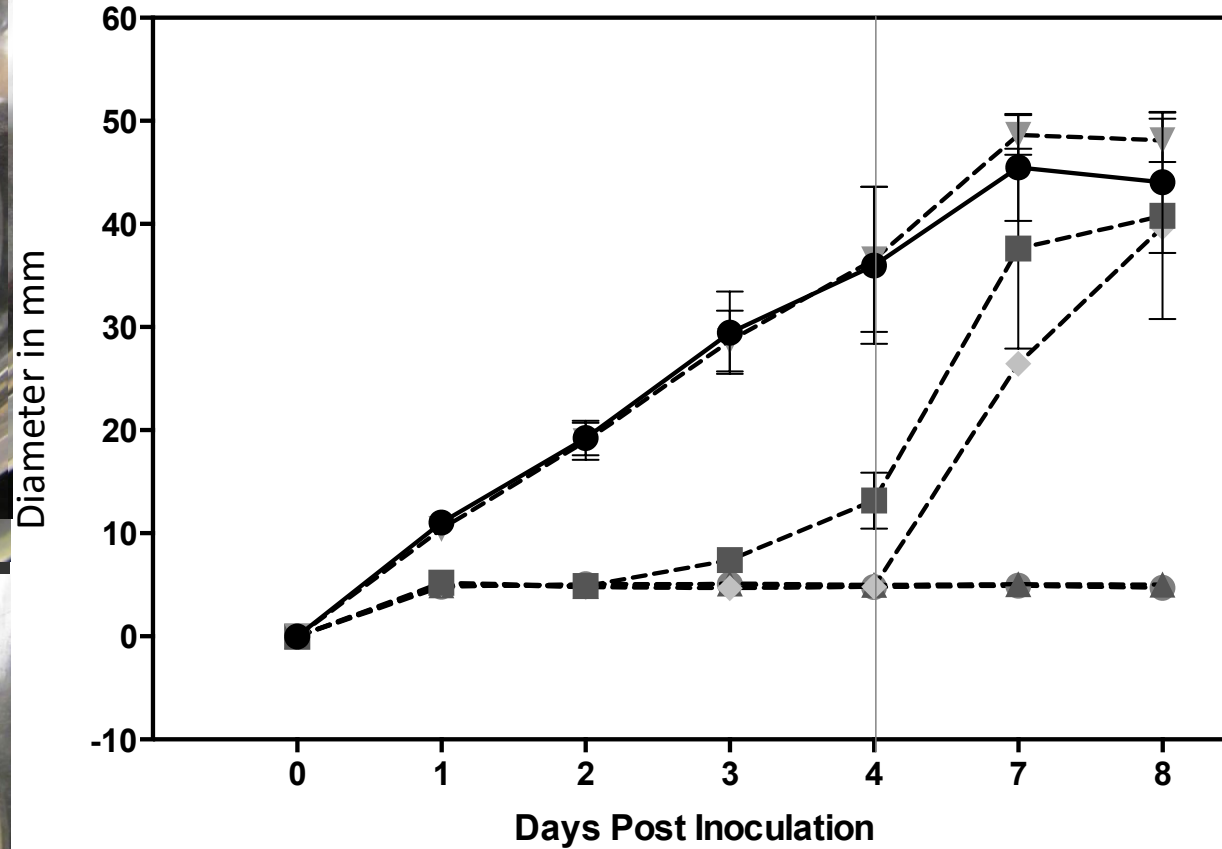
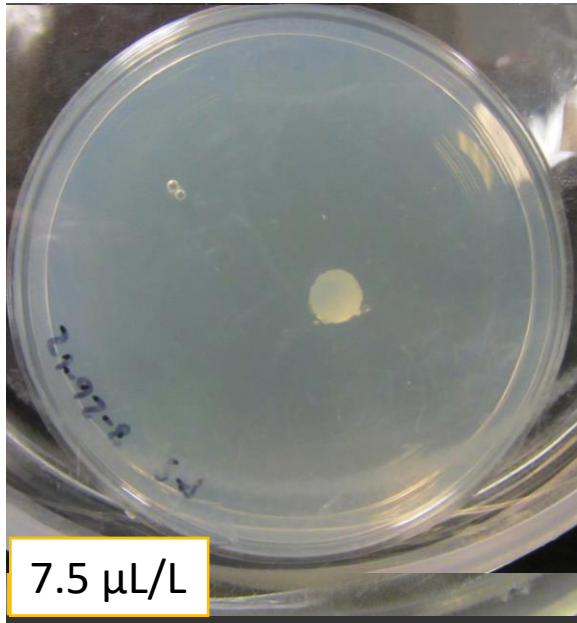
(●) UTC (▼) IS Check (■) 2.5  $\mu\text{L/L}$  (◆) 5  $\mu\text{L/L}$  (▲) 7.5  $\mu\text{L/L}$  (◆) 10  $\mu\text{L/L}$



## Results: *Late blight*

2.5  $\mu\text{L/L}$  was capable of capable of inhibiting *P. infestans* growth completely *in vitro*.

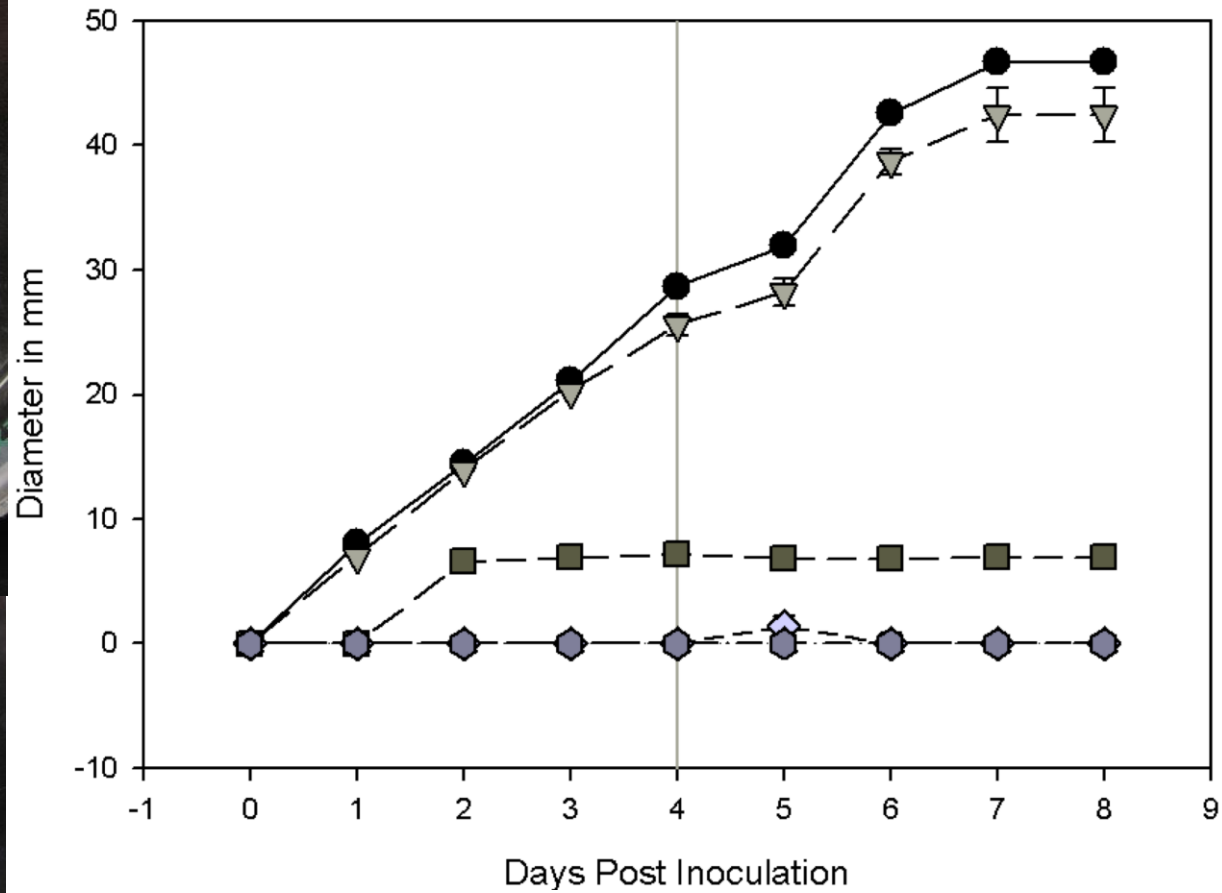
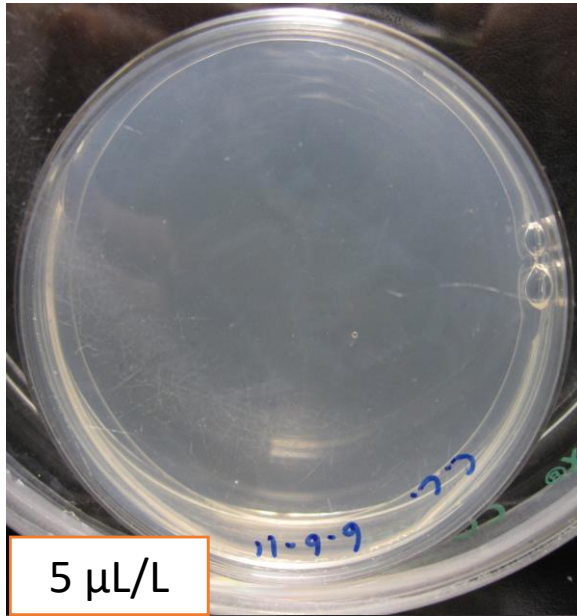
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## Results: *Fusarium dry rot*

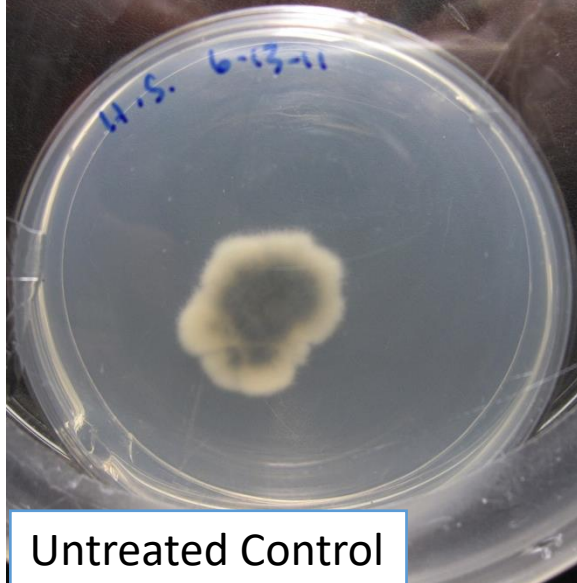
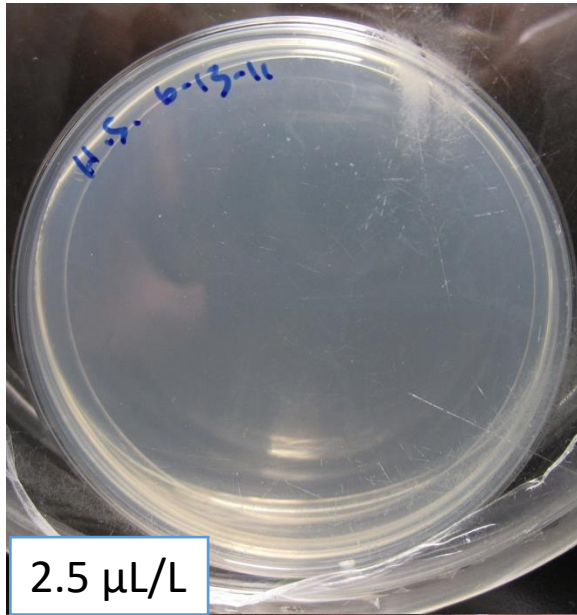
7.5  $\mu\text{L/L}$  was capable of inhibiting *F. sambucinum* growth completely *in vitro*, with 2.5 and 5  $\mu\text{L/L}$  able to slow the growth of the pathogen *in vitro*.

(●) UTC (▼) IS Check (■) 2.5  $\mu\text{L/L}$  (◆) 5  $\mu\text{L/L}$  (▲) 7.5  $\mu\text{L/L}$  (◆) 10  $\mu\text{L/L}$

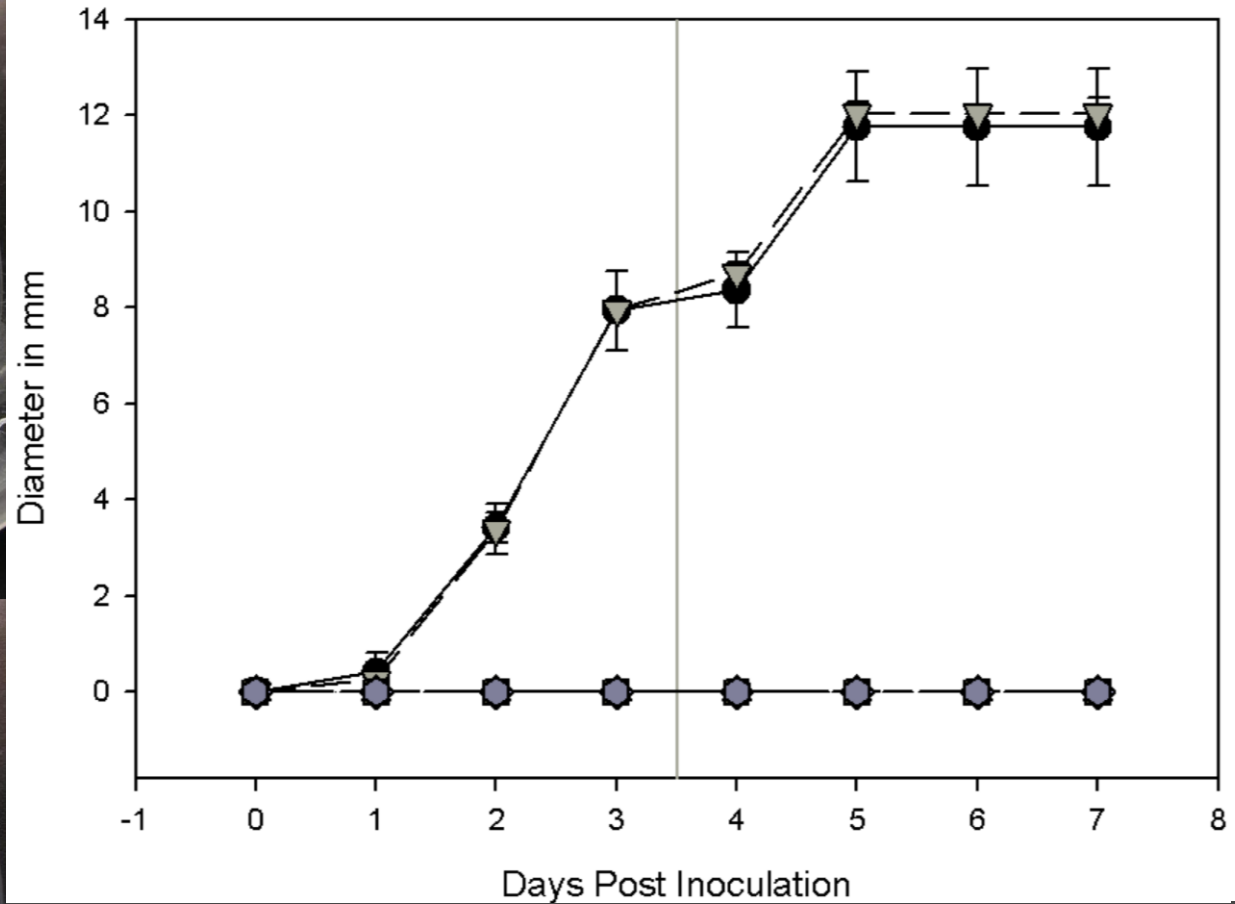


Results: *Black dot*

2.5  $\mu\text{L/L}$  of 2E-hexenal was capable of slowing *C. coccodes* growth, but 5  $\mu\text{L/L}$  completely inhibited growth *in vitro*.



(●) UTC (▼) IS Check (■) 2.5  $\mu\text{L/L}$  (◆) 5  $\mu\text{L/L}$  (▲) 7.5  $\mu\text{L/L}$  (●) 10  $\mu\text{L/L}$



Results: *Silver scurf*

2.5  $\mu\text{L/L}$  of 2E-hexenal was able to inhibit the growth of *H. solani* completely *in vitro*.

# Storage Rot Inoculation Methods

Tubers were inoculated by immersion in an inoculum suspension of either *Phytophthora erythroseptica* (pink rot) or *Pythium ultimum* (Pythium leak), or *Fusarium sambucinum* (dry rot)



# Inoculation results



# Storage Trial Methods

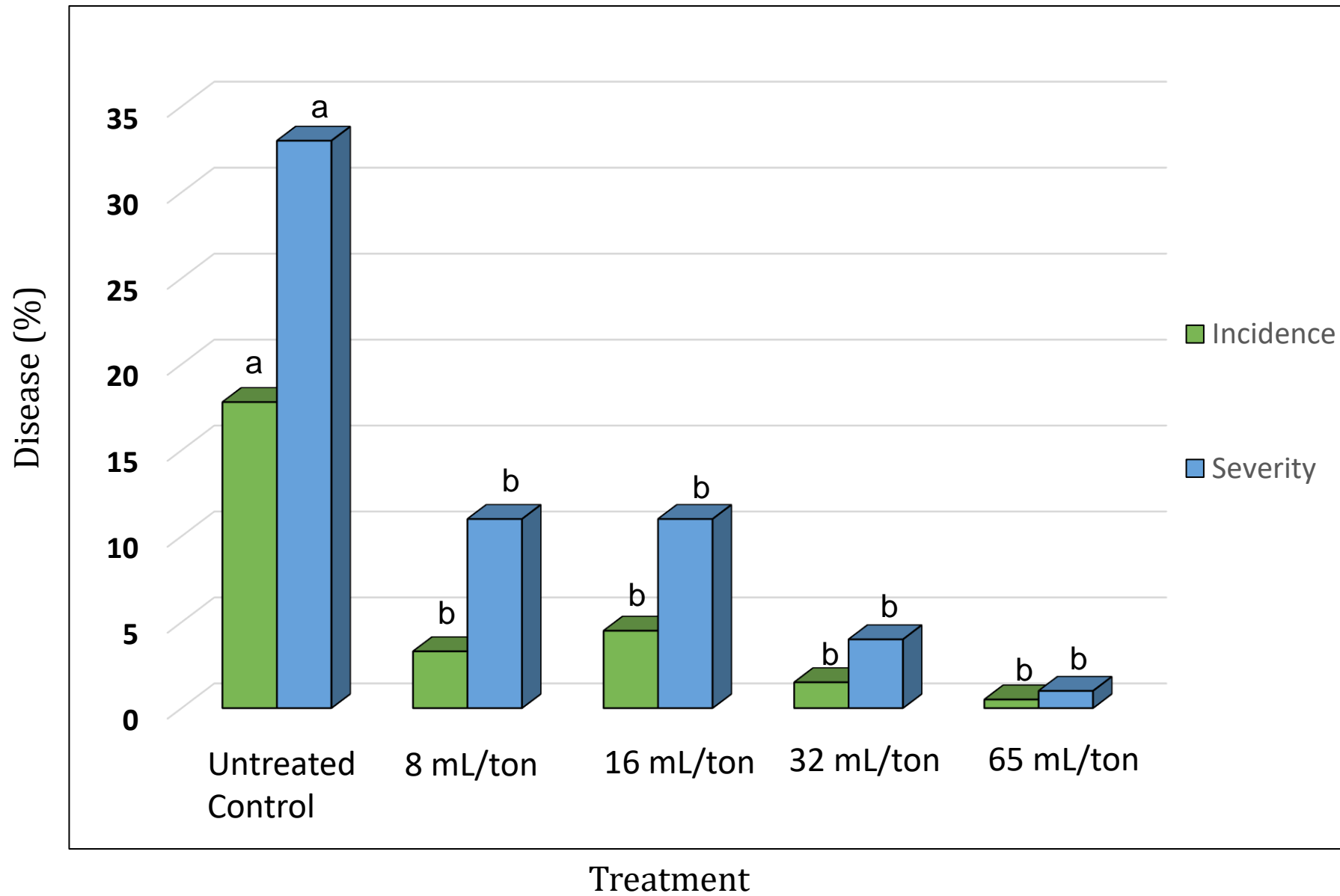
- Treatments consisted of 4 replicates of 25 tubers (cv. Russet Burbank) in a fully randomized design. For Fusarium dry rot experiments all tubers were inoculated. For Pythium and pink rot experiments one inoculated tuber was placed with 25 healthy tubers.
- Onion sacks containing the tubers were placed in 113-liter barrels and treated with *2E*-hexenal.
- Tubers were fumigated for 16 hours. After fumigation, tubers were stored at 16 C for 3 weeks before being rated.
- Treatment rates of *2E*-Hexenal were 0 (control), 8 mL/ton, 16 mL/ton, 32 mL/ton and 65 mL/ton.
- Tubers were rated for volume of internal rot (severity) and incidence.



# Treatment apparatus



# Pink rot Results



# Pink rot example results

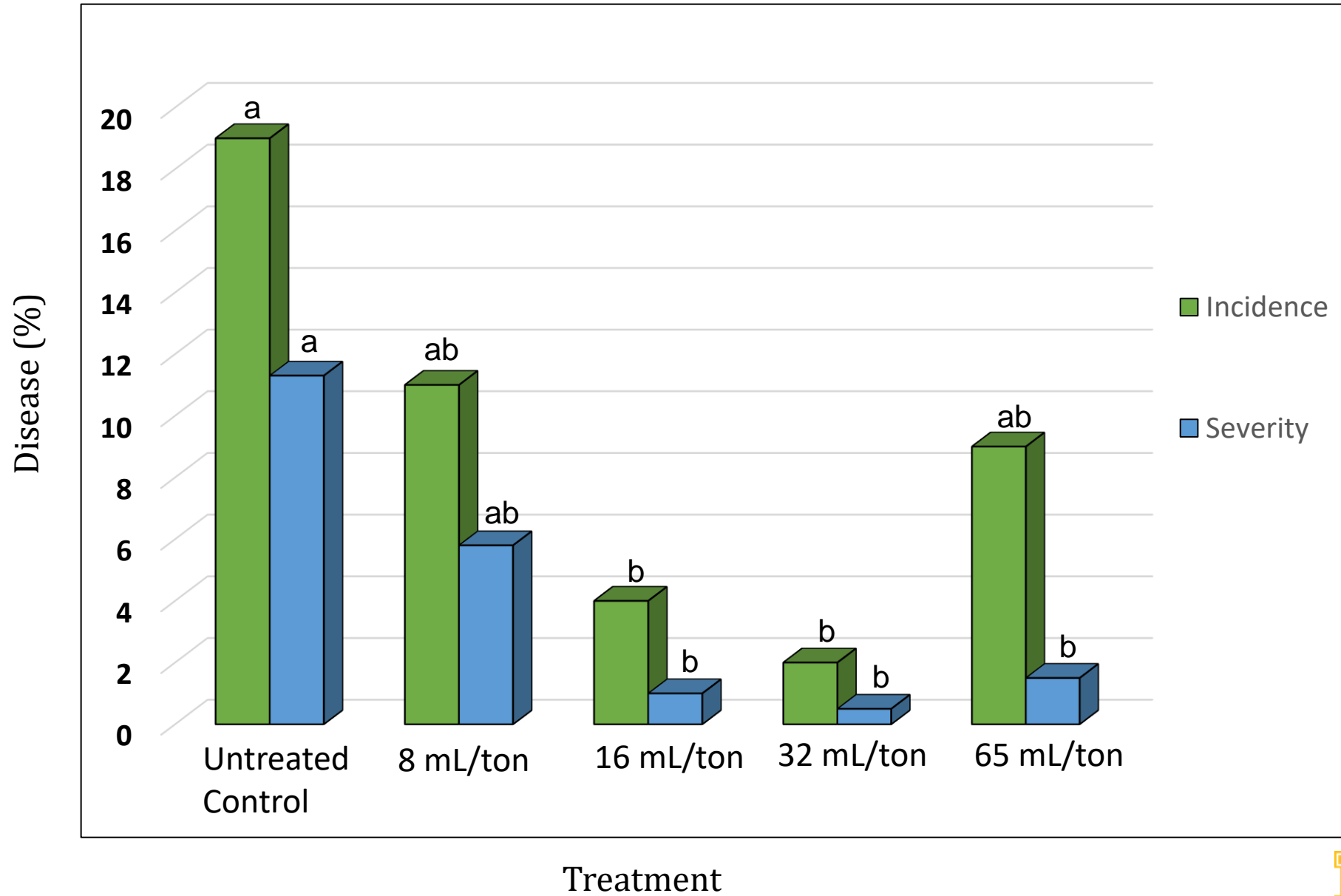


Untreated



Treated (32 ml/tonne)

# Pythium Results



# Pythium example results

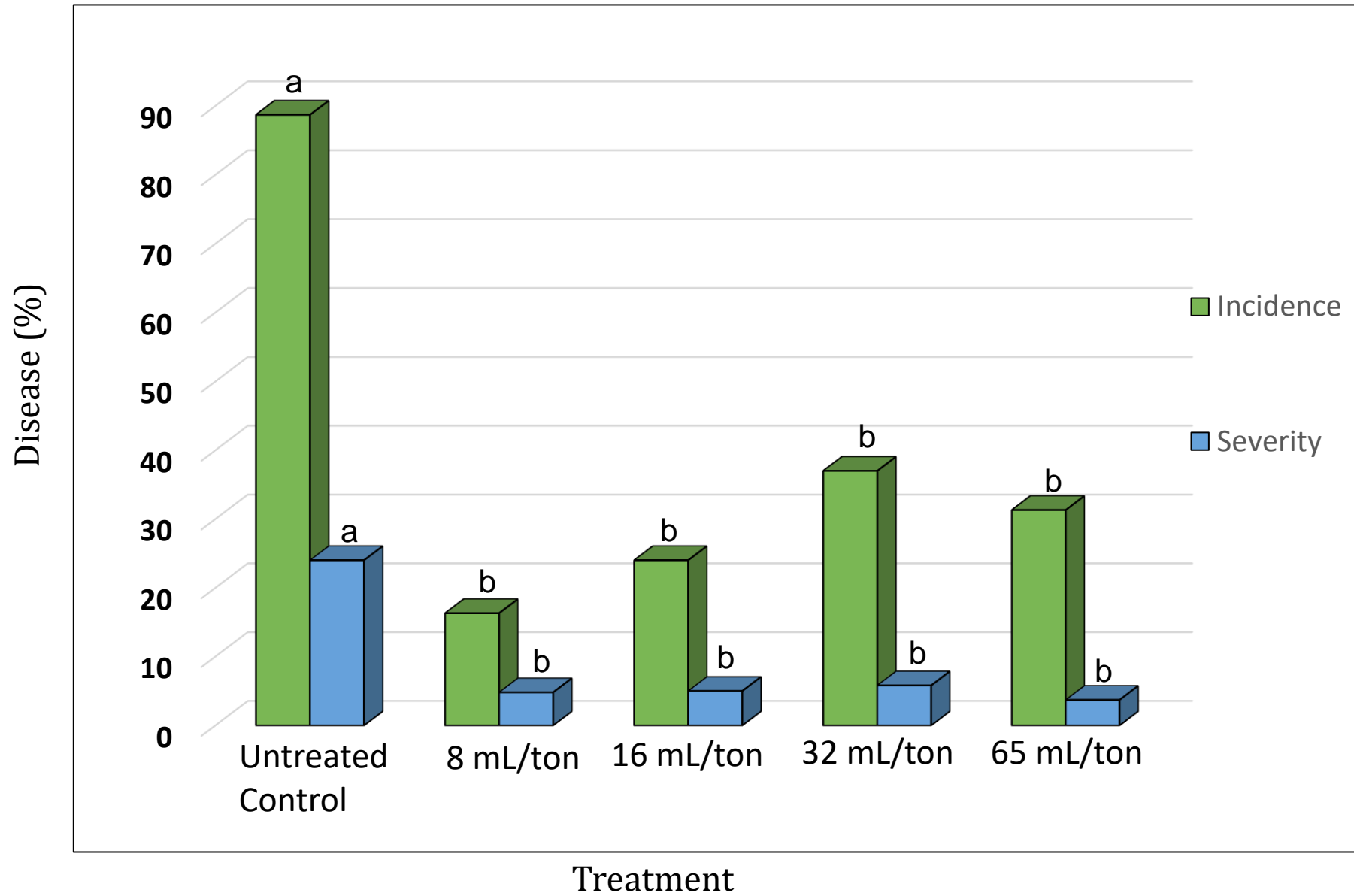


Untreated



Treated (16 ml/tonne)

# Fusarium Results



# Dry rot example results



Untreated



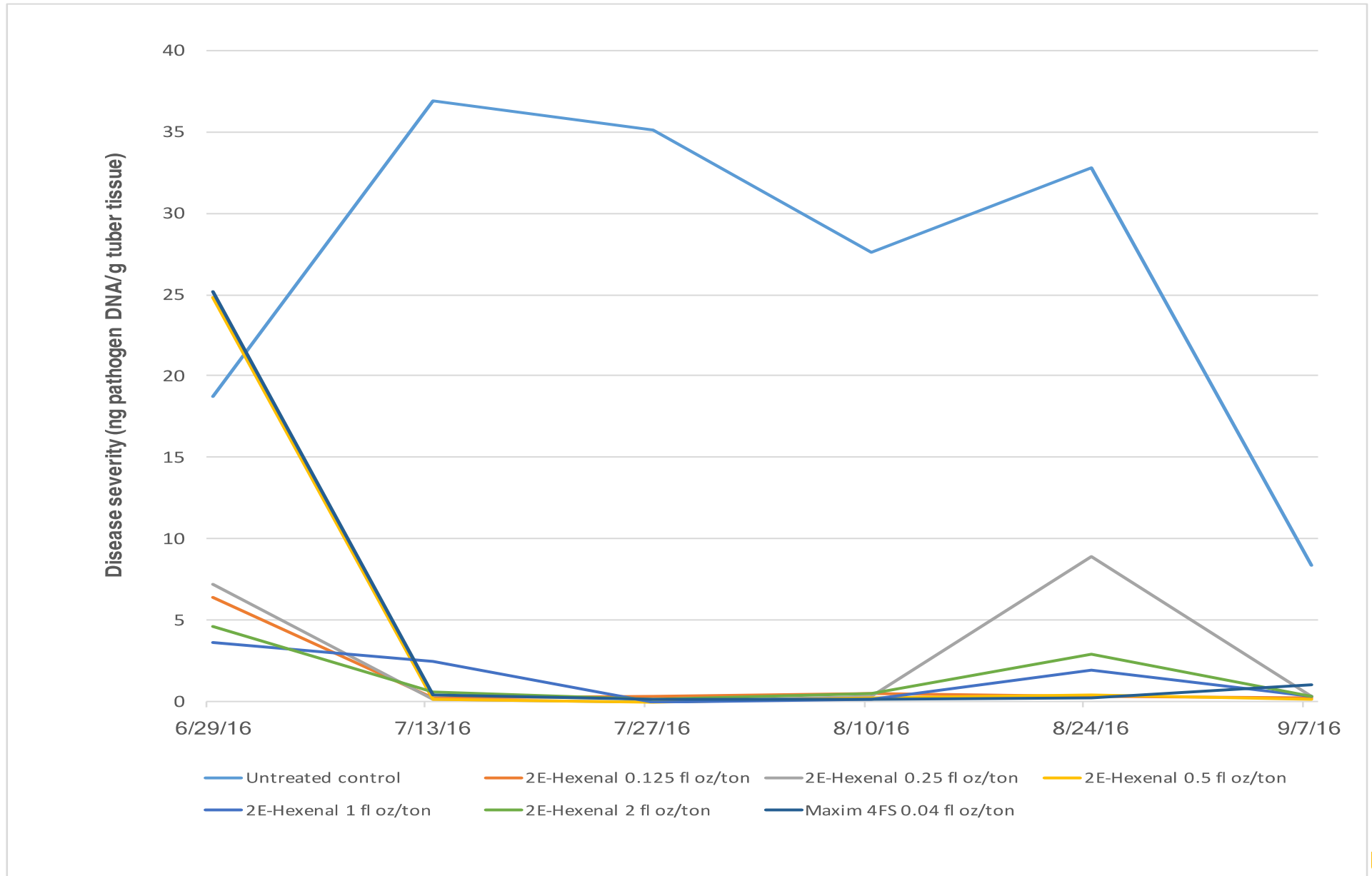
Treated (8 ml/tonne)

# Blemish disease methods

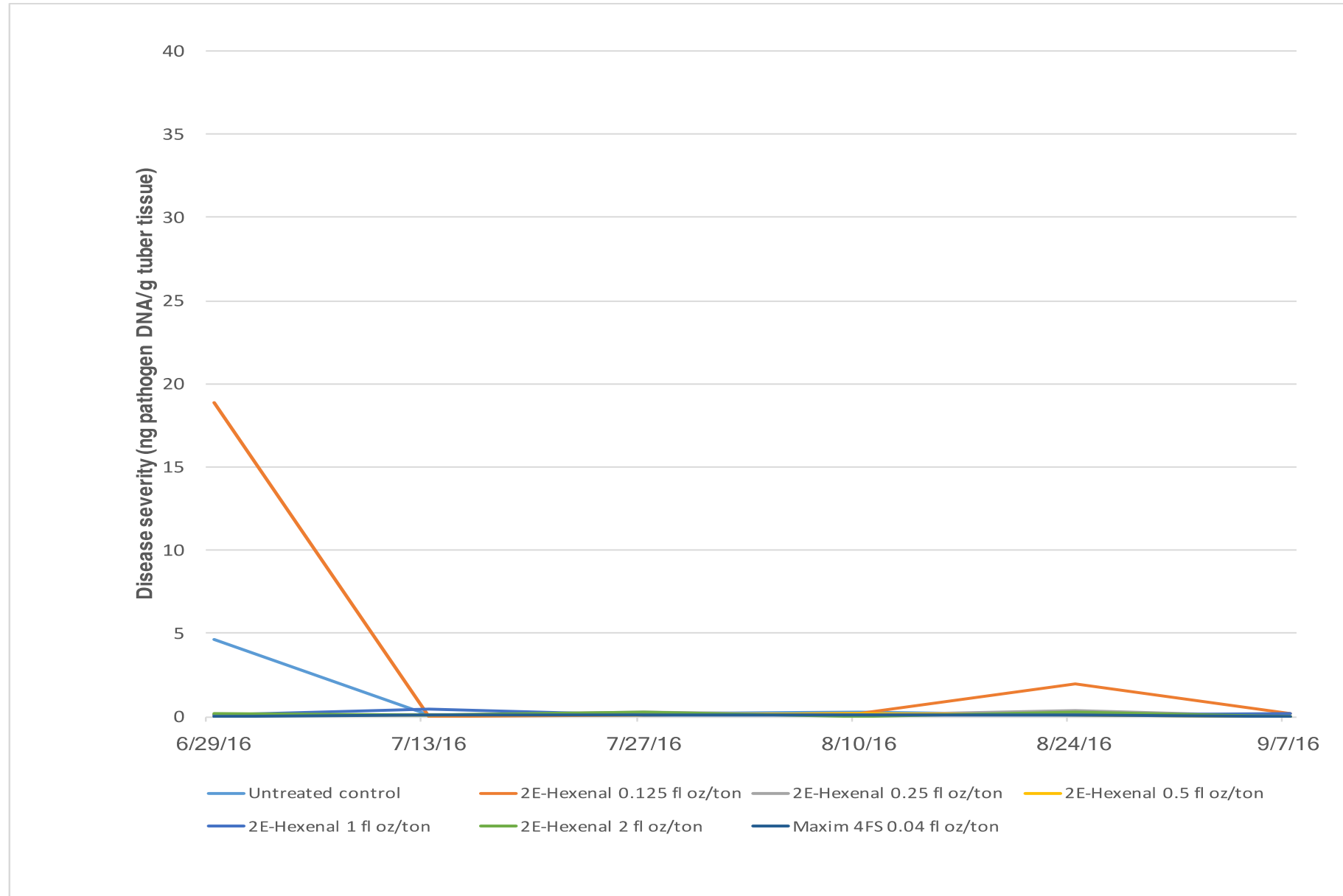
- Treatments consisted of 4 replicates of 25 tubers in a fully randomized design.
- Tubers (cv. Melody) naturally infected with black dot or silver scurf were used.
- Onion sacks containing the tubers were placed in 113-liter barrels and treated with *2E*-hexenal.
- Tubers were fumigated for 16 hours. After fumigation, tubers were stored at 12 C for 12 weeks before being rated.
- Treatment rates of *2E*-Hexenal were 0 (control), 8 mL/ton, 16 mL/ton, 32 mL/ton and 65 mL/ton.
- Levels of pathogen DNA on the tuber surface was quantified by qPCR every two weeks.



# Silver Scurf results



# Black dot results



# Overall Conclusions

- *In-vitro*, as little as 2.5  $\mu\text{L/L}$  of 2*E*-hexenal is effective at inhibiting growth of all potato storage pathogens.
- 2*E*-hexenal significantly reduced disease incidence and severity in tubers placed in storage with tubers inoculated with pink rot or pythium or inoculated with Fusarium dry rot.
- Over a 3-month period, the levels of silver scurf DNA in naturally infected tubers were significantly lower in those treated with 2*E*-hexenal.
- For black dot, there was no significant difference between treated and non-treated tubers.



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