

# Effects of a Commercial Soil Amendment on Plant Parasitic Nematodes

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# Soil Ecology and Biological Control

*The main scientific goal of our lab is to learn how to more sustainably manage agricultural systems to control pests and improve soil health.*

## Pest Detection

qPCR-Carrot  
nematodes and  
Orchard  
nematodes

## Cultural Control

Soil  
Amendments -  
compost, food  
hydrolysate

## Innundative Control

Biobased  
pesticides,  
Insect parasitic  
nematodes

# Applied nematode ecology: Ecological effects of an organic Amendment - stabilized food hydrolysate

- Unsold food collected from supermarkets in California.
- Digested with enzymes to produce a stabilized product consisting of amino acids, simple sugars, fatty acids, and minerals
- Marketed as a liquid fertilizer product to improve soil health.



# Stabilized food hydrolysate

*Fertilizer or organic matter amendment?*



Bacterial activity

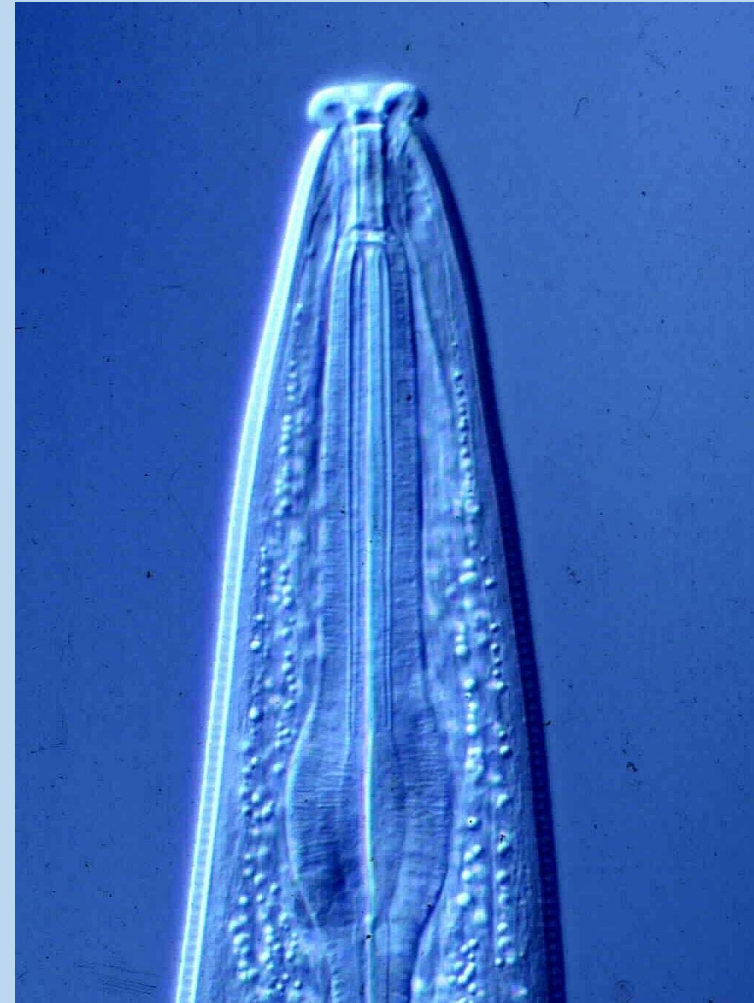
Organic matter breakdown

Nutrient availability

*How does H2H affect soil biological communities?*

# Nematode life history strategies

- Bacterial feeders
  - Indicate microbial nutrient processing and N mineralization
- Fungal feeders
- Omnivores
- Predators
- Plant Parasites



# Raised bed experiment



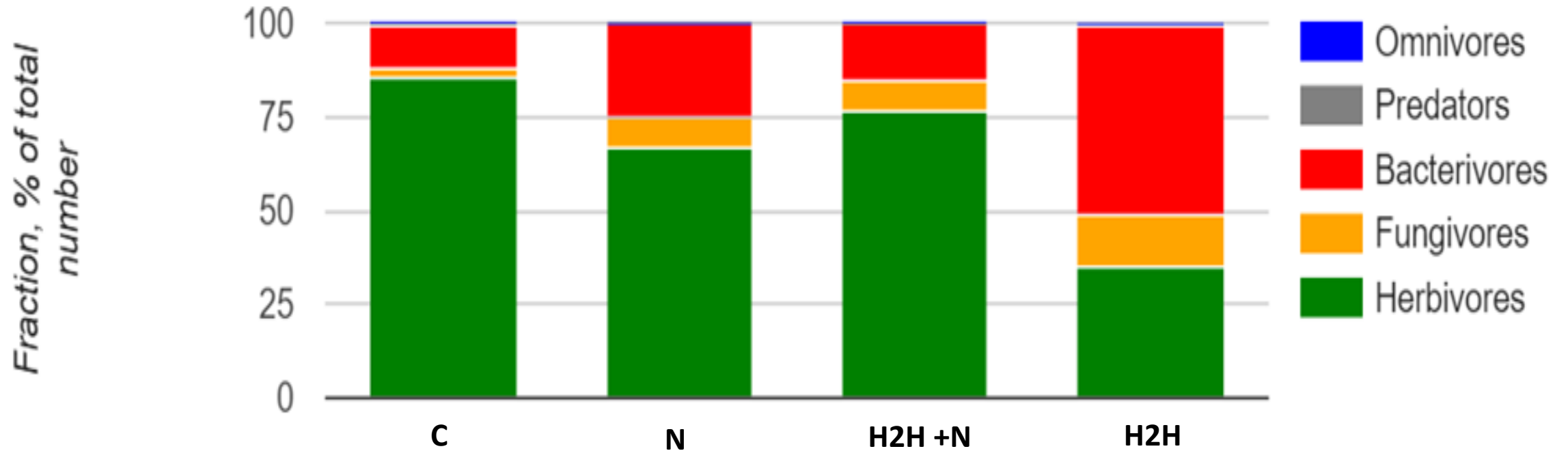
- Four replicated blocks of each treatment
- 11 plants/block

Name	Material	Rate	Source
H2H	Stabilized food hydrolysate	30 gal/acre	California Safe Soil, McClellan CA
UAN-32	Nitrogen fertilizer (7.75% Ammoniacal Nitrogen, 7.5% Nitrate Nitrogen, 16.50% Urea Nitrogen)	150 lbs N/acre	Simplot, Lathrop CA
H2H+UAN-32 (1:1)	Stabilized food hydrolysate + Nitrogen fertilizer (7.75% Ammoniacal Nitrogen, 7.5% Nitrate Nitrogen, 16.50% Urea Nitrogen) (1:1)	(15 gal/acre) + (75 lbs N/acre)	California Safe Soil, McClellan CA; Simplot, Lathrop CA
Control	Water	3.45 gph	Davis, CA



Family	Genus	Feeding habit
Rhabditidae	<i>Rhabditis</i>	bacterivores
	<i>Diploscapter</i>	bacterivores
Diplogasteridae	<i>Diplogaster</i>	bacterivores
Cephalobidae	<i>Cephalobus</i>	bacterivores
	<i>Eucephalobus</i>	bacterivores
	<i>Acrobeles</i>	bacterivores
	<i>Acrobeloides</i>	bacterivores
Plectidae	<i>Cervidellus</i>	bacterivores
	<i>Plectus</i>	bacterivores
Monhysteridae	<i>Anoplectus</i>	bacterivores
Prismatolaimidae	<i>Monhystera</i>	bacterivores
Alaimina(Alaimidae)	<i>prismatolainus</i>	bacterivores
Aphelenchoididae	<i>Alaimus</i>	bacterivores
Aphelenchidae	<i>Aphelenchoides</i>	fungivores
Dorylaimidae	<i>Aphelenchus</i>	fungivores
Tylenchidae	<i>Dorylaimus</i>	Omniovores
	<i>Tylenchus</i>	Herbivores
	<i>Filenchus</i>	Herbivores
Paratylenchidae	<i>Tetylenchus</i>	Herbivores
Tylenchorhynchidae	<i>Paratylenchus</i>	Herbivores
Pratylenchidae	<i>Tylenchorhynchus</i>	Herbivores
Heteroderidae	<i>Pratylenchus</i>	Herbivores
	<i>Meloidogyne</i>	Herbivores

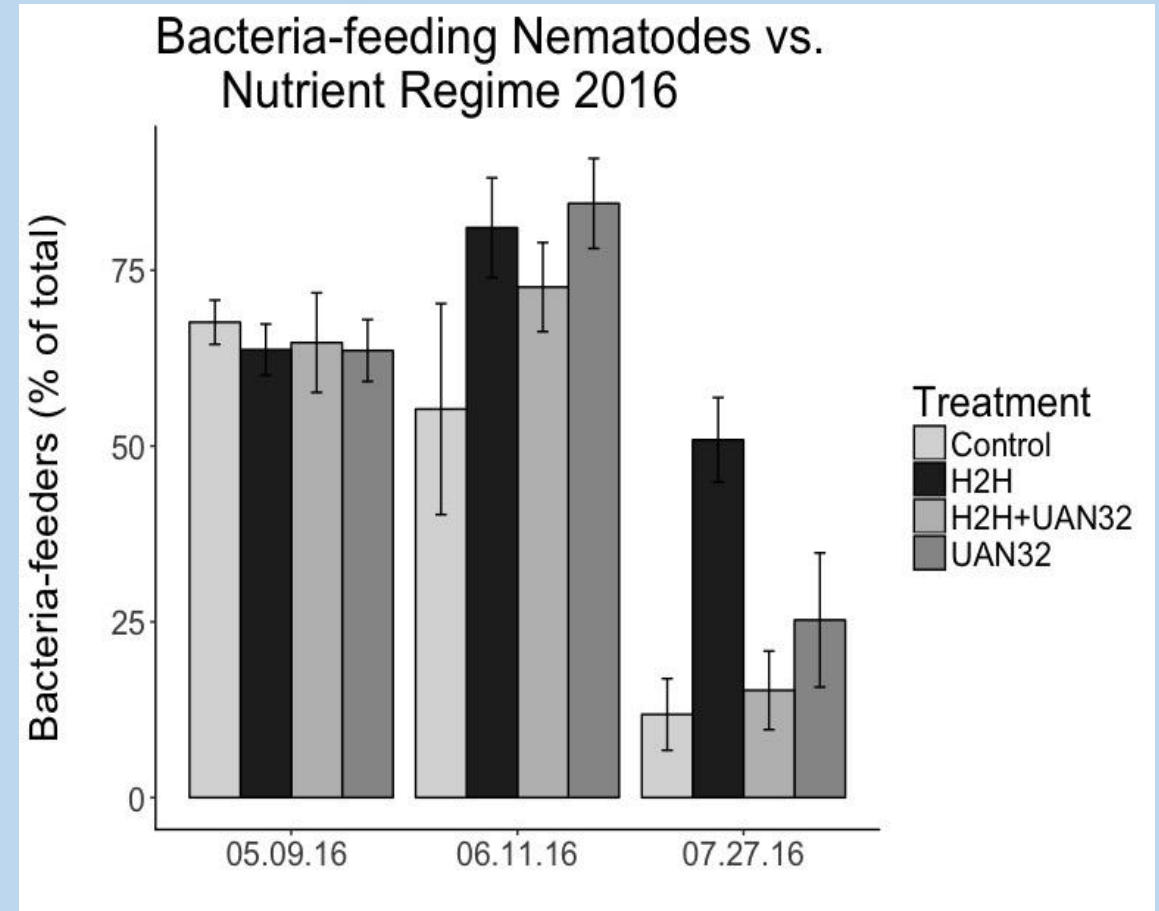
# Shifts in nematode community composition





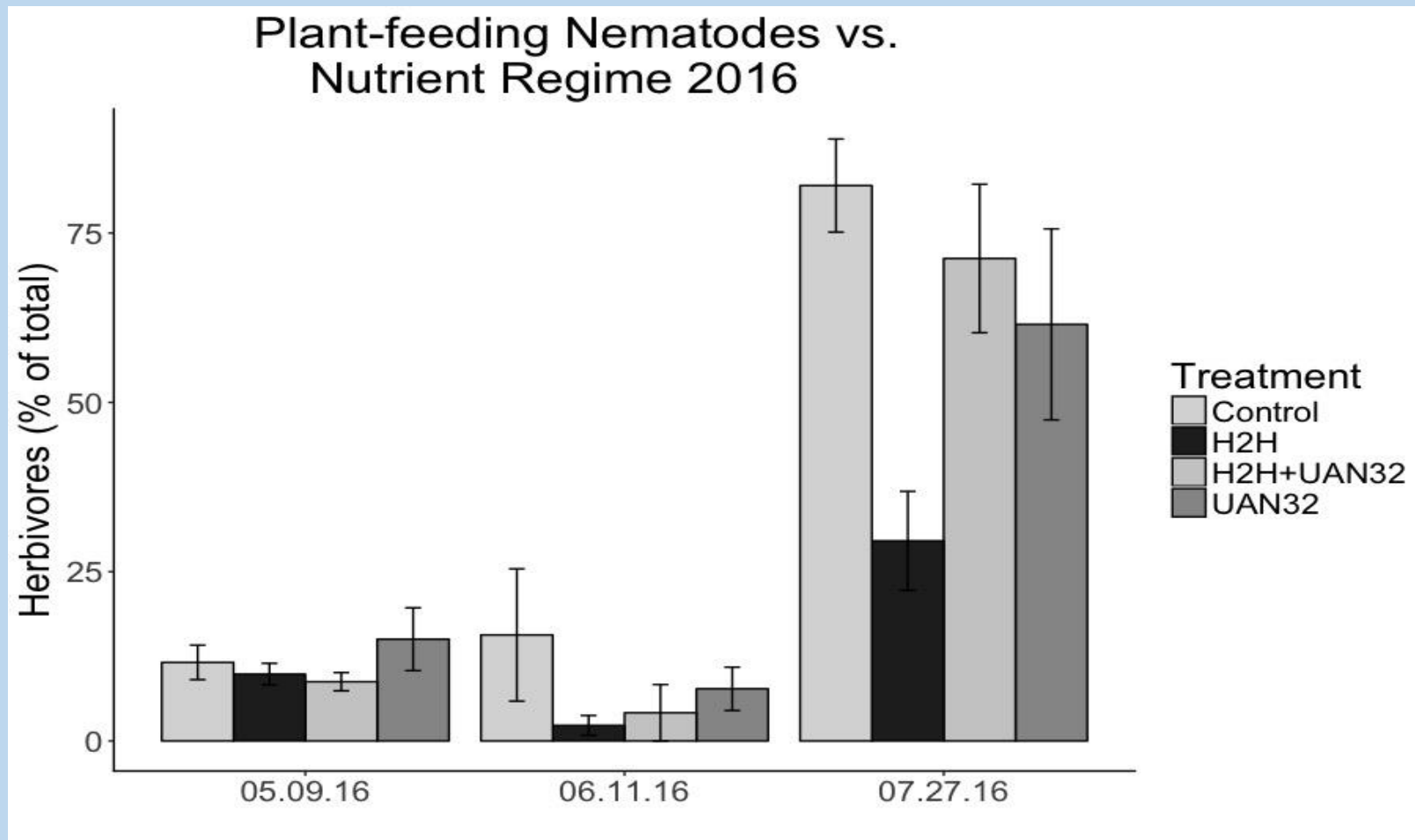
# 2016- Beneficial bacterial feeding nematodes

- Higher percent bacterial feeders with H2H compared to controls (P=0.03).
- Mainly *Acrobeloides* and *Cephalobus*.
- Responsive to inputs of high quality organic matter such as plant root exudates and detritus in the soil.



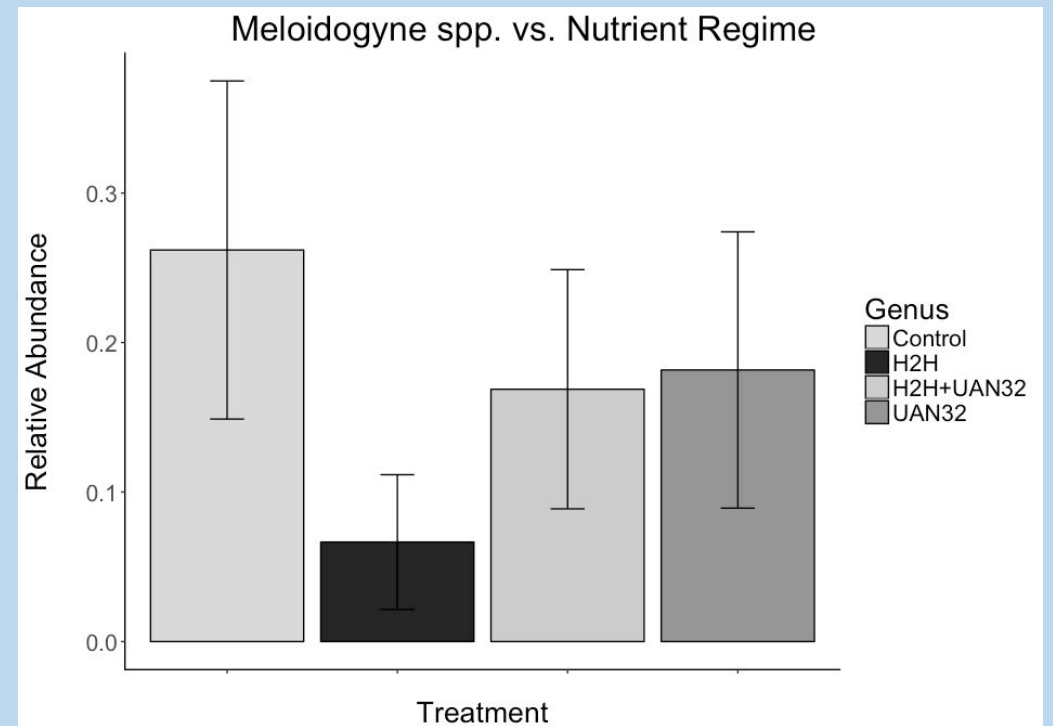
# Plant parasitic nematodes

- Decreased abundance of herbivores in H2H treated plots compared to controls ( $P=0.03$ ).



# Root knot abundance/damage

- Root knot nematodes made up the majority of the plant parasites.
- Root knot abundance and galling decreased with H2H.



# Next steps...

- *How can an amendment increase some nematodes but decrease others?*
- Food web mediated nematicidal effects
- Organic amendments alter soil communities to become more antagonistic to nematodes
  - Inputs stimulate microbial biomass and bacterial feeding nematodes
  - Bacterial feeding nematodes transport “helpful” microbes through soil
  - These suppress pests and promote root growth

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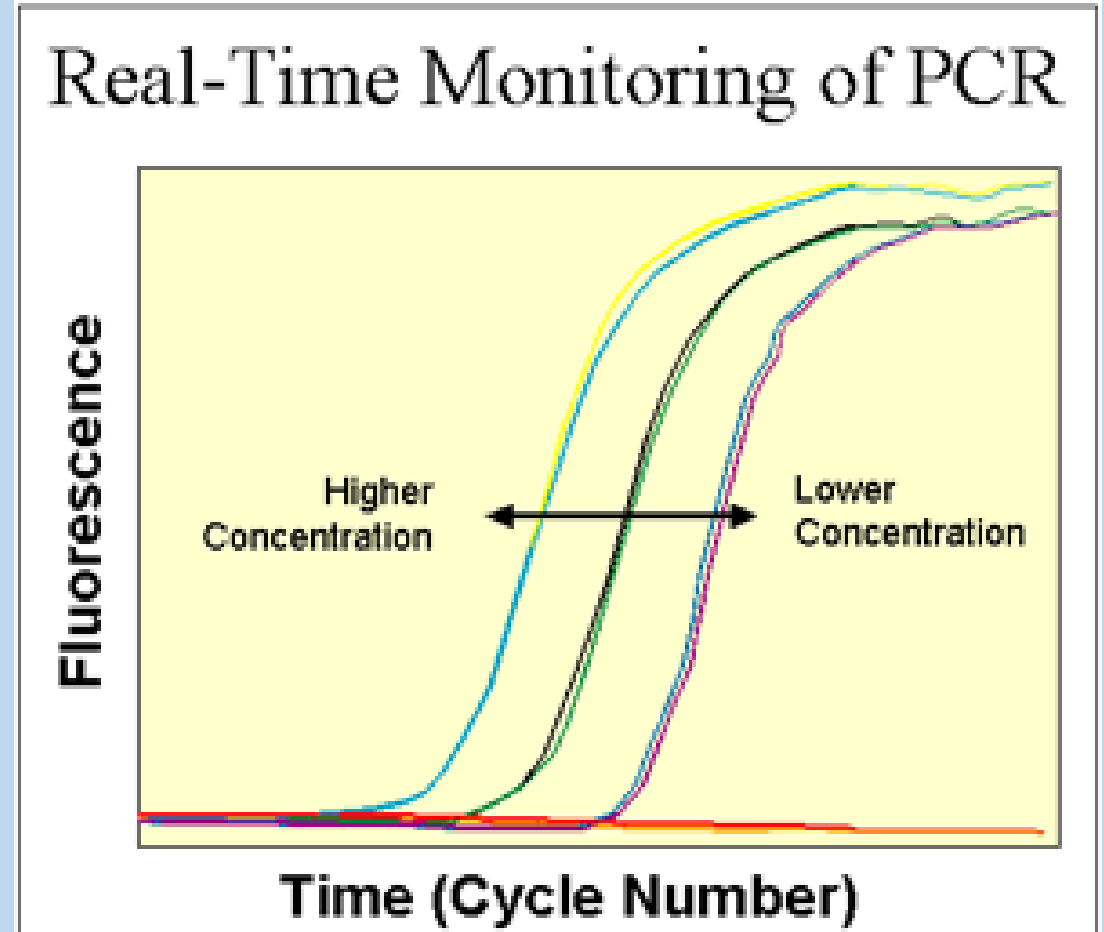
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# Nematode diagnostics using real time PCR (qPCR)

- Simultaneously quantifies and identifies nematodes
- Compares the intensity of the amplified signal to a standard curve calculated from known densities.
- Rapid and inexpensive pest identification and quantification.



# Comparison of traditional and molecular methods

## **Current method**

- Nematode extraction from soil
- Counting and identification (requires training)
- Subjective
- Inefficiencies in soil extraction

## **Real time (qPCR) method**

- Nematode extraction from soil
- Nematode DNA extracted and target sequence amplified by qPCR
- Concentration compared to standard curve
- Objective
- Sensitive-Can detect as little as 1/10 nematode!

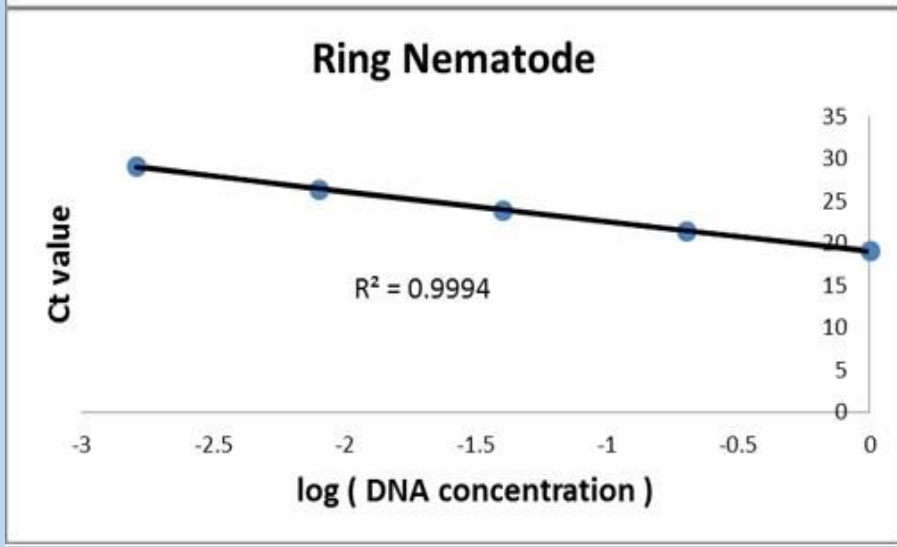
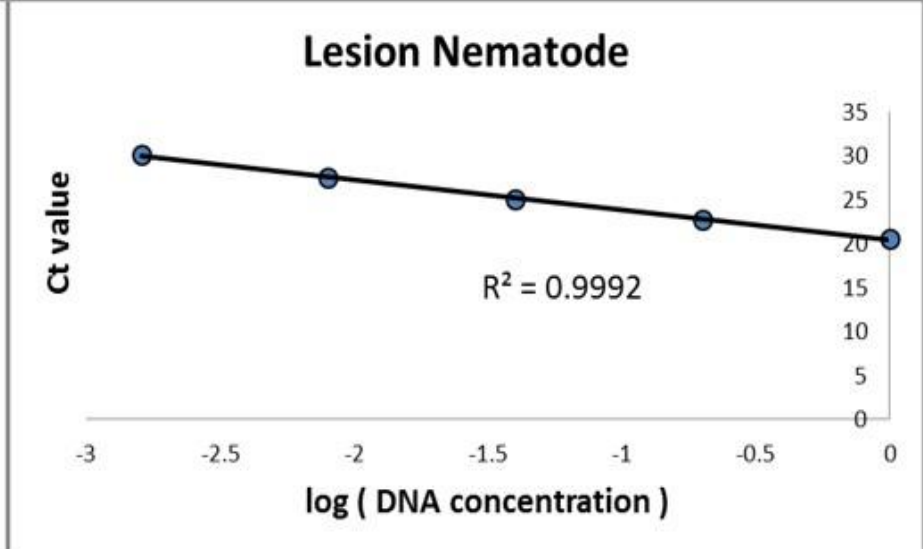
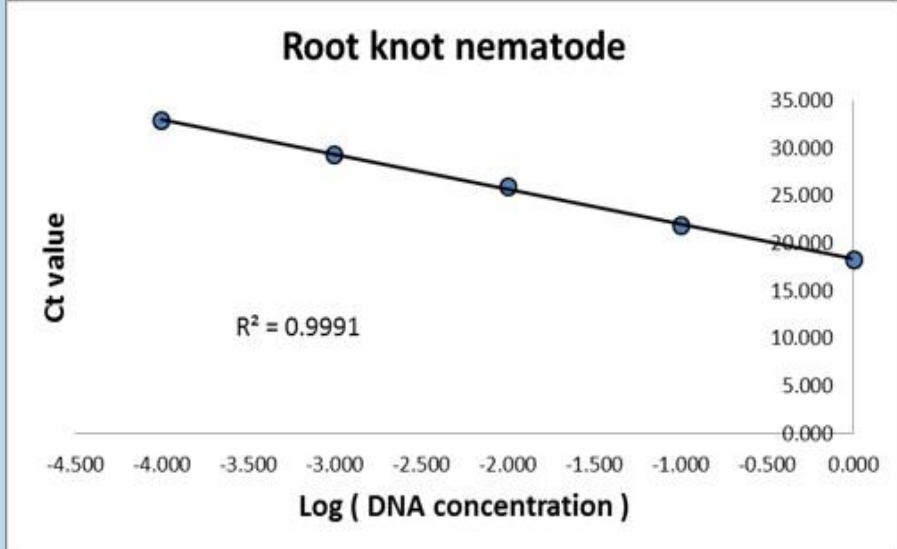


Figure 1. Standard curves generated using qPCR on serially diluted samples of 200 nematodes for each species. The x axis is the log of the %DNA concentration (total nematodes/200) and the y axis is the Ct value, or qPCR signal.



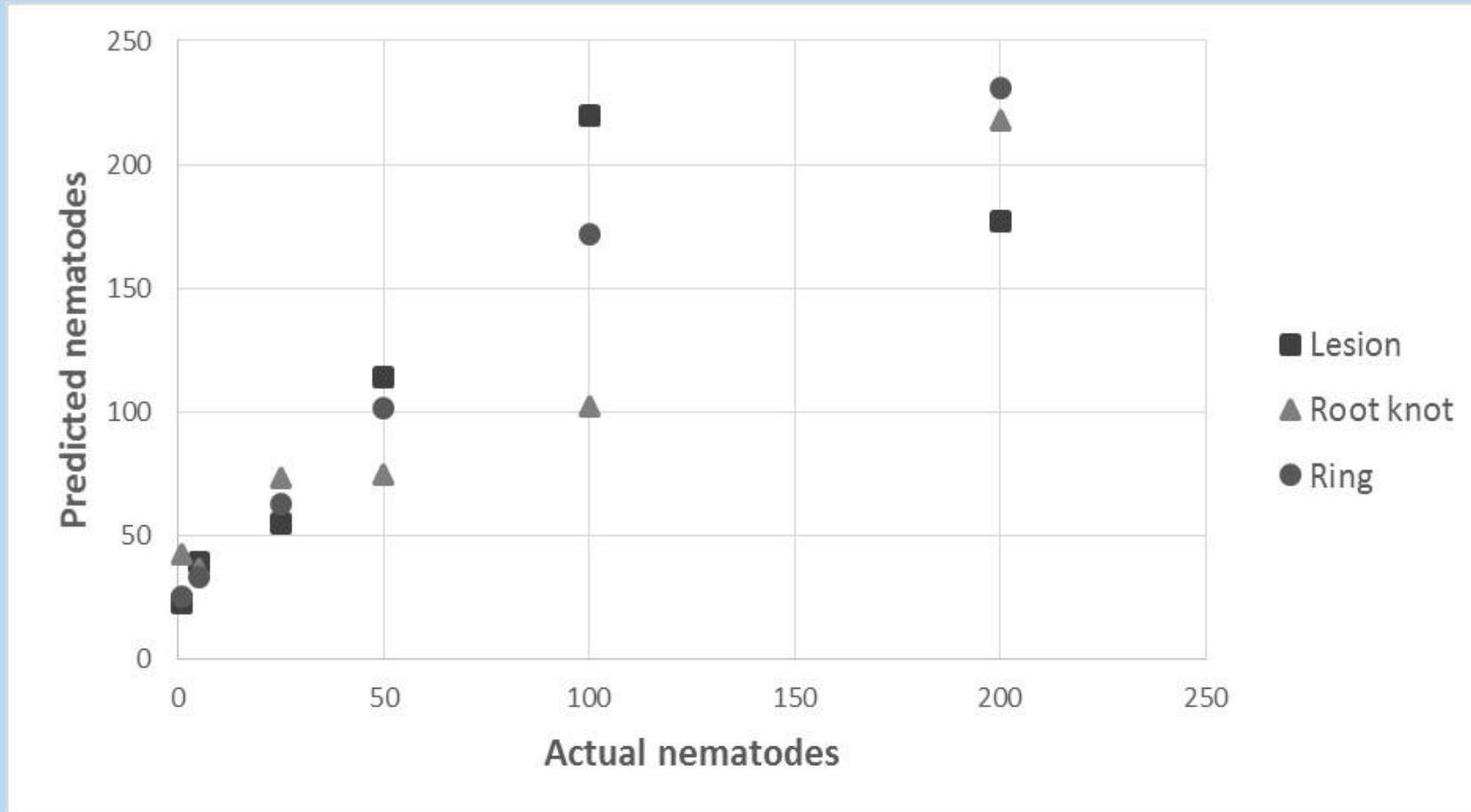


Figure 2. Relationship between known nematodes added to solution and nematode numbers predicted using the currently developed qPCR assay for three pest species of nematodes found in almonds, lesion, root knot and ring.

# Rapid detection and damage threshold analysis-decision making tools for nematode management in carrots

- Develop a qPCR panel for root knot nematodes *M. incognita*, *M. hapla*, *M. javanica*, and *M. arenaria*.
- In field tests comparing the qPCR assay to traditional methods of microscopic quantification. Other studies have found an 85% increase in assay sensitivity.
- Use qPCR derived nematode densities to model carrot damage thresholds in greenhouse trials.



# Thank you!

## Questions?

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