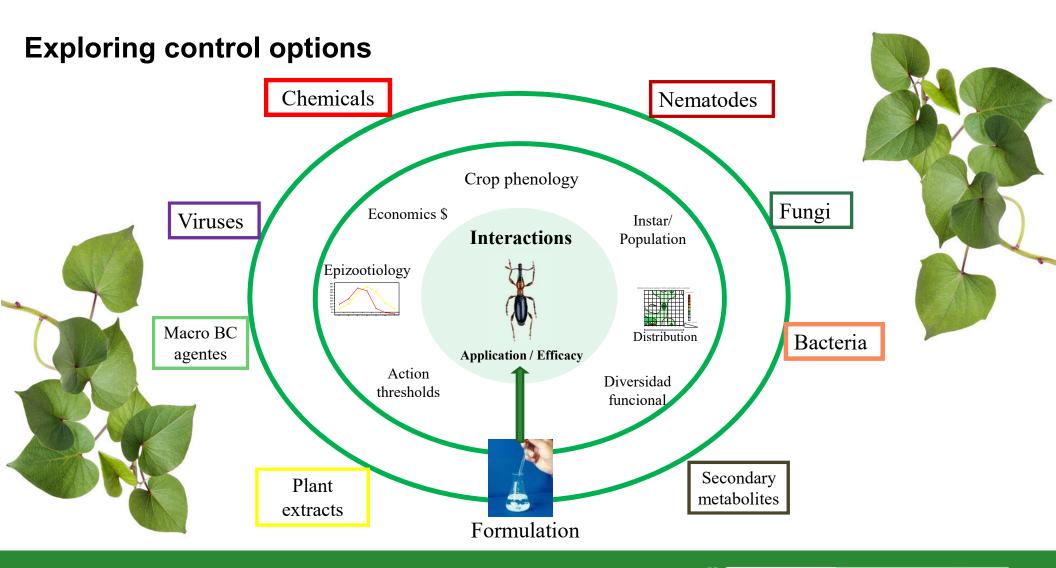
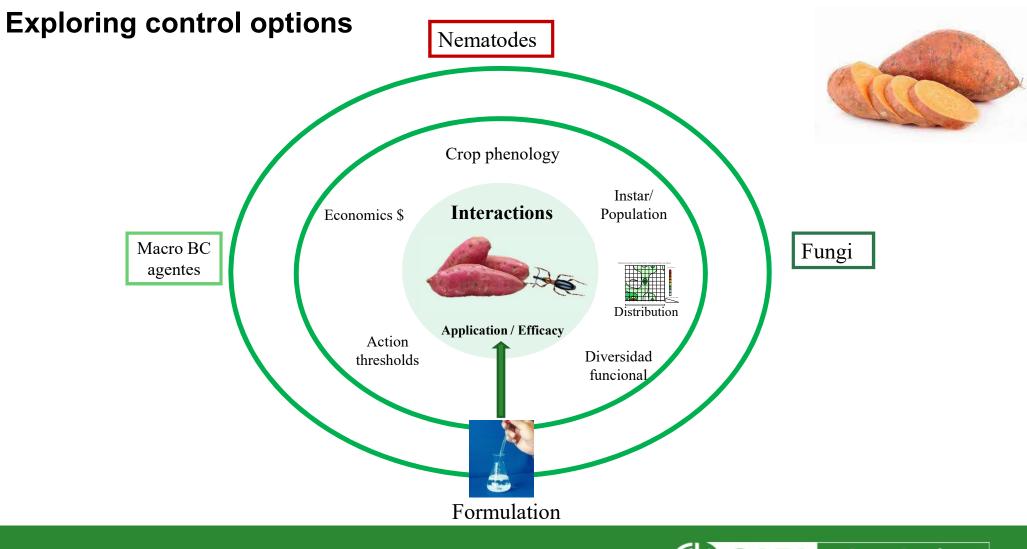
Biological control of sweet potato weevils: Current status and perspectives for the Caribbean

Eduardo Hidalgo - CABI e.hidalgo@cabi.org October 2021

(D) CABI plantwise



(b) CABI plantwise



(b) CABI plantwise

Quick facts about sweet potato weevils (SPW) and their biocontrol agents

- There are 2 SPWs reported in the Caribbean: Cylas formicarius and Euscepes postfasciatus
- Only a few biocontrol agents have been reported in the world:
 - Parasitoids
 - Predators
 - Entomopathogenic fungi
 - Entomopathogenic nematodes



Photo: Cook Island-Bishop Museum

Euscepes postfasciatus



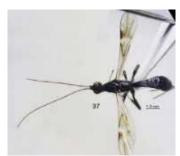
Cylas formicarius



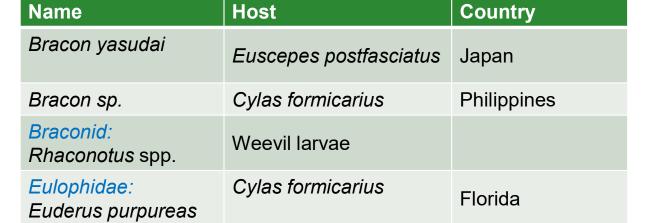
Parasitoids



Bracon yasudai



Rhaconotus sp.



- N
 - Euderus sp.



- Not found in all the countries
- Difficult mass production
- Sample for local acurrence in the Caribbean
- If found: keep conditions for natural augmentation



Predators

Predatory ants	Hosts	Country
Tetramorium guineense	Conoraliatia produtora	Cuba
Pheidole megacephala	Generalistic predators	
Iridomyrmex humilis	Generalistic predators Damaging for humans and livestock	Argentina



Tetramorium guineense



Pheidole megacephala



Field manipulation of ant nests



Nest of Tetramorium guineense in banana rotting stems and leaves. Source: Cisneros F. and alcazar, J., 2001

- Highly effective / Non especific predators
- Attack maily adult weevil
- Weed control is needed to reduce pray populations (ea. Aphids)
- Pheidole has show a negative interaction with mealybugs in pinaple



Field manipulation of ant nests

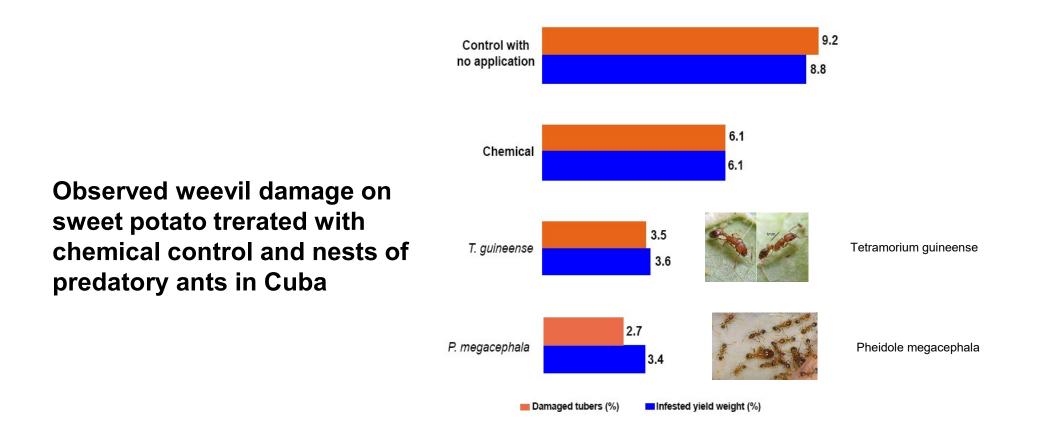




Banana pseudostem trap used in Cuba for collecting ant colonies. Source: Cisneros F. and alcazar, J., 2001

- Ants colonies can be captured with traps (3 to 12 days until the queen moves into the trap)
- Place the ant nests in the sweetpotato plantation 30 days after sowing (eary in the morning)
- Previous irrigation may be necessary (ants need moist soil)
- In Cuba: 100 nests /ha protected under the folliage

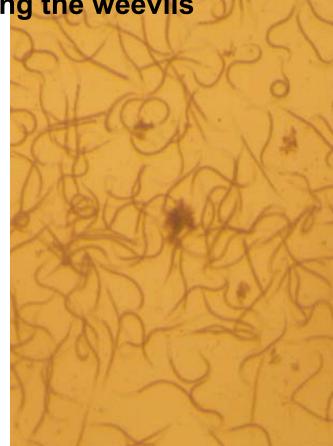






Using entomopathogenic nematodes for controlling the weevils

- Families: Steinernematidae and Heterorhabditidae
- Carry and introduce symbiotic bacteria (*Xenorhabdus* and *Photorhabdus*).
- Its host range includes both weevil species
- They can be grown on a large scale
- They can kill in 48 hours
- Can be stored and applied with conventional methods



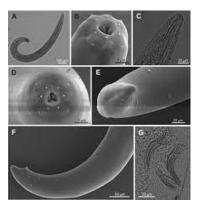
(b) CABI plantwise

Entomopathogenic Nematod species reported

Species	Effectivity	Country
Heterorhabditis karii		Kenya
Heterorhabditis indica	Higher effectivity (Larvae)	Kenya
Heterorhabditis bacteriophora		
Steinernema carpocapsae	(les effective) Larvae	Florida
Heterorhabditis sp. H1-24	Not entioned (Adults)	Cuba



Heterorhabditis



Steinernema



Are nematodes a feasible option in the Caribbean countries?





Industrial production of nematodes (Liquid culture)

Gaugler and Han (2002) reported commercial-scale (c.10,000-l bioreactors) production costs of US\$31 for *S. carpocapsae* and US\$42 for *H. bacteriophora* per hectare (2.5×109 nematodes/ha).

costs to end-users remain greater than the alternative pest management tactic in most markets. They concluded that growers seemed unlikely to pay a premium to use nematodes when there were familiar, easy-to-use, low-cost alternatives.



10k liter biorreactor: USD 100000.00



Industrialized production of nematodes Distribution of costs

The high product cost of EPNs is due to the relatively expensive and **lengthy processes** involved in their **mass production**, **formulation**, **storage** and **transport**.

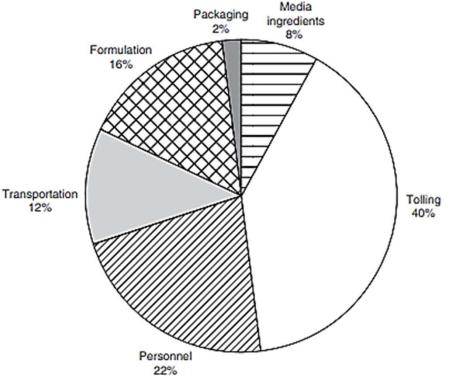


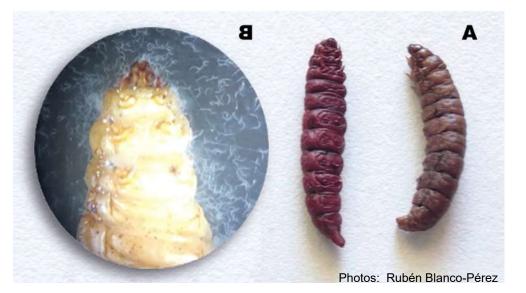
Fig. 14.2. Production cost breakdown for *Heterorhabditis bacteriophora* (HbNJ strain) for mass culture at a toll or contract manufacturer using a 3000-litre bioreactor (S. Franceschini, Italy, 2000, unpublished data).



In vivo production method

- in vivo Galleria process, yields between 0.5 e5and 4 e5 IJs/larva
- The *in vivo* process is regarded of lacking economy of scale
- Lack of improved quality while increasing scale, the keep small production?
- *in vivo* nematode production is sensitive to biological variations

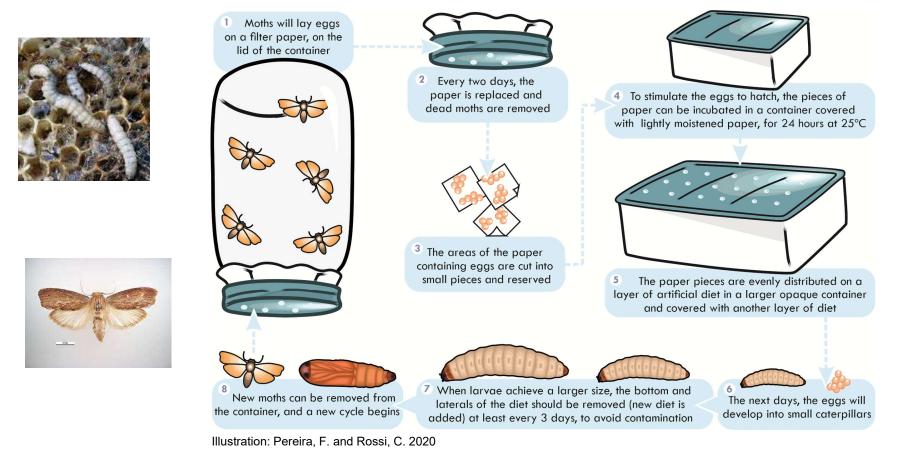
(Grewal et al.,2005).



Galleria mellonella larvae inoculated with nemoatodes



Rearing method for Galleria mellonella



(b) CABI plantwise

White trap technique for yielding nematodes from parasitised larvae

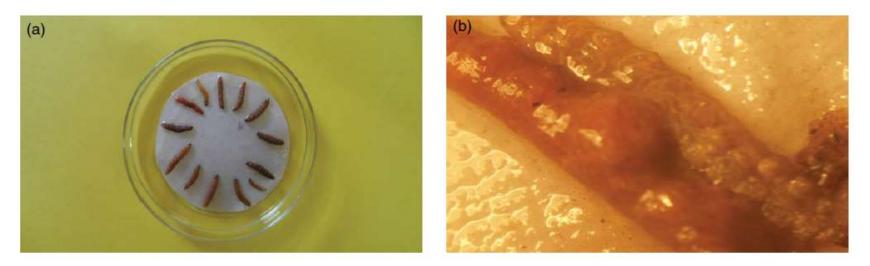


Fig. 13.3. (a) White trap method for harvesting EPNs emerged from insect cadavers; (b) infected juveniles (IJs) inside the bodies of EPN-infected insect larvae.



Size of the host and length of the cycle



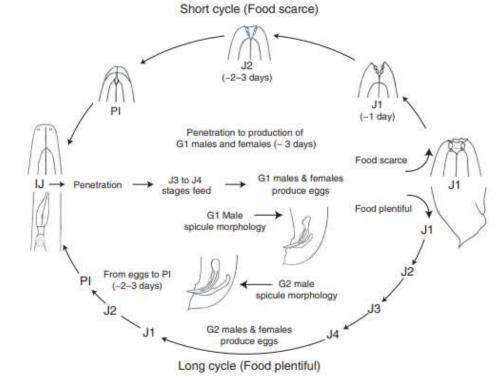
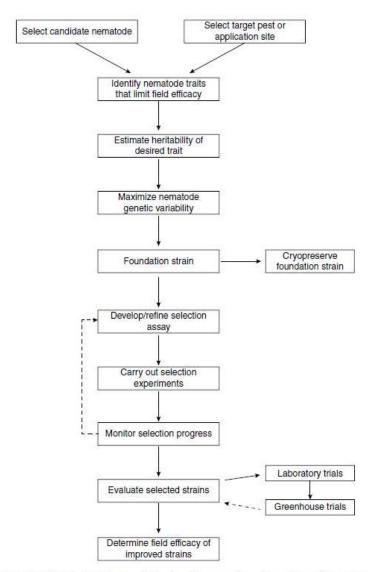




Photo by : Chigurupati Sai Prasanth





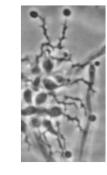
Planning a selection proces for nematode strains

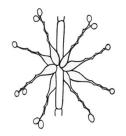
Fig. 26.1. A schematic illustration to design a selection breeding course for entomopathogenic nematodes. (From Gaugler et al., 1989; Burnell, 2002.)

(b) CABI plantwise

Entomopathogenic fungi

Species	Hosts	Country
Beauveria bassiana	<i>Cylas</i> and <i>Euscepes</i> (adults)	China
Metarhizium anisopliae		up to 43% controlof C. puncticollis applying 28 days after planting

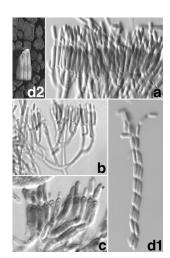




Beauveria bassiana



Photo from jorgeiv93.wordpress.com



Metarhizium anisopliae



Photo from Dotaona R. et al. 2017



Entomopathogenic fungi

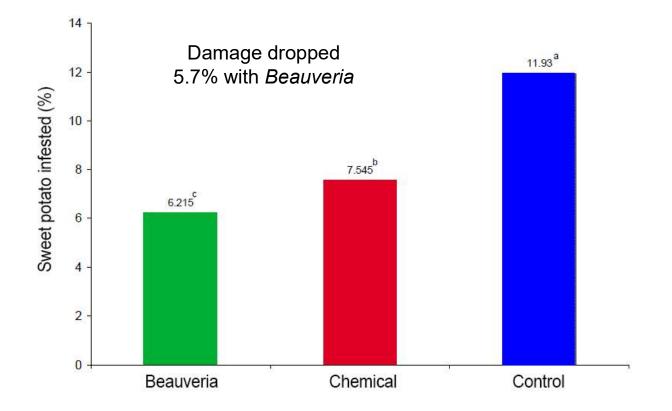
- Beauveria bassiana and Metarhizium anisopliae have shown effective against adult weevils
- Metarhizium produces more conidia at field level
- 48 to 72 hours for killing
- Application methods:
 - Plant dipping (5% conc.)
 - Spraying: 15 days after sowing (10e12 conidia/ha every 7-10 days until establishment)



Photos frm Dotaona R. et al. 2017

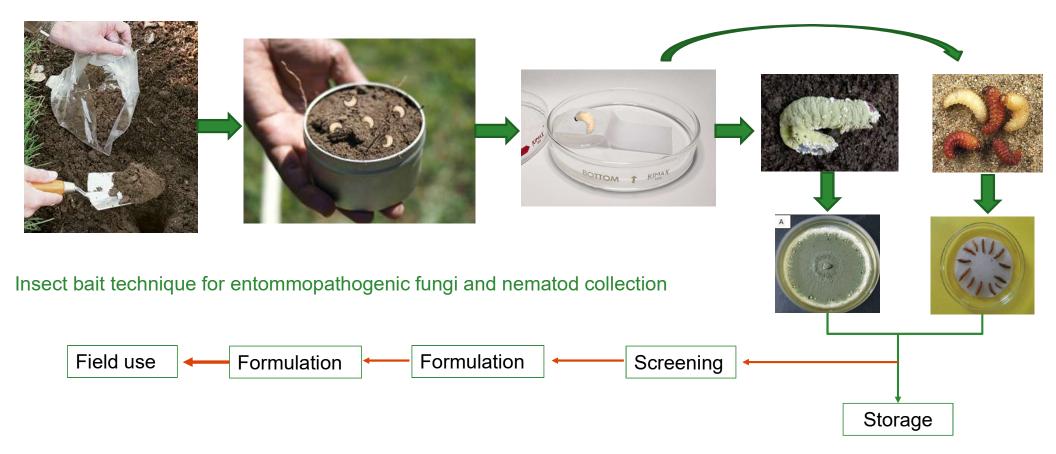


Observed effect of the application of B.bassiana for the control of SPW in Cuba

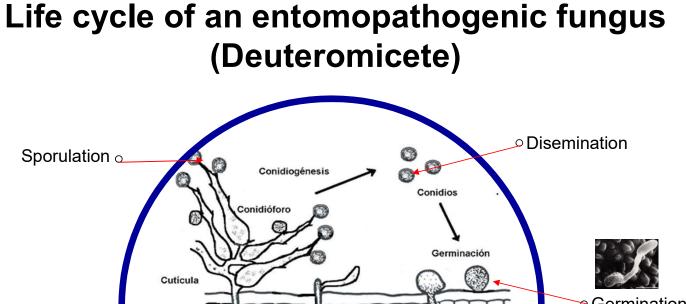


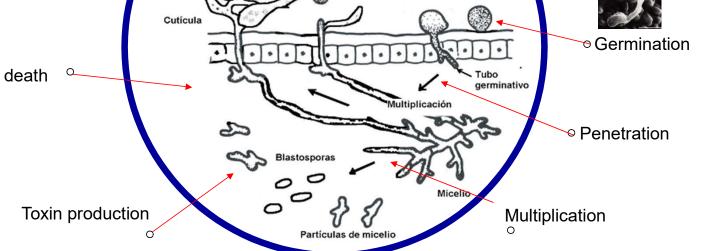


Generating local strain collections of entomopathogenic fungi and nematodes





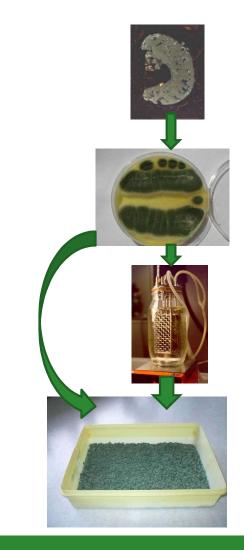






How to produce entomopathogenic fungi?

- 1. Isolate from insect or directly from soil using bait insects or media with antibiotics
- 2. Use monosporic isolates to avoid contaminants and assure genetic homogeneity
- Inoculate the solid substrate (rice) with a conidial suspension of aprox. 1x10⁶ con/ml, 10ml per 100g of substrate in polypropylene bags.
- 4. Incubate at 24-26 °C for 10 to 15 days
- 5. For large production, multiply the inoculum with a 3 day liquid fermentation in nutrient broth







- Consider the use of the 3 main bicontrol agents identified as effective (predatory ants, entomopathogenic fungi and nematodes)
- Predatory ants:

-

- Check the existence of *Pheidole* and *Tetramorium* species in the islands
- Validate the pseudostem trapping method for collecting ant colonies





Photo by : Chigurupati Sai Prasanth

- Entomopathogenic nematodes:

- Generate a strain bank isolated from representative soils/environments of the country
- \odot Screen against weevil larvae and adults
- Consider Small scale production at community level
- Analyse the economics and feasibility for a large-scale production system for the region





Entomopathogenic fungi

- Generate a strain bank isolated from representative soils/environments of the country
- Screen against weevil eggs, larvae, pupae and adults
- Consider Small scale production at community level
- Analyse the economics and feasibility for a large-scale production system for the region



- Parasitoids



- \odot Run field samplings for parasitoid presence
- If found:
 - \circ Identify
 - \odot Run studies of efficacy
 - \odot Procure adequate environment for augmentation





CABI is an international intergovernmental organisation, and we gratefully acknowledge the core financial support from our member countries and lead agencies including:

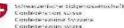


Ministry of Agriculture, People's Republic of China









Swiss Agency for Development

