

New opportunities for the integration of microorganisms into biological pest control systems in greenhouse crops

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Abstract Biological pest control with mass-produced arthropod natural enemies is well developed in greenhouse crops and has often resulted in the evolution of complex ecosystems with persistent populations of multiple arthropod natural enemy species. However, there are cases where arthropod natural enemies are either not effective enough, not available, or their use is rather costly. For these reasons, biological control based on microorganisms, also referred to as ‘microbials’, represents a complementary strategy for further development. Although commercially available microbials have been around for quite some time,

research on and the applied use of combinations of arthropod natural enemies and microbials have remained relatively under explored. Here, we review current uses of entomopathogenic fungi, bacteria and viruses, and their possible direct and indirect effects on arthropod natural enemies in European greenhouses. We discuss how microbials might be combined with arthropod natural enemies in the light of new methodologies and technologies such as conservation biological control, greenhouse climate management, and formulation and delivery. Furthermore, we explore the possibilities of using other microorganisms for biological control, such as endophytes, and the need to understand the effect of insect-associated microorganisms, or symbionts, on the success of biological control. Finally, we suggest future research directions to optimize the combined use of microbials and arthropod natural enemies in greenhouse production.

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Key message

- The application of microbials for pest control in greenhouse crops should be integrated with the use of arthropod natural enemies. Here we review the current uses of entomopathogenic fungi, bacteria and viruses, and their possible direct and indirect effects on arthropod natural enemies.
- New approaches in the use of conservation biological control, greenhouse climate management, formulation, delivery and endophytic microorganisms could increase the various ways in which microbials can interact with arthropod natural enemies, and these interactions can be both positive and negative for pest control.

- A better understanding of these interactions offers new opportunities to optimize and further develop biological pest control.

Introduction

Biological control of arthropod pests by arthropod natural enemies has been used successfully in greenhouse crops for decades (Pilkington et al. 2010). The protected environment of high-value greenhouse crops is particularly suitable for the effective functioning of commercially produced natural enemies and, globally, the majority of arthropod natural enemy species sold are used for augmentation in greenhouse crops (van Lenteren 2012). However, despite this success, there are still many cases where arthropod natural enemies are not used due to high costs or low efficacy.

Biological control agents based on entomopathogenic microorganisms (viruses, bacteria, fungi, etc.), also referred to as microbials, are often promoted as an alternative or back-up treatment when arthropod natural enemies are unavailable or not sufficiently effective (Chandler et al. 2011). The application of entomopathogens must be compatible with the arthropod natural enemies that are used in the same biological control system, and side effects of entomopathogens have therefore been studied extensively (Roy and Pell 2000). However, recent studies are increasingly exploring the wider properties of microorganisms, which suggest new opportunities for their use in biological control systems (Lacey et al. 2015). For example, several entomopathogenic fungi can also colonize plant tissues as endophytes and affect pests systemically via the plant (Vega et al. 2009). Furthermore, microorganisms that are symbionts of pests can influence the successes of both arthropod natural enemies and entomopathogens and therefore biological control (Zindel et al. 2011).

In this review, we discuss the need to use microbials and endophytes for pest control on the most important greenhouse crops. Greenhouse crops offer unique opportunities to design and optimize ecosystems through releases of arthropod natural enemies and by manipulating the greenhouse climate. We believe it is important to consider how new applications of microbials fit into such a designed ecosystem. We particularly address the means by which microbials and endophytes can be used to support and enhance biological control by arthropod natural enemies in different greenhouse cropping systems and discuss new developments, knowledge gaps and future prospects for the use of microbials in greenhouse crops. When discussing registered microbial products, we have focussed on

entomopathogenic baculoviruses, bacteria and fungi that are currently registered for use in Europe. These groups are often considered as examples of biopesticides (Glare et al. 2012), alongside natural compounds and minerals. Here we will only consider living microorganisms, as this category of biopesticides requires application approaches that differ substantially from chemical pesticides and other nonliving biopesticides. Entomopathogenic nematodes are often also considered as microbials by the biocontrol industry, but despite their obvious importance, they will also be excluded from this review as, strictly speaking, they are not microorganisms but animals that use associated bacteria to kill their hosts (Kaya and Gaugler 1993). The majority of commercially available microbial products are based on the species *Bacillus thuringiensis* Berliner (Bt), but as this species has already been extensively reviewed (Sanahuja et al. 2011; Vachon et al. 2012), it will not be discussed in great detail here.

Current status of microbials used for pest control in greenhouse crops

In Europe in 2010, the estimated sales of microbials based on entomopathogens, such as bacteria, viruses and fungi, was 42 million Euro, of which the majority (58 %) could be assigned to *B. thuringiensis* (Glare et al. 2012). Although the market for microbials is expected to increase substantially (Glare et al. 2012), the number of officially registered products in Europe remains limited (Table 1).

Microbials based on subspecies of *B. thuringiensis* are registered for specific insect pests; the product based on *B. firmus* is only registered for control of nematode pests (Table 1). *Bacillus thuringiensis* forms spores that contain crystals, predominantly comprising one or more Cry and/or Cyt proteins (also known as delta-endotoxins) that lyse gut cells when consumed by susceptible insects (Gill et al. 1992). Bacterial insecticides have to be consumed in order to confer control of the pest. After ingestion, the insect gut becomes paralyzed due to the action of bacterial toxins, feeding stops and eventually the pest dies (Vachon et al. 2012). There are currently three Bt strains approved in the EU (Table 1).

The entomopathogenic fungi are a diverse assemblage of species with one thing in common: they infect and cause disease in insects and other arthropods. The majority are found within two groups: the order Hypocreales within the phylum Ascomycota, and the phylum Entomophthoromycota (Blackwell 2010; Hibbett et al. 2007; Humber 2012). In contrast with other microorganisms, entomopathogenic fungi infect their hosts by directly breaching the cuticle to enter the insect haemocoel. Their ability to invade without

Table 1 Registered microbials for greenhouse crops in Europe (Gwynn 2014)

Classification/species	Isolate/strain	Commercial name	Target pests
Bacteria			
<i>Bacillus thuringiensis</i> subsp. <i>aizawai</i>	ABTS-1857	XenTari, Turex	Caterpillars
<i>B. thuringiensis</i> subsp. <i>israelensis</i> (serotype H-14)	AM65-52	Vectobac (Gnatrol)	Fungus gnats
<i>B. thuringiensis</i> subsp. <i>kurstaki</i>	PB 54	Belthirul	Caterpillars
<i>Bacillus firmus</i> Werner	I-1582	Bacillus firmus I-1582 WP5	Nematodes
Fungi			
<i>Beauveria bassiana</i> (Balsamo) Vuillemin	ATCC 74040, GHA	Naturalis, Botanigard	Primarily whiteflies and thrips
<i>Isaria fumosorosea</i> Wize (formerly <i>Paecilomyces fumosoroseus</i> Wize)	Apopka 97, FE9901	PreFeRal Nofly	Primarily whiteflies
<i>Lecanicillium muscarium</i> Petch (formerly <i>Verticillium lecanii</i> Zimmerman)	Ve-6	Mycotal	Primarily whiteflies
<i>Metarhizium brunneum</i> Petch (formerly <i>Metarhizium anisopliae</i> Metschnikoff)	Bipesco 5, F52	Taerain Met52EC	Primarily whiteflies and thrips
<i>Purpureocillium lilacinus</i> (Thom) Samson (formerly <i>Paecilomyces lilacinus</i>)	251	BioAct	Root-knot nematodes
Viruses			
<i>Helicoverpa armigera</i> nuclear polyhedrosis virus	HearNPV	Helicovex	<i>Helicoverpa armigera</i>
<i>Spodoptera exigua</i> nuclear polyhedrosis virus	Florida isolate (SeNPV-F1)	Spod-X	<i>Spodoptera exigua</i>
<i>Spodoptera littoralis</i> nuclear polyhedrosis virus	SpliNPV	Littovir	<i>Spodoptera littoralis</i>

the requirement for ingestion is a great advantage for infecting phloem-feeding insects, such as aphids and whiteflies, which do not ingest microorganisms on the leaf surface.

Beauveria bassiana sensu lato, *Isaria fumosorosea* (formerly *Paecilomyces fumosoroseus*), *Metarhizium anisopliae* sensu lato. (Bischoff et al. 2009) and *Lecanicillium* (formerly *Verticillium*) species have all been reported to be effective, when sprayed in suspension, against thrips, aphids, whiteflies and weevils in greenhouse crops (de Faria and Wraight 2007; Khan et al. 2012; Skinner et al. 2012). Currently, there are five products based on entomopathogenic fungi registered in the EU, all of them based on species from the Hypocreales (Table 1). Entomopathogenic fungi in the order Entomophthorales are known for their ability to cause dramatic epizootics that rapidly reduce host populations (Pell et al. 2010). This attribute, which is more rarely seen in species from the Hypocreales, means they have the potential to be more effective biological control agents than commercially available microbials from the order Hypocreales. However, a major constraint to their augmentation as biological control agents relates to difficulties in their in vitro mass production, storage and formulation (Pell et al. 2001).

This is undoubtedly the reason why the biocontrol industry has been unable to develop them as commercial microbials for augmentation (Ravensberg 2011). However, as explored further in this review, the possibility of integrating them through conservation biological control approaches seems promising.

Although a number of different types of virus infect pest arthropods, commercial microbial products are predominantly based on just one virus family, the Baculoviridae. Three baculovirus species are registered as biological control agents of lepidopteran pests of greenhouses in the EU (Table 1). Baculoviruses are generally very host specific, infecting only one or a few closely related insect species. However, two extreme examples in the context of contrasting specificity are *Spodoptera exigua* multiple nuclear polyhedrosis virus (SeMNPV), which infects only one species, *Spodoptera exigua* Hubner (the beet armyworm), and *Autographa californica* MNPV, which infects species from more than 15 families of Lepidoptera (Cory and Myers 2003). High specificity for the host makes baculoviruses good candidates for biological control, as their application does not directly affect non-target insect species.

Combining microorganisms with arthropod natural enemies

Use of entomopathogenic microorganisms and endophytes in greenhouse crops could have various direct and indirect effects on existing biological control systems based on arthropod natural enemies, potentially leading to both positive and negative outcomes for overall pest control (Fig. 1). Similarly, the outcome of such interactions can also be influenced by symbionts in various ways. Positive interactions represent an opportunity to enhance the efficacy of existing biological control systems.

Microbials as a correction tool

Microbials are traditionally seen as an alternative or back-up treatment when arthropod natural enemies are unavailable or insufficiently effective (Chandler et al. 2011). For some pests, effective arthropod natural enemies are simply

not available or those that are available are considered to be too expensive and hence hardly used (Table 2). Microbials could be a sustainable alternative control method in these cases. For other pests, highly effective arthropod natural enemies are commercially available, but are not effective on all crops; many natural enemies maintain a close relationship with specific plants because of their plant feeding habits or oviposition requirements (Messelink et al. 2014). For instance, western flower thrips (*Frankliniella occidentalis* Pergande), can be controlled effectively by phytoseiid predatory mites in sweet pepper, which provides pollen and nectar, but not in many ornamental plants that lack these supplemental food resources (Messelink et al. 2014). In such scenarios, the use of microbials has potential as a complementary measure in biological control programmes, as long as any potentially negative direct and indirect effects of these microorganisms are considered.

Direct effects of microbials on arthropod natural enemies

Augmentative biological control is increasingly being combined with methods to conserve both the released and the naturally occurring species of arthropod natural enemies present (Messelink et al. 2014). Conservation approaches can result in the permanent presence of several interacting species of pests and natural enemies and such food webs require an ecosystem approach to manage (Janssen et al. 1998; Messelink et al. 2012). Application of additional control agents should, ideally, complement or support these persistent communities of arthropod natural enemies. The species diversity and persistence of arthropod natural enemies used to control pests in greenhouse crops strongly depend on the type of crop. This is particularly true for those natural enemies that maintain close relationships with certain host plants because of their particular plant-feeding behaviours and oviposition requirements. The predatory mirid bugs *Macrolophus pygmaeus* Rambur and *Nesidiocoris tenuis* Reuter, for example, perform well on hairy plants such as tomato and aubergine and, once present, often persist throughout the entire cropping cycle. Therefore, it is desirable that any microbial used is compatible with these persistent arthropod natural enemies, in order to conserve their populations. The most important and persistent groups of arthropod natural enemies include not only generalist predators, but also specialist species that can remain for a long time when they have reached equilibrium dynamics with their prey/host (Table 3). Such specialists include whitefly parasitoids and some spider mite predators. Sometimes achieving a stable equilibrium is attempted through deliberate releases of the pest itself (e.g. the ‘pest-in-first’ method; (Markkula and Tiitanen

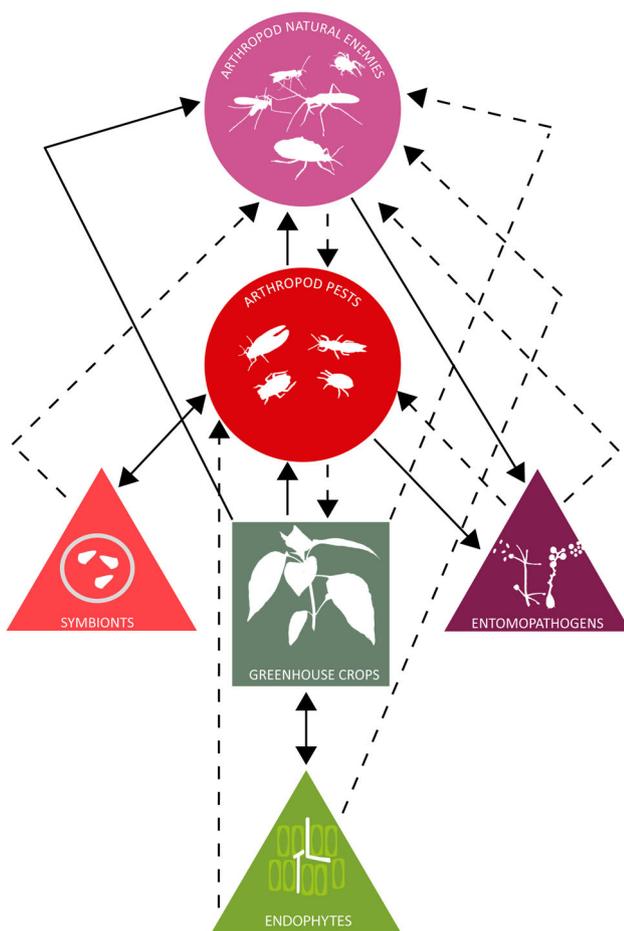


Fig. 1 Putative interactions between crops (*square*), arthropod pests and natural enemies (*circles*), and microorganisms (*triangles*). *Dashed arrows* indicate potential negative effects, while *solid arrows* indicate potential positive effects

Table 2 Key European greenhouse pests against which arthropod natural enemies are (i) not always effective, (ii) not used because they are considered as too expensive, (iii) are unavailable or (iv) are unknown (pers. obs. by the authors of this paper)

Common name	Scientific name	Crop
Insecta: Hemiptera		
Aphids	Several species, e.g. <i>Myzus persicae</i> Sulzer <i>Aulacorthum solani</i> Kaltenbach <i>Aphis gossypii</i> Glover <i>Macrosiphum euphorbiae</i> Thomas	Vegetables and ornamentals
Armoured scales	<i>Diaspis boisduvalii</i> Signoret <i>Aulacaspis rosae</i> Bouche	Ornamentals
Mealybugs	<i>Planococcus citri</i> Risso <i>Pseudococcus longispinus</i> Targioni-Tazzetti <i>Pseudococcus viburni</i> Signoret	Vegetables and ornamentals
Tarnished plant bug	<i>Lygus rugulipennis</i> Poppius	Aubergine
Common nettle bug	<i>Liocoris tripustulatus</i> Fabricius	Cucumber
Common green capsid	<i>Lygocoris pabulinus</i> Linnaeus	Sweet pepper
Whiteflies	<i>Trialeurodes vaporariorum</i> Westwood <i>Bemisia tabaci</i> Gennadius	Ornamentals
Insecta: Diptera		
Cabbage root fly	<i>Delia radicum</i> Linnaeus	Radish
<i>Lyprauta</i>	<i>Lyprauta chacoensis</i> Edwards <i>Lyprauta cambria</i> Chandler	<i>Phalaenopsis</i> spp.
Insecta: Lepidoptera		
Caterpillars	several species, e.g. <i>Chrysodeixis chalcites</i> Esper <i>Lacanobia oleracea</i> Linnaeus <i>Duponchelia fovealis</i> Zeller	Vegetables and ornamentals
Insecta: Thysanoptera		
Western flower thrips	<i>Frankliniella occidentalis</i>	Ornamentals
<i>Echinothrips</i>	<i>Echinothrips americanus</i> Morgan	Ornamentals
Acari: Prostigmata		
Tomato russet mite	<i>Aculops lycopersici</i> Masse	Tomato

The ranking is based on the Class and Order of species

1976). In such cases, it is important that the application of microbials against the same or other target pests does not disrupt the populations of arthropod natural enemies that increase resilience of cropping systems through their continuous presence.

Microbials could have potential negative effects on non-target organisms, although most are considered to be highly host-specific, such as the *B. thuringiensis* subspecies, viruses of particular species of Lepidoptera and fungi from the phylum Entomophthoromycota (not registered, but often occurring spontaneously in greenhouse crops, G.J Messelink & C. Tkaczuk, pers. obs.). In contrast, entomopathogenic fungi within the order Hypocreales (Ascomycota), although isolate and species dependent, can have wider host ranges than other pathogens and could potentially kill non-target arthropod natural enemies (Roy and Pell 2000). However, these potential side effects are

likely to vary significantly depending on the type of arthropod natural enemy. For example, greenhouse studies with predatory mirid bugs showed no negative effects of two commercial isolates of *B. bassiana* (GHA and ATCC 74040) on predator populations (Hamdi et al. 2011; Labbé et al. 2009). In contrast, densities of predatory *Orius* spp. were significantly reduced due to infection by the GHA isolate of *B. bassiana* (Shipp et al. 2012), although in other studies there were no or only weak side effects on *Orius* spp. predators (Hamdi et al. 2011; Pourian et al. 2011). Predatory mites generally, both in laboratory and greenhouse trials, lack vulnerability to commercial isolates of the entomopathogenic fungi *B. bassiana*, *M. brunneum* and *I. fumosorosae* (Ludwig and Oetting 2001; Vergel et al. 2011; Wu et al. 2014). In the laboratory, leafminer and whitefly parasitoids can be highly susceptible (30–70 %) to *B. bassiana* (Shipp et al. 2003), but in greenhouse trials

Table 3 Important arthropod natural enemies in greenhouse crops that are often continuously present (Heinz et al. 2004)

Type of natural enemy: common name	Type of natural enemy: Scientific name	Crop
Generalists		
Predatory mirid bugs	<i>Macrolophus pygmaeus</i> <i>Nesidiocoris tenuis</i>	Tomato, aubergine and sweet pepper
Predatory anthocorid bugs	<i>Orius</i> spp.	Sweet pepper
Phytoseiid mites	Several species, e.g. <i>Neoseiulus cucumeris</i> Oudemans and <i>Amblyseius swirskii</i> Athias-Henriot	Sweet pepper, cucumber, aubergine, cut flowers and potted plants
Specialists		
Whitefly parasitoids	<i>Encarsia formosa</i> Gahan, <i>Eretmocerus eremicus</i> Rose & Zolnerowich, <i>Eretmocerus mundus</i> Mercet	Sweet pepper, tomato, cucumber, aubergine, poinsettia, gerbera and roses
Leafminer parasitoids	<i>Diglyphus isaea</i> Walker, <i>Dacnusa sibirica</i> Telenga	Tomato and gerbera
Predatory mites	<i>Phytoseiulus persimilis</i> Athias-Henriot <i>Neoseiulus californicus</i> McGregor	Strawberry, sweet pepper, roses and chrysanthemum

only low levels of infection were observed in whitefly parasitoid populations (Labbé et al. 2009; Shipp et al. 2012). In a recent study, the parasitoid *Trybliographa rapae* Westwood was shown to be susceptible to *B. bassiana* and *M. brunneum*, but further experiments showed that it was also able to recognize and avoid *M. brunneum* suggesting these two biological control agents are compatible (Rännbäck et al. 2015). Similarly, a new study has shown that the mortality of different arthropods for the control of the western flower thrips ranged from 3 to 61 % when combined with entomopathogenic fungi in laboratory tests (Saito and Brownbridge 2016). However, their results also indicate that compatibility and overall increased effects are observed when both biological control agents are applied. Therefore, the majority of studies suggest that microbials are compatible with arthropod natural enemies, but caution should be practiced with the application of entomopathogenic fungi with broad host ranges.

Indirect effects of microbials on natural enemies

The application of microbials to crops can have unexpected effects on arthropod natural enemies through changes in behaviour of the pests, the arthropod natural enemies or through changes in pest densities or pest diversity (Roy and Pell 2000; Roy et al. 2006). This latter aspect is particularly relevant for generalist predators that feed on multiple pest species. Changes in the density of one pest species due to the application of a target-specific microbial can indirectly affect other pest species, simply because of the availability of food or because the predators perform better on mixed

diets of pests (Messelink et al. 2008, 2010; Muñoz-Cárdenas et al. 2014). Such predator-mediated effects should be taken into account when applying microbials against a specific pest species.

Other negative interactions can occur as a result of avoidance. Predatory bugs may, in some cases, avoid preying on infected prey or plants treated with microbials, which could increase their prey searching time and reduce predation rates (Labbé et al. 2009; Meyling and Pell 2006; Pourian et al. 2011). Several parasitoid species seem to be able to detect whether their host is infected by an entomopathogenic fungus, such as the whitefly parasitoid *Encarsia formosa* (Fransen and van lenteren 1993) and the aphid parasitoid *Aphelinus asychis* Walker (Mesquita and Lacey 2001). This is positive for the survival of the parasitoid's offspring, but at the same time may increase searching time and thereby reduce efficacy. Other parasitoid species are not able to detect infected living hosts, as was reported for *Aphidius ervi* Haliday and aphids infected by *Pandora neoaphidis* Remaudiere & Hennebert (Baverstock et al. 2005). This could have a tremendous negative effect on the efficacy of this parasitoid, as parasitoids developing in infected aphids will not develop into adults. Natural enemies may also predate on infected hosts, thereby reducing the efficacy of microbial applications (Pell et al. 1997; Roy et al. 2008). All these complex interactions may lead to nonadditive effects of combined treatments. Sometimes it may be possible to avoid this by adapting the timings of microbial applications and releases of arthropod natural enemies in order to minimize potential negative effects. For example, parasitoids might be more

effective when their release time after a microbial application is long enough to allow the separation of entomopathogen-infected hosts from uninfected hosts (Fransen and van lenteren 1993).

Positive, potentially synergistic, effects can also occur when the arthropod natural enemies enhance the dispersal of entomopathogens. Down et al. (2009) showed enhanced control of aphids when conidia of the entomopathogenic fungi *Lecanicillium longisporum* Zimmerman were disseminated by *Orius laevigatus* Fieber. Furthermore, the presence of coccinellid predators increased the proportion of aphids becoming infected by the fungal pathogen *P. neoaphidis* due to increased transmission (Roy et al. 2001; Wells et al. 2011). Some predators may further increase dispersal of conidia of aphid pathogens by inducing the production of winged morphs (Müller et al. 2001). Predators and parasitoids may also facilitate infection by microbes when their foraging activity increases movement of the pest, increasing the likelihood that pathogen propagules come into contact with their hosts (Roy and Pell 2000).

Finally, microbes and predators may act synergistically when sublethal doses of the pathogen make the pest more vulnerable to predation, perhaps due to reduced defence responses or through extended larval development times. Such effects have been observed for Colorado potato beetle (*Leptinotarsa decemlineata* Say) larvae that were more successfully attacked by the predator *Perillus bioculatus* Fabricius after treatment with sublethal doses of *B. thuringiensis* (Cloutier and Jean 1998). Similarly, microbes can make pests more vulnerable to some biopesticides such as neem (Mohan et al. 2007). To our knowledge, there are no studies that categorically demonstrate synergistic or additive effects from combined treatments of microbes and predators in greenhouse crops, but, based on the studies mentioned, this could be a promising area for future research. This might be particularly interesting for pests, such as western flower thrips (Almeida and Janssen 2012) and some caterpillars, e.g. *C. chalcites* (G.J. Messelink pers. obs.), that have strong defensive responses to their arthropod natural enemies.

Increasing efficacy with new formulation and application techniques

The success of a biological control programme with microbes depends on how much of the agent used reaches the target pests. For this reason, the way that microbes are formulated and applied is particularly important because the pathogen must be able to survive under greenhouse conditions but must also reach the intended pest (Lohse et al. 2015). Formulation plays an important role in delivering the pathogen to the target environment.

Formulated microbes are typically prepared as technical concentrates, wettable powders or oil dispersions (de Faria and Wraight 2007). Some adjuvants and other ingredients can improve the persistence of microbes in the environment by protecting them from inactivation by sunlight (Reddy et al. 2008; Shapiro 1992). The most promising technology to date is encapsulation within a matrix, which provides a stable microenvironment and protects the microbial from biotic and abiotic stress factors (contamination, soil antagonists, temperature, dryness, UV light, mechanical stress) (Vemmer and Patel 2013). This extends the shelf life and maintains metabolic activity for prolonged time periods not only during storage but also after application resulting in a reduction in the dose and number of applications required.

Traditionally microbes have been applied in a similar way to pesticides (Chandler et al. 2011), with associated problems of how to ensure that microbial formulations survive passage through spraying devices and the potential negative impacts of massive inundative applications of microbes on beneficial invertebrates, such as arthropod natural enemies and pollinators. To avoid massive applications of microbes, techniques for new applications have been developed. One such alternative option is seed treatment. Keyser et al. (2014) found that when seeds were coated with *Metarhizium* species, the fungi were able to disperse via the roots and induce infections in insects feeding on the roots. Specifically, in laboratory and greenhouse experiments, *Tenebrio molitor* Linnaeus larvae were exposed to roots of wheat plants that had been grown from seeds inoculated with conidia of either *M. brunneum* or *M. robertsii*. All four *Metarhizium* isolates tested maintained pathogenicity towards *T. molitor* larvae for up to 4 weeks after being dispersed by roots through both artificial substrates and nonsterile soil. Based on these results, the authors proposed that plant–root associations improved the mobility of entomopathogenic fungi in the soil and increased their likelihood of encountering susceptible insect hosts. Another way to achieve this plant–root colonization is the inoculation of growing substrates with *Metarhizium* sp., which appeared very effective in controlling black vine weevil larvae, *Otiorhynchus sulcatus* F., up to a year after application (Bruck 2005, 2010). Adding entomopathogens to growing media can also be useful for controlling soil-dwelling stages of pests that do not attack roots, such as pupae of western flower thrips (Ansari et al. 2008). With further research, treating seeds or growing media with entomopathogenic fungi has the potential to be successfully applied in a wide range of greenhouse conditions.

Another interesting technique to achieve more targeted application of microbes is the auto-dissemination approach, which aims to promote the transmission of

infective pathogen propagules within and between insect populations. For example, the ability of bumblebees to disseminate conidia of *B. bassiana* from hive-mounted dispensers to greenhouse crops shows potential for control of whiteflies, thrips and *Lygus* species on tomatoes and sweet pepper (Al-mazra'awi et al. 2006; Kapongo et al. 2008). Arthropod natural enemies may also contribute to the dispersal of microbials as described previously (Roy and Pell 2000). Promising results were achieved with predatory mites; when dusted with *B. bassiana* prior to their release, *A. swirskii* and *N. californicus* increased mortality in the pest *Diaphorina citri* Kuwayama, while maintaining low mortality of their own species (Zhang et al. 2015).

Auto-dissemination can be targeted even more specifically by using devices that contain both entomopathogens and attractive species-specific semiochemicals; the pests enter the device in response to the semiochemical, become contaminated by the pathogen (which will eventually cause their death), exit from the device and spread the infection amongst conspecifics (Baverstock et al. 2010; Lacey et al. 2015; Vega et al. 1995, 2007). To increase the chances of infection, traps are baited with highly attractive semiochemicals that indicate the presence of food, mates or oviposition sites. This method is often known as 'lure and infect' as opposed to 'lure and kill' because the pests do survive for a short time during which they interact with other individuals promoting transmission and the development of epizootics. Combinations of pheromone lures and entomopathogenic fungi have been successfully used to control insects such as bark beetles (*Ips typographus* Linnaeus), weevils (*Cylas formicarius* Fabricius, *Cosmopolites sordidus* Germar), moths (*Plutella xylostella* Linnaeus), greenbugs (*Plautia crossota* stali Scot), thrips (*Megalurothrips sjostedti* Trybom) and aphids (*Phorodon humuli* Schrank) under field conditions (Hartfield et al. 2001; Kreutz et al. 2004; Lopes et al. 2014; Mfuti et al. 2016; Pell et al. 1993; Roditakis et al. 2008; Tinzaara et al. 2007; Tsutsumi et al. 2003; Vickers et al. 2004; Yasuda 1999). However, the use of pheromones has limitations because in many cases aggregation pheromones, which attract both sexes, are unknown and sex pheromones tend to be sex-specific i.e. only one sex is attracted and controlled. Nevertheless, other olfactory cues such as food and host odours also have potential as attractants (Klein and Lacey 1999; Lecuona et al. 2005; Renn et al. 1999). For instance, combining the codling moth granulovirus (CpGV) with apple-associated yeasts increased the mortality of codling moth (*Cydia pomonella* Linnaeus) under laboratory and field conditions (Knight and Witzgall 2013). The potential for combining microbials with semiochemicals for pest control under greenhouse conditions is evident, especially considering that key greenhouse pests such

as thrips, moths and aphids are susceptible to pathogens and there is a vast amount of knowledge concerning their chemical communication already available (Elimem et al. 2014; Furlong and Pell 2001; Niassy et al. 2012; Pickett et al. 1992). However, for lure and infect strategies to be successful in greenhouse crops, more research is required to develop and optimize lure and infect devices and to evaluate such systems under greenhouse conditions.

Conservation of microbials in greenhouse crops

Conservation biological control relies on modification of the environment or management practices, to protect and encourage natural enemies that are already present within the system (Landis et al. 2000). Various methods are used for conservation of arthropod natural enemies in greenhouse crops. For example, the use of banker plants that provide alternative hosts or food resources (Messelink et al. 2014), but conservation of microbials has not received much attention. This is probably because microbials have traditionally been used to react to outbreaks of pests rather than as a preventive control measure. However, in the review paper of Pell et al. (2010) several methods to conserve fungal entomopathogens and stimulate natural epizootics are suggested. These include: (1) preventing intense soil disturbance as many fungal entomopathogens have at least part of their life cycle in the soil, (2) preventing frequent use of fungicides, (3) encouraging non-pest alternative hosts and (4) encouraging other arthropods for dispersal of conidia. Although these practices are not widely used in greenhouse crops, some organic growers are exploring the use of banker plants to encourage non-pest aphids infected by entomophthoralean fungi and thereby induce natural epizootics of these fungi (Messelink & Dinu, personal observations). There are several options for the use of non-pest aphids as reservoirs for these specialized aphid pathogens; for example the grain aphid, *Sitobion avenae* (Shah et al. 2004), which is commonly used as a banker plant system for aphid parasitoids (Huang et al. 2011) could also support multiplication of specialist aphid pathogens. Because these aphids are adapted to monocotyledonous plants, they cannot survive on dicotyledonous crops grown in greenhouses, thus using such aphids poses no risk and is relatively simple. Furthermore, the banker plant itself may contribute to the survival and persistence of entomopathogens. Hairy leaves may provide a better microclimate and leaf waxiness could increase adhesion and germination of conidia on the insect cuticle (Cory and Hoover 2006). Besides the provision of alternative hosts on banker plants, companion plants in general could be considered to support the survival of microbials. An additional benefit of these banker and companion plants could be that they provide pollen and nectar for arthropod

natural enemies and bees that could simultaneously aid dispersal of entomopathogens from the banker to the crop as they move back and forth between them, but this idea requires further elaboration.

Uniquely, greenhouses also provide the opportunity to use UV-blocking screens that can increase the survival of microbials. Several studies have shown the damaging effects of UV light on entomopathogenic fungi and viruses (Braga et al. 2001, 2002; Cory and Hoover 2006) and that survival of fungal entomopathogens is better in greenhouses with UV-blocking screens than in greenhouses without screens (Diaz and Fereres 2007). Not only plastic covers, but also greenhouse glass can partly or totally filter out UV-B wavelengths (Hemming et al. 2012), which will be beneficial for the survival of entomopathogens. Finally, by combining conservation tools with the selection of microbial isolates with particular traits that make them easier to conserve after application (such as persistence), could increase their biological control potential (Cory and Franklin 2012).

Balancing climatic adaptation for microbials and arthropod natural enemies

Greenhouse systems have been developed to protect crops from unfavourable environmental conditions and, depending on the geographical area, these systems range from low-tech plastic tunnels to high-tech glasshouses (van Henten et al. 2006). Globally, various technologies for managing temperature, humidity, light and CO₂ levels are increasingly used to optimize the environment for crop growth (Montero 2011). The most advanced technologies are mainly driven by the need to save energy and reduce fossil fuel consumption (Vadiee and Martin 2014). This need has resulted in new concepts such as the closed greenhouse system where cooling by ventilation is replaced by mechanical cooling and excess solar energy is collected and stored to be reused to heat the greenhouse (De Gelder et al. 2012). Implementing such new techniques enhances the possibilities to completely manage the greenhouse climate. Interestingly, these new techniques also offer new opportunities to temporarily adapt the greenhouse climate for other purposes, such as optimizing pest control with microbials. For example, increasing greenhouse humidity levels significantly increased pest control with *B. bassiana* (Shipp et al. 2003). Such decisions obviously need to be considered carefully, as some climatic benefits for microbials may have detrimental effects on crop growth, arthropod natural enemies or may favour some plant diseases. Also, it needs to be mentioned that the relative humidity in the microclimate can be significantly different to the ambient conditions. For example, high humidity levels at the leaf surface may be sufficient for infection by

entomopathogenic fungi, even when the ambient relative humidity levels are lower. One important advance in greenhouse climate management is the development of temperature integration regimes for reducing energy consumption, which allow higher temperatures during the day to be tolerated and compensated for by lower temperatures during the night (Körner and Challa 2003). Experiments showed that an extra benefit of this regime could be a reduced influx of thrips from outside into the greenhouse, because vents are opened less frequently than when traditional climate management is used (Jakobsen et al. 2006). The lower temperatures at night and in the early hours of the morning may also increase efficacy of Entomophthorales (Milner et al. 1984), but further research is needed to determine whether the higher day temperatures might be detrimental for the survival of these fungi and other microbials. Potential negative effects on arthropod natural enemies also require further evaluation. For example, lower night temperatures might be detrimental for arthropod natural enemies that are night active, such as the aphid predatory midge *Aphidoletes aphidimyza* Rondani, which requires a minimum temperature to be flight-active (Markkula et al. 1979). Diurnal temperature ranges of more than 15 °C could also be detrimental for biological control of spider mites by *Phytoseiulus persimilis* (Vangansbeke et al. 2015).

Overall it is a huge advantage to have the opportunity to adapt the greenhouse climate in order to optimize microbial efficacy, particularly when the potential damage by the target pest is larger than the potential crop losses that may occur due to less favourable climatic conditions, plant diseases or reduced efficacy of some arthropod natural enemies. Not only temperature and humidity, but also artificial light in greenhouses might influence pest control (Johansen et al. 2011), but the effects on microbials are relatively unknown. While it is known that the survival of fungal conidia is significantly reduced when exposed to UV-A and UV-B irradiation (Yao et al. 2010), and as we mentioned earlier, there are opportunities to enhance pest control by fungal microbials by using UV-blocking covers (Costa et al. 2001). This technique seems to be compatible with natural enemies, as no negative effects have been found to date (Dáder et al. 2015; Doukas and Payne 2007). Hence, there are several methods now available to adapt the greenhouse environment and these warrant further study. For example, which combinations of artificial light and greenhouse climatic conditions are optimal for pest control with microbials and arthropod natural enemies? It is not only the direct environmental effects on microbials, but also the indirect effects through changes in pest behaviour that could affect these results (Johansen et al. 2011).

Endophytes for pest control

A relatively new field of research in pest control with microorganisms aims to make use of fungal species that act as endophytes. The term endophyte refers to fungi and bacteria that develop within plant tissues without causing any conspicuous symptoms of disease in the plant (Wilson 1995). Besides the class II fungal endophytes (sensu Rodriguez et al. (2009)), which have a broad host plant range and are able to survive in the environment without a host, entomopathogenic fungi are also known to colonize plant tissues (see recent review by Vidal and Jaber (2015)).

So far several studies provided evidence that entomopathogenic fungi are able to confer at least partial resistance to their host plants when colonizing plant tissues. For instance, prior inoculation of tomato plants with the endophytic fungus *Acremonium strictum* Gams resulted in a significant decrease in the performance of pests such as greenhouse whiteflies (*Trialeurodes vaporariorum* (Westwood)) and the cotton bollworm (*Helicoverpa armigera* (Hübner)) compared with noninoculated plants (Jallow et al. 2004; Vidal 1996). Other examples include control of the diamondback moth, *Plutella xylostella* (Linnaeus), on cabbage inoculated with an endophytic strain of *A. strictum* (Raps and Vidal 1998); reduction in populations of *Aphis gossypii* Glover on squash plants previously inoculated with an endophytic isolate of *Fusarium oxysporum* Snyder & Hanssen (Martinuz et al. 2012); and retarded development of *Thrips tabaci* Lindeman on onions inoculated with several species of fungal endophytes (Muvea et al. 2014). The underlying mechanisms mediating the effects on herbivorous insects remain, however, to be investigated in detail. The effects were hypothesized to be due to metabolic products produced by the endophytic fungi or by plant metabolites induced by their presence. Compounds produced by Class II fungal endophytes are mainly extracellular enzymes like proteases, lipases and cellulases, and metabolites like beauvericin, oosporein, fumonisin, harzianolide, butenolide and fusaric acid (Gurulingappa et al. 2010, 2011; Ownley et al. 2008; Vega et al. 2009).

However, some natural enemies also feed on the plants harbouring endophytic fungi and could attack herbivorous insects that may be experiencing negative effects due to fungal endophytes. This appeared to be the case for the omnivorous predatory bug *M. pygmaeus* on tomato plants previously inoculated with a nonpathogenic isolate of *F. oxysporum* (Messelink et al., unpublished data). Surprisingly, this endophyte still enhanced pest control by deterring the predator from feeding on the plant, thereby increasing its feeding activity on the prey. The parasitoid *Bracon hebetor* Say, when parasitizing larvae of *Spodoptera litura* Fabricius feeding on cauliflower inoculated by *Aspergillus* spp., experienced prolonged development

time of the larvae and reduced parasitism rates by the adults (Kaur et al. 2015). Thus, the use of Class II endophytic fungi for control of herbivorous pests in greenhouses needs to be tested on a case by case basis for each plant-herbivore-natural enemy-interaction.

One of the most striking recent discoveries with regard to endophytic fungi is that several, if not all, isolates of entomopathogenic fungi can also act as endophytes. Most research on entomopathogenic fungi has focused on their virulence and the mechanisms they use to infect the host directly through the cuticle (Vega et al. 2008). However, endophytic isolates of *B. bassiana*, *M. brunneum* and *L. muscarium* that are active within plants have shown potential to control insects such as the corn borer *Ostrinia nubilalis* Hübner (Bing and Lewis 1991); the coffee berry borer *Hypothenemus hampei* Ferrari (Posada et al. 2007); the banana weevil *Cosmopolites sordidus* (Akello et al. 2008); the aphids *Acyrtosiphon pisum* Harris and *Aphis fabae* Scopoli (Akello and Sikora 2012); the American bollworm *H. armigera* (Qayyum et al. 2015) and the pea leafminer *Liriomyza huidobrensis* Blanchard (Akutse et al. 2013).

Bacterial endophytes have been studied mainly for their potential as plant growth promoters and for their ability to induce systemic resistance to plant pathogens (Kavino et al. 2007; Ryan et al. 2008; Sturz et al. 2000). Direct control of pests has not been reported for bacterial endophytes. Indirect effects caused by the induction of systemic resistance and by changes to the chemical profile of plants have been suggested as having the potential to influence the performance of some pests (Kloepper and Ryu 2006; Valenzuela-Soto et al. 2010).

The fact that bacterial and fungal endophytes might prime plants for resistance to both pests and diseases makes them formidable as biological control agents. For example, endophytic isolates of *F. oxysporum* have the ability to suppress aphids and also nematodes, Oomycota and other plant diseases (Fuchs et al. 1997; Hallmann and Sikora 1996; Kim et al. 2007; Martinuz et al. 2012). The use of endophytes that decrease performance by extending pest developmental time might also provide an opportunity to increase the impact of arthropod natural enemies in greenhouses. Irrespective of the mechanism, the longer it takes the pest to develop and complete its life cycles, the more exposed it is to parasitoids and predators. Finally, the use of endophytes introduced at an early stage of plant development (such as seeds coated with endophytes) circumvents the intrinsic problems of microbial use such as exposure to detrimental environmental factors, competition with other microorganisms and the challenge of synchronizing microbials with herbivore presence. However, entomopathogenic fungi are obviously not able to fully colonize all tissues of fast growing plants (Behie et al.

2015), resulting in several plant parts not being protected, and the interactions between specific entomopathogenic fungal isolates, the plant cultivar and the growing media (Tefera and Vidal 2009) seems to influence the extent to which these fungi can mediate herbivore preference and performance. Climate regimes in greenhouses may be beneficial for most fungi because they provide specific environmental conditions (constant higher temperatures and humidity), and most probably are also favourable for endophytic establishment in plants. However, isolates specifically adapted to these conditions also need to be selected to increase colonization rates of plant tissues. Thus, a screening process to find the optimal combination of the different agents is mandatory for the implementation of this strategy. The use of endophytic organisms as biological control agents might become even more complicated as a result of the findings of Yan et al. (2015) who demonstrating a ‘fight for niches’ within single plants, for regions not yet colonized by other endophytic organisms. Endophytes are known for their complex metabolic capabilities (Gutierrez et al. 2012) and some of the compounds they produce which are aimed at maintaining a multipartite symbiosis with competing microorganisms (Schulz et al. 2015), might also have adverse effects when consumed by humans. Thus, the method and timing of the inoculation of the crop plant, and the previous history of microbial colonization events, plays an important role in the potential for success with this strategy. Some of these problems may be addressed by the development of specific formulations for endophytes that enhance colonization rates and provide a headstart for these organisms within seedlings. However, these aspects need to be rigorously tested to gain a better understanding of the potential of microorganisms as biological control agents.

Insect-associated microorganisms

In recent years, it has been increasingly recognized that microbial symbionts affect the ecology, life history and evolution of their arthropod hosts in many different ways (Zchori-Fein and Bourtzis 2012). A lot of the key greenhouse pests are exclusively sap feeders, and like other insects with limited diets, harbour a wide range of microbial associates (Zchori-Fein and Bourtzis 2012). All phloem and xylem-feeding pests harbour maternally inherited, intracellular, nutritional bacterial symbionts; these are ‘obligate’ or ‘primary’ symbionts that provide their hosts with essential nutrients missing in the plant sap and are often restricted to specialized organs called bacteriomes. In addition, sap-feeders also commonly harbour facultative or ‘secondary’ symbionts which are involved in nearly all aspects of their host’s biology, including protecting them against natural enemies (Oliver et al. 2003;

Teixeira et al. 2008) and heat stress (Montllor et al. 2002); their performance on crops (Hosokawa et al. 2007); and in increasing their ability to transmit plant viruses (Gottlieb et al. 2010; Van den Heuvel et al. 1994). These ‘secondary’ symbionts also manipulate the reproduction of their hosts in ways that increase the frequency of female hosts becoming infected (e.g. Werren et al. (2008)). Furthermore, sap-feeders are usually associated with extracellular gut microbes, which may have nutritional or metabolic functions (Kikuchi et al. 2011; Werren et al. 2008). Although studies on host-microbial symbionts in insects have focused on bacteria, archaeal and fungal symbioses have also been reported (Gibson and Hunter 2010).

Zindel et al. (2011) summarized the various ways by which symbiotic microorganisms may dramatically affect all phases of augmentative biological control, from the mass rearing of natural enemies to their efficiency in the field. For example, symbiotic bacteria may induce cytoplasmic incompatibility, resulting in sterile eggs when a symbiont-infected female mates with an uninfected male. A mixed population of symbiont-infected and noninfected individuals would thus reproduce more slowly than one in which all members carried the symbiont or where all members were symbiont-free. In addition, symbiotic microorganisms can increase or decrease the survivorship of an insect host under extreme environmental conditions or influence the transmission capacity of disease-vectoring arthropods (Zindel et al. (2011) and refs there in).

Amongst their other phenotypes, insect microbial associates may protect their hosts against natural enemies such as parasitoids, bacteria, viruses and fungi, and these ‘defensive’ interactions have been thoroughly reviewed (Oliver et al. 2014). Most relevant to this review are examples concerning greenhouse pests with no effective biological control solutions (Table 2), such as aphids and whiteflies. The pea aphid, *Acyrtosiphon pisum*, has four different symbionts belonging to the bacterial genera *Regiella*, *Rickettsia*, *Rickettsiella* and *Spiroplasma*, which induce defences against the entomopathogenic fungus *P. neoaphidis*. The presence of these distantly related symbionts in aphids reduces both their susceptibility to *P. neoaphidis* infection and sporulation on *P. neoaphidis*-infected aphid cadavers (Łukasik et al. 2013). Although information on the symbionts of aphid species that are greenhouse pests is rather scarce, bacteria such as *Rickettsia* have been reported from the cotton aphid, *A. gossypii*, and the potato aphid *M. euphorbiae*, although the effects of these symbionts on their phenotypes has not been determined. Recently Hendry et al. (2014) showed that bacterial symbionts from the genus *Rickettsia* greatly reduced the efficacy of the entomopathogenic bacterium *Pseudomonas syringae* Van Hall against the sweet potato whitefly, *B. tabaci*. There is every reason to assume that similar symbiotic interactions may

occur in other insects colonizing greenhouse crops, and this needs to be taken into account when microbials are produced and applied for pest control in greenhouses and other crop systems.

Symbionts are, so far, not deliberately added to biological control systems, but this might be a field for future research. For example, aphid biotypes that contain particular symbionts might be used to select for parasitoids that are adapted to these symbionts, in order to enhance their performance in the field.

Conclusions and recommendations

Although greenhouse crop production seems ideal for the use of biological control agents, experience has taught us that there are a range of interactions that must be considered when incorporating organisms into a closed system. For decades, research focussed on understanding the tritrophic interactions between plants, herbivores and their arthropod natural enemies (Kennedy 2003; Price et al. 1980; Vet and Dicke 1992). However, it is now clear that tritrophic interactions are influenced by all the other components of the ecosystem, and as a result, research has shifted into understanding crop-pest management from the perspective of multitrophic interactions (Van der Putten et al. 2001). As we have shown here, the combined use of microbial biological control with the use of arthropod natural enemies must take into account direct and indirect effects on each side.

Understanding the complexity of ecological interactions between different types of biological control agents is itself a subject that requires further research. In terms of biological control, studies need to focus on unveiling direct and indirect effects of the application of microbial biological control agents and microbial communities within insects (symbionts) and plants (endophytes) on arthropod natural enemies. Although most cases show evidence of compatibility between different types of agents, it is necessary to pinpoint the essential conditions that ensure the success of their combination. We consider that the following research areas should be prioritized: (1) optimizing the efficacy of the existing and new microbial products and (2) determining how to combine both microbial and arthropod natural enemies with available technologies. Optimization of microbial efficacy is an ongoing goal achieved by new formulation and conservation techniques and by selection of specific and virulent isolates with the most desirable characteristics to be effective and persist under greenhouse conditions. Combining microbials with the existing and new technologies is already promising since greenhouse climate control and new delivery methods are already available. The use of endophytes and

manipulation of symbionts to improve control of pests could also represent elegant solutions to many of the current problems with microbials, but further research is required.

Understanding of the ecological consequences of using microbials in combination with other biological control agents needs closer examination, especially for organic greenhouse cropping systems that have evolved into complex ecosystems with persistent populations of multiple arthropod natural enemy species. To our surprise, there are only a limited number of studies involving microbials and arthropod natural enemies under greenhouse conditions, confirming that this area is fertile ground for research. In our opinion, such studies deserve more attention as they may help to identify complementary and synergistic interactions between microbials and arthropod natural that increase the opportunities to enhance and further develop biological pest control.

Authors contribution

All authors contributed to the writing of this review. FG and GM edited and finalized the manuscript.

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