# Advances in biostimulants as an integrated pest management tool in horticulture

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## **1** Introduction

Plant biostimulants are microorganisms, microbial derivatives, and materials of mineral, marine, or botanical origin that stimulate plant physiological responses including induced systemic resistance (ISR), and improve nutrient or water uptake, promote plant growth and health, or impart tolerance to biotic and abiotic stressors, without having nutritional or pesticidal properties. This definition covers both the technical and the regulatory aspects of biostimulants. However, there are some exceptions to the biostimulant products that are available in the market, due to the inherent nutritional or pesticidal properties of some of the active ingredients or their sources. For example, silicon has biostimulant properties, but when applied as calcium or potassium silicate,

plants receive calcium and potassium as additional nutrients along with silicon (Savvas and Ntatsi, 2015). Some protein hydrolysates have several nutrient components in addition to biostimulant activity (Baglieri et al., 2014; Visconi et al., 2015). Similarly, certain microorganisms have both biostimulatory and biopesticidal properties and depending on the strain or application procedure, they are registered or used for one of these purposes (Kim et al., 2017; Fiorentino et al., 2018; Dara, 2019a). Crop habitat is a dynamic environment with complex interactions among various biotic and abiotic elements. Sometimes, biostimulatory and biopesticidal properties overlap, interchange, or are activated or synergized by their interactions with other influencing factors. To avoid under-exploring the full potential of biostimulants, this chapter does not involve the regulatory aspects but focuses only on the biostimulatory properties of various active ingredients and how they can be used in integrated pest management (IPM) programs as a part of cultural control. The IPM section briefly describes the new IPM model (Dara, 2019b), the role of biostimulants in improving crop production and protection, and lists various categories of biostimulants. The following sections focus on the role of biostimulants in enhancing crop growth, health, and yield, improving soil health, inducing natural resistance, suppressing diseases and arthropod pests, and imparting tolerance to abiotic stresses. Various interactions of biostimulants with other biotic and abiotic factors in the crop environment, the different strategies for successfully using biostimulants in food production, and the challenges and future needs in promoting the use of biostimulants will be discussed in the concluding sections.

#### 2 Integrated pest management

The traditional definition of IPM was limited to pest monitoring, various control options, and making treatment decisions based on economic thresholds with a focus on protecting the ecosystem (Peterson et al., 2018). In many cases, the rotational use of pesticides with different modes of action, mainly intended for managing pesticide resistance, is considered as practicing IPM. In general, a heavy emphasis on minimizing pesticide applications with a focus on environmental safety, a lack of consideration for factors that influence pest management practices at the farm level, and various other reasons have limited the widescale implementation of IPM strategies. The transformation of agriculture as a global enterprise, the technological advances in multiple disciplines of crop production, the growing world populations and their food needs, and an increasing demand for sustainably produced food warranted a practical IPM model that is economically viable, environmentally sustainable, and socially acceptable. As a result, the new IPM model has been developed to meet the needs of modern-day agriculture (Dara, 2019b). The new model provides

a template for an integrated crop production system that makes the best use of traditional and modern technologies, incorporates information management and effective decision-making, and emphasizes the importance of research and outreach to maintain farm profitability and food affordability (Fig. 1). The knowledge of pests, the vulnerable stages of their life cycle, and the influence of various agronomic practices on pest populations and their damage are critical in the new IPM model as they help devise and implement effective strategies. Good agronomic practices such as using biostimulants not only improve crop health but also lessen the burden on crop protection directly or indirectly. Biostimulants help crop production in one or more of the following ways (Fig. 2):

- 1 Stimulate ISR and prime the plants for biotic and abiotic stressors.
- 2 Improve water and nutrient use efficiency and thus reduce the input costs and associated environmental risks from leaching or runoff,



**Figure 1** The new integrated pest management (IPM) model provides a template for various crop and pest situations to produce crops in an economically viable, socially acceptable, and environmentally sustainable manner (Dara, 2019b).

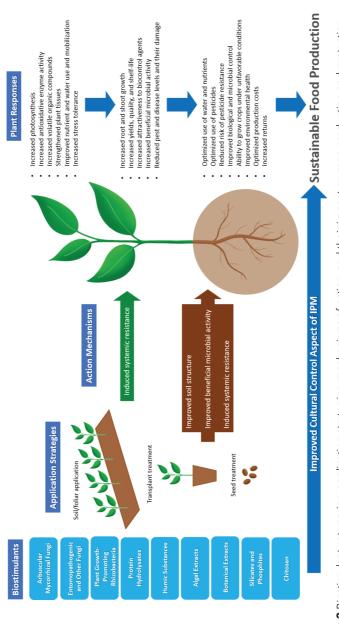


Figure 2 Biostimulant categories, application strategies, mechanisms of action, and their impact on crop production and protection.

enhance plant vigor and yields and thus compensate for losses due to pest damage.

- 3 Help optimize fertilizer inputs and prevent overuse of certain nutrients that increase a plant's attractiveness to pests.
- 4 Strengthen plant tissues and reduce pest damage or proliferation.
- 5 Release metabolites or enhance the production of plant defense chemicals that negatively influence pest populations or attract natural enemies.
- 6 Contribute to the reduction of pest management practices, pesticide resistance, and the potential negative impact on the environment.
- 7 Build soil structure and promote natural beneficial microbial populations that further improve crop health in the short term and improve soil health in the long term.
- 8 Contribute to carbon sequestration and a reduction in the use of synthetic agricultural inputs.

One key area of IPM is focusing on preventive measures and minimizing the reliance on curative measures. Biostimulants are good agronomic inputs that contribute to these preventive measures as a part of cultural control. Biostimulant use has increased in recent years as a result of increased market availability, research and outreach efforts, the need for improved crop production practices, and the interest in sustainably produced food items. Examples of some research studies with commercially available or developmental products that have demonstrated the benefits of various categories of biostimulants, including arbuscular mycorrhizal fungi (AMF), entomopathogenic and other fungi, plant growth-promoting rhizobacteria (PGPR), protein hydrolysates, humic substances, algal and botanical extracts, and minerals are discussed in the following sections.

## 3 Enhancement of crop growth, health and yield

Biostimulants are primarily used for improving crop growth, health, and yields through one or more mechanisms. Increased nutrient and water uptake through enhanced root growth, increased solubility or bioavailability of nutrients, higher beneficial microbial activity, and improved metabolic activity are among various mechanisms that aid plant growth and health.

## 3.1 Arbuscular mycorrhizal fungi (AMF)

AMF have been used in crop production for a long time for their biostimulatory properties. *Glomus aggregatum* significantly enhanced the growth of grape rootstocks with improved root morphology (Aguín et al., 2004). Inoculation

of *Glomus mosseae* resulted in higher biomass and concentration of some nutrients in shoots, and higher single-fruit weight in cucumbers (Wang et al., 2008). *Glomus versiforme* inoculation improved osmotic adjustment, leaf water potential, and soluble sugar content, and increased the transpiration and photosynthetic rates among other changes, resulting in improved plant growth of tangerine (Wu and Xia, 2006). A review by Rouphael et al. (2015) discussed yield and growth improvement of several horticultural crops growing under nutrient-deficient conditions with the help of AMF.

#### 3.2 Entomopathogenic and other fungi

Although not primarily used as biostimulants, entomopathogenic fungi such as *Beauveria bassiana*, *Cordyceps fumosorosea*, and *Metarhizium* spp. appear to have biostimulatory properties such as promoting plant growth through improved nutrient and water uptake and alleviating plant stresses. A strain of *Beauveria bassiana* is currently available in the United States as a soil amendment, for its biostimulant properties. A recent review of entomopathogenic fungi described various non-entomopathogenic roles and how they can be exploited as holistic tools (Dara, 2019a). When entomopathogenic fungi are applied for arthropod pest control, they also benefit plants through their endophytic and mycorrhiza-like roles and improve plant growth and health. Additionally, entomopathogenic fungi could also serve as the substitutes of AMF for crops that have a poor relationship with AMF.

The mycoparasitic fungal species of *Trichoderma* are commonly used as biofungicides, but some of them also have biostimulatory properties and are sold as biofertilizers. A biofertilizer enriched with *Trichoderma harzianum* significantly increased tomato yields, quality, and nutritional value and contributed to the reduction of synthetic fertilizers (Molla et al., 2012). A biofungicidal strain of *T. harzianum* increased the access of AMF to non-host brassica plants and prompted plant growth (Poveda et al., 2019). A *Trichoderma*-based biofertilizer enhanced the soil enzyme activity that improved nutrient availability and promoted the growth, quality, and yield of flowering Chinese cabbage (Ji et al., 2020). Similarly, positive results were seen in yield or other parameters from field studies conducted in California strawberries with biostimulants that have *T. harzianum* and *T. virens* or *T. harzianum* and *Bacillus amyloliquefaciens* (Dara, 2020, 2021), and a tomato study with a combination of products where one of them contained *T. harzianum* and some PGPR (Dara and Lewis, 2019).

#### 3.3 Plant growth-promoting rhizobacteria (PGPR)

The rhizosphere environment supports an abundance of microorganisms due to the presence of organic matter, root exudates, nutrients, and moisture.

Among these microorganisms are the PGPR, which regulate plant hormones, produce antibiotics and organic compounds, fix nitrogen, solubilize phosphate, improve soil structure, increase nutrient and water uptake, and alleviate plant stresses (Glick, 2012; Backer et al., 2018). Similarly, root exudates also influence soil microbial communities (Bakker et al., 2012). Thus, both plants and soil microorganisms influence and depend on each other. Examples of PGPR genera that are commonly used in crop production include Agrobacterium, Azospirillum, Azotobacter, Bacillus, Burkholderia, Delfitia, Paenibacillus, Pseudomonas, Rhizobium, Serratia, and Streptomyces that enhanced plant growth and yields in multiple studies. For example, Azospirillum brasilense improved root and shoot growth in strawberry (Pedraza et al., 2010), Bacillus subtilis increased shoot development and yields in apricot (Esitken et al., 2002), Pseudomonas fluorescens improved nutrient uptake and promoted plant growth and yields in broccoli (Tanwar et al., 2014), a consortium of Pseudomonas putida, Camanomonas testosterone, Citrobacter freundii, and Enterobacter cloacae and a nutrient program with a product containing Azotobacter chroococcum, B. subtilis, Bacillus megaterium, Bacillus mycoides, and T. harzianaum improved tomato yields (Dara and Lewis, 2019).

#### 3.4 Protein hydrolystates

Protein hydrolysates contain peptides and amino acids derived from enzymatic or chemical hydrolysis of animal- or plant-derived materials and have biostimulant effects that help plants in multiple ways (Colla et al., 2015). Significant improvement in marketable strawberry yields was seen when a hydrolysate of grocery store food waste was used as a substitute for the grower-standard nutrient program at 50% or 100% through drip application in a commercial field in California (Dara, 2017). An Italian study showed that protein hydrolysates improved the biomass of the aerial parts and fruit diameter, both under wellwatered and water-stressed conditions (Meggio et al., 2020). In another Italian study, a protein hydrolysate derived from legume seeds increased antioxidant activities, total soluble solids, mineral and nutritional value, photosynthetic activity, and yields of tomato (Rouphael et al., 2017). Such a positive impact on tomato yields and quality parameters was also seen in another study from a legume-based protein hydrolysate, seaweed extract, and a tropical plant extract (Colla et al., 2017). Similarly, a soy protein hydrolysate improved strawberry fruit yields comparable to synthetic fertilizers in a Californian study (Dara, 2020).

#### 3.5 Humic substances

Another group of biostimulants comprises humic substances, which are biologically derived organic matter in soil and water that regulate critical ecological and environmental processes and have a major impact on plant growth and metabolism (Nardi et al., 2002; Canellas et al., 2015). Reducing soil compaction, increasing nutrient use efficiency and microbial activity, and improving plant biomass are some of the benefits of humic substances with their hormone (auxin)-like activity (Nardi et al., 2002; Rose et al., 2014). In a Cuban study, humates from vermicompost decreased total carbohydrate content, increased protein content, nitrate uptake, and crop yields, and shorted the crop cycle by 21 days in lettuce (Hernandez et al., 2015). In a potted plant study in Italy, humic-like substances derived from alkaline hydrolysis of tomato plant residue increased the nitrogen assimilation in kidney bean (Baglieri et al., 2014). A meta-analysis by Rose et al. (2014) showed that exogenously applied humic substances increased the root biomass by 21% and the shoot biomass by 22%. Similarly, Seyedbagheri (2010) reported from 17 years of research studies in Idaho, USA, that a 11-20% yield increase was seen in potato from various humic acid products. An extensive review by Canellas et al. (2015) summarized the positive effects of humic substances in several horticultural crops including fruits such as apricots, grapes, pineapple, and strawberry, and vegetables such as broccoli, beans, eggplant, garlic, lettuce, onion, pepper, potato, and tomato with an increase in root dry weight by up to 124% and increase in yield by 80% in some of these crops.

#### 3.6 Algal extracts

Extracts from several species of micro- and macroalgae have been used in crop production. Species of Aphanizomenon, Chlorella, Dunaliella, and Spirulina are among the microalgae used for their biostimulant and biofertilizer properties. Species of Ascophyllum, Durvillea, Ecklonia, Saragassum (brown algae), Ulva (green algae), and Gelidium (red algae) are some of the common macroalgae or seaweeds used for improving plant growth or relieving them from stresses. A bioassay conducted in Brazil by immersing onion seedlings in a combination of the microalga Scendesmus subspicatus and humic acid showed synergism and promoted plant growth and improved bulb caliber, sugar and protein content, and yields (Gemin et al., 2019). An Indian study showed faster germination and plant growth, and improved root and shoot development of tomato from seed treatment; and higher chlorophyll content, and increased root and shoot development from foliar application of a microalgal consortium (Supraja et al., 2020). In an Italian study, Barone et al. (2018) showed that extracts of the microalgae Chlorella vulgaris and Scenedesmus quadricauda upregulated several genes of sugar beets putatively involved in sulfate starvation, and triggered primary and secondary metabolic pathways and intracellular transport, resulting in improved root development and nutrient uptake. Similarly, treating tomato and pepper plants with the polysaccharide extract of *Spirulina platensis* resulted in as much as a 30% increase in plant size, 230% increase in root weight, and 100% increase in size and number of nodes, in a Moroccan study (Elarroussi et al., 2016). Microalgae also have various plant hormones such as auxins, cytokinins, and gibberellins and influence several physiological processes related to plant growth, development, maturity, yields, and abiotic stress tolerance (Ronga et al., 2019).

Several macroalgal biostimulants are available around the world, and the majority are based on *Ascophyllum nodosum* (Sharma et al., 2014). Improved plant growth, chlorophyll content, nutritional value, yield, or other benefits were seen from macroalgal biostimulants in apple (Spinelli et al., 2009), onion (Araujo et al., 2012), orange (Fornes et al., 2002), peanut (Featonby-Smith and van Staden, 1987), strawberry (Dara and Peck, 2018), and various other crops. A review by Battacharyya et al. (2015) summarized the benefits of macroalgal extracts in several fruit and vegetable crops, with increased germination, plant growth, chlorophyll content, fruit yield and quality, nutritional value, and early maturity among the several parameters measured.

#### 3.7 Botanical extracts

Several botanical extracts are commonly used as biopesticides, and some of them have biostimulatory properties. Field studies conducted in California in strawberry and tomato with cold-pressed neem extract containing neem oil and azadirachtin, carboxylic acids from rice hulls, and extracts of black walnut and willow showed yield improvements (Dara and Peck, 2018; Dara, 2019c; Dara, 2021). Godlewska et al. (2019) evaluated raw extracts of dandelion, mugwort, and other plants applied to white head cabbage and observed an increase of shoot and root growth by 246% and 106%, respectively. Ali et al. (2019) reported that a single foliar pre-transplant application of garlic bulb extract improved plant growth and antioxidant activity of eggplant.

#### 3.8 Silicates and phosphites

In addition to macronutrients and micronutrients that ensure optimal plant growth and health, products based on silicon and phosphorus have biostimulatory effects of improving mechanical strength, modifying leaf architecture for improved exposure to sunlight, enhancing the activity of antioxidative enzymes to reduce oxidative damage, and improving water use efficiency, mineral uptake, and photosynthesis, among others (Gómez-Merino and Trejo-Téllez, 2015; Savvas and Ntatsi, 2015). Several studies have demonstrated the benefits of using silica-based products in improving plant growth or yields. For example, the application of silica clay improved the fresh weight of onion leaves by 46% (Araujo et al., 2012). Potassium silicate

application resulted in a numerical increase of tomato yields by 27% (Dara and Lewis, 2019). Substrate application of potassium silicate and foliar application of sodium silicate improved plant and flower characters of gerbera (Kamenidou et al., 2010). A combination of silicon- and seaweed-based products improved strawberry yields by 16% (Dara and Peck, 2018). Similarly, phosphite improved the yield parameters in celery, lettuce, onion, pepper, potato, strawberry, and tomato when applied as phosphorous acid (Rickard, 2000; Bertsch et al., 2009; Estrada-Ortiz et al., 2012) and emergence, early growth, and mycorrhizal colonization in potato seed tubers and root and shoot growth in strawberry when applied as potassium phosphite (Glinicki et al., 2010; Tambascio et al., 2014).

## 3.9 Chitosan

Chitosan is a polysaccharide derived from chitin in the exoskeleton of shellfish. It is also present in the arthropod exoskeleton and in fungi. Similar to other biostimulants, chitosan also induces plant defenses and imparts tolerance to various stresses. A review by Pichyangkura and Chadchawan (2015) discussed increased antioxidant activity in apricot, lychee, grapes, navel oranges, sweet basil, and tomato, and improved growth parameters in artichoke, chilli pepper, Indian spinach, okra, watermelon, and several ornamental crops, with chitosan application.

## 4 Enhancement of soil health

Soil is more than dirt and is full of life. Organic matter and microbial communities are the major components of soil, which influence its water holding capacity and nutrient content. Soils rich in organic matter and beneficial microorganisms improve the physical, chemical, and biological properties of soil, which in turn influence plant growth and health. Healthy soils are also important for nutrient cycling and water conservation. Most biostimulant materials enrich soils with organic molecules, beneficial microorganisms, or minerals. For example, humic substances, being a major component of soil organic matter, have a major impact on soil structure, microbial activity, and overall soil health. Humic substances also influence heavy metal mobilization in soil with improved leachability or plant uptake (Shahid et al., 2012; Canellas et al., 2015). Certain microorganisms such as *B. subtilis* can also be used for the bioremediation of soils (Nadhirawaty and Titah, 2019). Microalgae can remove antibiotics, pharmaceuticals, and personal care products from water and such benefits can be extended for improving soil health as well (Xiong et al., 2017; Kiki et al., 2019; Hena et al., 2021). AMF, PGPR, and other biostimulants fix, mobilize, solubilize, or increase the bioavailability of nutrients in soils and thus improve

soil health (Pindi and Satyanarayana, 2012; De Pascale et al., 2017). Soil health and soilborne disease incidence are closely related, and improved soil health through organic and microbial amendments and other cultural practices is important for disease suppression (Janvier et al., 2007). Similarly, improved soil quality and health through organic amendments and other practices not only improves water and nutrient availability, but also increases the resilience of soil against adverse environmental conditions and contributes to disease suppression (Lal, 2016).

## **5 Inducing natural resistance**

Plants respond to various beneficial and harmful stimulants of both a biotic and an abiotic nature and trigger defense mechanisms to naturally induce resistance to stresses. The mechanism where the salicylic acid pathway is activated in response to virulent and avirulent pathogens, nonpathogenic microorganisms, chemicals, and other stressors is called systemic acquired resistance (SAR) (Dara, 2021). The mechanism where jasmonic acid and ethylene pathways are activated in response to beneficial microorganisms is called ISR. Pathogenesis-related proteins produced in both mechanisms also help plants perform well under stress. While SAR is more of a curative or reactive strategy, ISR is a proactive strategy. Many biostimulants activate ISR and prime plants for potential stressors. Several studies have demonstrated induced resistance from biostimulants including algal extracts (Raghavendra et al., 2007; Sharma et al., 2014), AMF (Jung et al., 2012; Cameron et al., 2013), Trichoderma (Yoshioka et al., 2012), PGPR (Zehnder et al., 2001), protein hydrolysates and humic substances (Nardi et al., 2016), and chitosan (Nandeeshkumar et al., 2008). Primed plants have enhanced responses to stressors and thus mitigate their negative impact (Pastor et al., 2014). Although some biostimulatory active ingredients may also exhibit pesticidal properties, ISR is the primary mechanism in crop protection and in their role in IPM

## **6** Suppression of diseases

#### 6.1 Arbuscular mycorrhizal fungi and other microorganisms

Numerous studies have demonstrated the role of biostimulants in disease suppression through ISR and other mechanisms. Larkin (2008) reported that an AMF inoculant containing six *Glomus* spp., *Gigospora margarita*, and *Paraglomus brazianum* reduced stem canker and black scurf caused by *Rhizoctonia solani* in potato by 17-28%. Larkin (2008) also found that when aerobically composted tea was used alone and in combination with a mixture of six *Bacillus* spp., *Streptomyces griseoviridis*, *T. harzianum*, humic acids, and

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seaweed and yeast extracts in a crop rotation with barley/ryegrass, the incidence of stem canker and black scurf, and the common scab caused by *Streptomyces scabiei* declined by 18-33%, with a 20-23% yield increase. In another study, the *T. harzianum* strain T39 reduced symptoms of downy mildew caused by *Plasmopara viticola* by triggering resistance locally and systemically (Perazzolli et al., 2008), similar to that seen in resistant genotypes (Perazzolli et al., 2012). Similarly, a strain of *T. harzianum*, in the presence of root rot-causing *Fusarium solani*, upregulated several genes involved in plant defenses in olive (Amira et al., 2017).

## 6.2 Plant growth-promoting rhizobacteria

Wei et al. (2011) demonstrated that fortifying an organic fertilizer with two strains of B. amyloliquefaciens reduced the incidence of tomato wilt caused by Ralstonia solanacearum in greenhouse and field studies. Omara et al. (2017) reported that treating soybean seedlings with PGPR Methylobacterium aminovorans and M. rhodinum, Bradyrhizobium japonicum, B. megaterium var. phosphaticum, and the mycoparasitic fungus Trichoderma viride suppressed damping off caused by R. solani. These treatments also increased the nodule numbers; dry weight of nodules; nitrogen, phosphorus, and potassium content; and seed yield. Recep et al. (2009) observed that 17 strains of B. amyloliquefaciens, Bacillus atrophaeus, Bacillus pumilus, Bacillus macerans, Burkholderia cepacia, Flavobacter balastinium, and Pseudomonas putida inhibited some of the dry rot-causing Fusarium oxysporum, Fusarium culmorum, and Fusarium sambucinum, in vitro. However, only the OSU-7 strain of B. cepacia controlled the disease under storage conditions. Other studies demonstrated similar disease suppression including the control of shot-hole disease (Coryneum blight) in apricot with B. subtilis (Esitken et al., 2002), tomato yellow leaf curl virus with Enterobacter asburiae (Li et al., 2016), F. oxysporum and Fusarium guttiforme with chitinase from Chromobacterium violaceum (Sousa et al., 2019), and Verticillium dahliae with cyclodextrin glycosyltransferase, an antimicrobial protein, from Bacillus cereus YUPP-10 (Zhou et al., 2021).

## 6.3 Protein hydrolysates

Soybean and casein hydrolysates reduced downy mildew infection in grape by 76% and 63%, respectively (Lachhab et al., 2014). Both hydrolysates boosted the plant defenses by upregulating the genes responsible for pathogenesis-related proteins and the stilbene synthase enzyme involved in the production of resveratrol, a defensive phytoalexin. In another study, guar hydrolysate suppressed *Pdosphaera xanthii*, the causal agent of powdery mildew, in zucchini (Cappelletti et al., 2017).

#### 6.4 Humic substances

Disease suppression is one of the benefits of humic substances in addition to their impact on improving crop health and yields. Hernandez et al. (2015) observed reduced disease incidence in lettuce when vermicompost was applied. Increased phenylalanine ammonia lyase activity, which improves plant defenses, was thought to be responsible for disease suppression. Suppression of *Pythium* damping off in cucumber, *Rhizoctonia* root rot in radish, and *Verticillum* wilt in strawberry were observed from vermicompost application (Chaoui et al., 2002). Similarly, the combination of vermicompost and wood vinegar (pyroligneous acid) reduced *V. dahliae* in cucumber (Saberi et al., 2013). Several other studies demonstrated the impact of humic substances in improving crop health through disease suppression (Olivares et al., 2017; Jindo et al., 2020)

#### 6.5 Algal and botanical extracts

A review by Shukla et al. (2021) explained the role of seaweed extracts in priming plants and triggering defense mechanisms against plant pathogens. Seaweed polysaccharides and their derivate oligosaccharides activate the salicylic, the jasmonic acid, and the ethylene signaling pathways that increase the production of pathogenesis-related proteins and other compounds with antimicrobial activity, and offer protection from pathogens (Vera et al., 2011). For example, the seaweed polysaccharides ulvan, carrageenan, alginate, and laminarin improved total polyphenol and lignin contents, inhibited fungal growth, and reduced Verticillium wilt in olives (Salah et al., 2018). Extract of A. nodosum increased the activity of peroxidase, polyphenoloxidase and other enzymes, and several pathogenesis-related proteins in carrots, resulting in enhanced resistance to Alternaria radicina and Botrytis cinerea (Jayaraj et al., 2008). In a different study, laminarin suppressed scab (Venturia inaequalis) in apple by inducing plant resistance, but supplemental fungicidal applications were necessary to achieve the desired control (Van Hemelrijck et al., 2013). Foliar application or seed treatment with the extracts of brown algae Cystoseira myriophylloides, Fucus spiralis, and Laminaria digitata significantly reduced disease severity by V. dahliae and Agrobacterium tumefaciens in tomato through induced resistance (El Modafar et al., 2012).

Many botanical extracts are primarily used as biopesticides rather than biostimulants. Some products based on soybean and other vegetable oils are used as both biostimulants and biopesticides. However, in general, there are fewer examples where botanicals have been used exclusively as biostimulants and have indirectly contributed to pest or disease suppression. Willow bark extract, which is used as a plant growth regulator and biostimulant, also has fungicidal properties (Deniau et al., 2019). Treating potato with Canada milkvetch extract, *P. fluorescens* DF37, and *B. pumulis* M1 provided up to 84% of *Verticillium* control with up to 24% of yield increase (Uppal et al., 2008), but it was not clear whether the disease suppression was a result of biostimulation or the pesticidal activity of these materials.

#### 6.6 Silicates and phosphites

Sodium silicate inhibited conidial germination and mycelial growth of *Fusarium sulphureum* in controlled dry rot of potato tubers (Li et al., 2009). Calcium silicate reduced downy mildew (*Peronospora manshurica*) in soybean (Nolla et al., 2006), and when applied with calcium chloride, white mold (*Sclerotinia sclerotiorum*) in dry beans (Paula Júnior et al., 2009). Foliar application of silicates might act as a physical barrier to pathogen infection, but root application of potassium metasilicate increased plant defenses and controlled downy mildew (*P. xanthii*) in cucumber (Liang et al., 2005). Silicon, along with *G. mosseae* and *Enterobacter* sp. UPMSSB7, suppressed white root rot caused by *Rigidoporus microporus*, similar to a fungicide treatment, while improving various growth parameters of rubber seedlings (Shabbir et al., 2021).

In addition to its nutritional importance, phosphite has a significant influence on disease suppression. Föster et al. (1998) reported a significant reduction in root and crown rot caused by *Phytophthora capsici* in pepper plants treated with phosphite. Shearer and Fairman (2007) observed that phosphite application resulted in a significant reduction in the mortality of *Banksia* spp. from *Phytophthora cinnamomi* infection. Silva et al. (2011) reported yield improvement and downy mildew reduction in soybean from potassium phosphite application. They also observed that potassium phosphite application supplemented with fungicidal treatments significantly controlled Asian soybean rust caused by *Phakospora pachyrhizi* and powdery mildew caused by *Microsphaera diffusa*. In another study, Zhao et al. (2013) found that applying phosphorus oxyanion solution to sweet oranges with huanglongbing, caused by *Candidatus Liberibacter* spp., reduced the severity of disease symptoms and significantly improved fruit yields.

#### 6.7 Chitosan

Numerous studies have demonstrated the effect of chitosan on improved enzymatic activity, production of phytoalexins and pathogenesis-related proteins, lignin synthesis, and enhanced plant defenses, resulting in reduced disease incidence and increased postharvest quality (El Hadrami et al., 2010). Pichyangkura and Chadchawan (2015) thoroughly reviewed the plant protection offered by chitosan from several studies in vegetables, fruits, and ornamentals. Control of *F. oxysporum* f. sp. *radices-lycopersici* (Lafontaine and Benhanou, 1996) and *Xanthomonas gardneri* (Jail et al., 2014) in tomato, *Alternata, Fusarium avenaceus, F. culmorum, Epicoccum purpurascens,* and *S. sclerotiorum* in lemon balm (Szczeponek et al., 2006), *B. cinerea* and *P. viticola* in grapes when applied with copper sulfate (Aziz et al., 2006), *Colletotrichum capsici* in bell pepper during postharvest storage when applied with lemongrass oil (Ali et al., 2015), *Aspergillus flavus* (De Oliveira Pedro et al., 2013), *Aspergillus ochraceus* (Meng et al., 2020), and *Phytophthora infestans* (Huang et al., 2021) *in vitro*, and *F. oxysporum* in potato (Ren et al., 2021) are some of the examples of chitosan application in disease management.

## 7 Suppression of arthropod pests

## 7.1 AMF and other microorganisms

AMF can have an impact on herbivore damage by modifying the morphological and biochemical characters of the plant by augmenting defense mechanisms and imparting resistance. A recent review by Selvaraj and Thangavel (2021) discussed the role of AMF as a part of IPM against pest insects. Coppola et al. (2019) found that Trichoderma atroviride P1 in tomato contributed to the control of Egyptian cotton leafworm, Spodoptera littoralis, and the potato aphid, Macrosiphum euphorbiae, through direct and indirect defense mechanisms. An increased oxidative burst reaction and the production of volatile organic compounds that attract the parasitoid Aphidius ervi parasitizing aphids, and an enhanced production of protective enzymes against the leafworm, were found to be the defense mechanisms. A similar impact of increased defenses and attraction of A. ervi to parasitize M. euphorbiae was also seen in tomato from T. harzianum application, in an earlier study (Coppola et al., 2017). Some species of Trichoderma and their strains can also control pest insects through entomopathogenic activity. For example, T. atroviride, Trichoderma citrinoviride, and T. harzianum inhibited 73-85% of egg hatch in the bean weevil Acanthoscelides obtectus, and the polyphagous long-horned beetle Xylotrechus arvicola (Rodríguez-González et al., 2017). A Trichoderma sp. alone and in combination with Bacillus thuringiensis effectively controlled the brinjal shoot and fruit borer Leucinodes orbonalis, the cotton aphid Aphis gossypii, and the cotton leafhopper Amrasca biguttula biguttula (Nawaz et al., 2020). Trichoderma longibrachiatum controlled L. orbonalis and increased eggplant yields by 56% (Ghosh and Pal, 2016).

## 7.2 Plant growth-promoting rhizobacteria

As the inducers of plant defense mechanisms, PGPR elicit responses that influence herbivore infestations. While some PGPR may have entomopathogenic

properties, most of them suppress pest populations through ISR (Disi et al., 2019). For example, Zehnder et al. (1997) reported that when cucumber plants were treated with *B. pumilus* INT-7, feeding damage by the cucumber beetle, Diabrotica undecimpunctata howardi, and the bacterial wilt caused by Erwinia tracheiphila, which the beetle transmits, were significantly reduced due to the increased synthesis of cucurbitacin. In cotton, treating plants with Bacillus spp. resulted in higher levels of gossypol and jasmonic acid, leading to reduced larval feeding of the beet armyworm Spodoptera exigua (Zebelo et al., 2016). In peppermint, treatment with B. amyloliquefaciens GB03 and P. putida SJ04 increased salicylic and jasmonic acid levels and essential oil production, similar to the response from the feeding of the sunflower looper Rachplusia nu (del Rosario Cappellari et al., 2020). Although this study did not measure the impact of PGPR on herbivores or herbivory, it demonstrated that PGPR elicit defense responses. Several studies also demonstrated the influence of PGPR on natural enemies. In arugula, treatment with B. amyloliquefaciens GBO3 and the presence of the fall armyworm Spodoptera frugiperda, and the diamondback moth *Plutella xylostella*, influenced the behavior of the predatory earwig Doru lutiepes (Bell et al., 2020). The presence of PGPR and herbivore damage had a synergism in attracting earwigs. In rice, reduced leaf folder damage and increased natural enemy activity were seen from the inoculation of P. fluorescens (Saravanakumar et al., 2008). Similarly, P. fluorescens WCS417r colonization of the thale cress Arabidopsis thaliana increased the activity of the parasitoid Microplitis mediator when the plants were infested with the cabbage moth Mamestra brassicae (Pangesti et al., 2015). PGPR suppressed the production of herbivore-induced terpene, methyl salicylate, and lilial, and enhanced plant growth and ISR.

#### 7.3 Humic substances

A review by Joshi et al. (2015) presented several examples of insect control with vermicompost or vermiwash applications. Aphids, beetles, earworms, hornworms, mealybugs, and mites were controlled on cabbage, corn, cucumber, eggplant, mustard, pepper, and tomato in multiple studies. Humic acid also controlled the root-knot nematode *Meloidogyne incognita*, *in vitro* and *in situ*, and promoted banana growth (Seenivasan and Senthilnathan, 2018).

## 7.4 Algal and botanical extracts

Some algal and botanical extracts have both biostimulatory and insecticidal properties. It can be difficult to distinguish their roles in certain situations, but some examples will be discussed where biostimulants have an impact on arthropods. An earlier report by Stephenson (1966) indicated the indirect

impact of a hydrolyzed seaweed product on aphids, red spider mites, and plant diseases, through increased plant resistance. Eckol, a phenolic extract of the brown seaweed *Ecklonia maxima*, increased various growth parameters, enzymatic activity, chlorophyll, carotenoid, and proline content, and reduced aphid infestations (Rengasamy et al., 2016). Increased myrosinase activity was responsible for the population reduction in the cabbage aphid *Brevicoryne brassicae*. González-Castro et al. (2019) found repellent and insecticidal properties of the extracts of the macroalgae *Caulerpa sertularioides*, *Laurencia johnstonii*, and *Sargassum horridum* against the Asian citrus psyllid *Diaphorina citri*.

Juglone, a quinoid compound extracted from walnut trees, has biostimulatory, algacidal, bactericidal, fungicidal, insecticidal, and herbicidal properties (Islam and Widhalm, 2020). Azadirachtin, which is commonly used as an insecticide and insect growth regulator, also has a biostimulatory effect (Dara, 2021).

#### 7.5 Silicates and phosphites

Silicates applied as fertilizers benefit plants by strengthening the tissues and acting as a mechanical barrier, and also trigger immune responses. Numerous studies have demonstrated a significant reduction in arthropod populations and their damage from silicate applications (Laing et al., 2006; Moraes et al., 2019). Some of the mechanisms of resistance offered to plants by silicon include the reduced penetration of plant tissues, increased wearing of mouthparts, reduced digestibility and palatability of plant material, and damage to the insect's midgut epithelium (Alhousari and Greger, 2018). Additionally, soluble silicon in plants also attracts natural enemies and enhances the biocontrol of pests (Bakhat et al., 2018). For example, silicic acid application reduced damage by the cucurbit beetle Diabrotica speciosa, made the plants less preferable, and increased plant growth in potato without affecting their natural enemies (De Assis et al., 2012). Potassium silicate supplementation for chrysanthemum reduced 54% of mining by the American serpentine leafminer Liromyza trifolii, although it also reduced the plant height by 10% (Klittich and Parrella, 2014). Silicon application through irrigation significantly reduced the numbers of twospotted spider mites (Tetranychus urticae) and the levels of powdery mildew (Podosphaera aphanis) in strawberry (Liu et al., 2020a).

Similar to silicates, phosphites also have an impact on pest suppression. Foliar application of potassium phosphite in potato reduced the number of potato tuber moth (*Phthorimaea operculella*) larvae by more than 50%, without affecting parasitoid populations (Mulugeta et al., 2018). Similarly, foliar application of a phosphite fungicide containing sodium, potassium, and ammonium phosphites reduced the survival and prolonged the development

of the Colorado potato beetle (*Leptinotarsa decemlineata*), and reduced the foliar damage while controlling diseases (Patterson and Alyokhin, 2014).

#### 7.6 Chitosan

Chitosan has the potential to control multiple pests including arthropods. Derivatives of chitosan had significant insecticidal activity against S. littoralis (Rabea et al., 2005), B. brassicae, the oleander aphid Aphis nerii, and the wheat aphid Schizaphid graminum (Sahebzadeh et al., 2017) in laboratory assays. Colloidal chitosan from seafood waste showed significant antifeedant activity against S. frugiperda and reduced the feeding damage (Moorthy et al., 2021). Zhang and Tan (2003) reported insecticidal activity of chitosan against the cotton bollworm (Helicoverpa armigera), P. xylostella, S. exigua, the bird-cherry oat aphid (Rhopalosiphum padi), A. gossypii, the grain aphid (Sitobion avenae), the green peach aphid (Myzus persicae), the mealy plum aphid (Hyalopterus pruni), and the rose-grain aphid (Metopolophium dirhodum), with very high mortality in some species. Alfy et al. (2020) found that chitosan nanoparticles were effective against S. littoralis, the desert locust Locusta migratoria, and M. incognita. Fan et al. (2020) also reported that fluorinated derivatives of chitosan effectively killed the second-instar larvae and inhibited the egg hatch of M. incognita.

Chitosan is not only used for its direct and indirect effect on pest suppression. Its role as a nanocarrier of pesticides for safe and targeted delivery is also explored (Maluin and Hussein, 2020). For example, chitosan helped improve RNA interference efficacy against *S. frugiperda* (Gurusamy et al., 2020). Chitosan was also used for the controlled release of spinosad for increased environmental durability (Li et al., 2020). The monoterpenes carvacrol and linalool, nanoencapsulated in chitosan, increased the control efficacy against the pests *H. armigera* and *T. urticae* (Campos et al., 2018). In a different study, chitosan alone and as chitosan-reduced silver nanocrystals were detrimental to *H. armigera* (Murugan et al., 2021).

## 8 Tolerance to abiotic stresses

When plants undergo abiotic stresses such as salinity and drought, they are more vulnerable to pests and diseases. If biostimulants can improve plant growth under abiotic stresses, plants can use their resources to withstand biotic stresses. At the same time, plant stresses can also influence soil microbial populations (Liu et al., 2020b). For example, *G. mosseae* increased the salt tolerance of citrus seedlings with improved plant and root growth, photosynthesis, and ionic balance (Wu et al., 2010). *G. versiforme* stimulated plant growth and improved biomass of tangerine under water stress (Wu and

Xia, 2006). Improved nutrient absorption, higher accumulation of carbohydrates in roots and foliage, change in root plasticity, and higher antioxidative enzyme activity were among the factors that were found to impart salt and drought tolerance (Wu and Xia, 2006; He et al., 2007; Wu et al., 2010). Dual inoculation of *Rhizobium* sp. and *Enterococcus mundtii* increased seed germination, plant height, biomass, chlorophyll content, and nutrient uptake in mung bean growing in saline conditions (Kumawat et al., 2021).

Plant-derived protein hydrolysates improved plant growth, nutrient utilization, and yields of lettuce growing in pots under saline conditions (Lucini et al., 2015). The diameter of grapes was significantly improved, even under water-stressed conditions, when a protein hydrolysate was applied (Meggio et al., 2020). Although the differences were not significant, protein hydrolysates helped maintain the weight and number of fruit clusters under water stress.

Canellas et al. (2015) discussed the role of humic substances in regulating secondary metabolism and alleviating various stresses. Increasing the accumulation of phenolic compounds, improving osmoregulation, and regulating peroxidase activity are among the various influences of humic substances on plants. Similarly, macroalgae-based biostimulants also stimulate the production of plant defense mechanisms and other physiological responses that alleviate stress (Sharma et al., 2014). Silicate fertilizers increase the yield, quality, and shelf-life of grapes grown in calcareous gray desert soil (Zhang et al., 2017). Phosphites induce adaptive plant responses to various stresses and improve plant performance (Trejo-Téllez and Gómez-Merino, 2018). For example, under heat stress, phosphite induced the accumulation of stress proteins, reduced oxidative stress, and maintained photosynthetic efficiency in potato seedlings (Xi et al., 2020).

## 9 Multitrophic interactions and influencing factors

The effect of biostimulants can vary widely depending on various factors. In addition to neutral and positive effects, some biostimulant materials can negatively impact growth and yields (Nardi et al., 2016). AMF *Glomus caledonium, Glomus fasciculatum,* and *G. mosseae* increased the number of leaves per plant in the oxeye daisy *Leucanthemum vulgare* but influenced the levels of parasitism of the ragwort leafminer *Chromatomyia syngenesiae*, by *Diglyphus isaea* (Gange et al., 2003). Depending on the AMF species and their combinations, parasitism was higher, lower, or neutral in AMF-colonized plants. *Trichoderma* spp. that produce the volatile organic compound trichodiene, suppressed plant defenses and increased the emergence of *A. obtectus* and its damage to beans (Rodríguez-González et al., 2018). Application of glycine betaine did not improve strawberry fruit yields, but a product containing nutrients and humates improved yields by 11% (Dara, 2019d). Soy protein hydrolysate

improved strawberry yields when applied alone but not in combination with a product containing *T. harzianum* and *T. virens* (Dara, 2020). A product containing soybean and corn oils stunted strawberry plant growth and reduced yields, probably from higher application rates (Dara, unpublished data). Similarly, an animal-derived protein hydrolysate reduced the weight of daughter plants in strawberry (Lisiecka et al., 2011). Seedling growth of cucumber was inhibited but the single-fruit weight and concentration of certain nutrients in shoots increased with *G. versiforme* (Wang et al., 2008). Overapplication of silicates reduced flower yields and caused growth abnormalities in ornamentals (Kamenidou et al., 2008).

Certain practices and other agricultural inputs might not be compatible with some biostimulants, especially if they have live microorganisms and require special storage, handling, and application. Both the biostimulant industry and the grower community should be aware of the potential negative impacts of biostimulants and work with the researchers to determine rates, frequencies, and application strategies that are appropriate for their crop situation.

## 10 Strategies of using biostimulants

Although numerous studies have demonstrated the potential of biostimulants in improving crop growth, health, and yields, it is always advisable to consider IPM or integrated crop management approaches that take advantage of multiple strategies and reduce the reliance on one or a few options. Arthropod pests and plant pathogens continuously evolve and develop resistance to various agricultural inputs, especially those that have pesticidal properties. Since some biostimulants also have a certain level of pesticidal properties, a continuous monitoring of potential negative interactions in the environment and an adjustment of the strategies as appropriate are recommended. In general, biostimulants can be used as seed or transplant treatments, or they can be applied throughout the crop cycle as seed treatments, to the soil, or as foliar sprays (Fig. 2). Based on several studies discussed in this chapter, biostimulants improved tomato seed germination, root and shoot growth, general health, yield, quality, and nutritional value, suppressed various pests and diseases, improved biocontrol activity, and reduced the need for synthetic fertilizers and the potential use of pesticides (Fig. 3). Using tomato as a model plant, the importance of biostimulants can be demonstrated in crop production and crop protection. In general, biostimulant supplements can enhance the current agronomic practices and build plant resilience to various biotic and abiotic stresses.

Building soil health should be one of the priorities for sustainable agriculture, and biostimulants play a major role in this area by adding organic matter, beneficial microorganisms, nutrients, and other essentials to the soil.

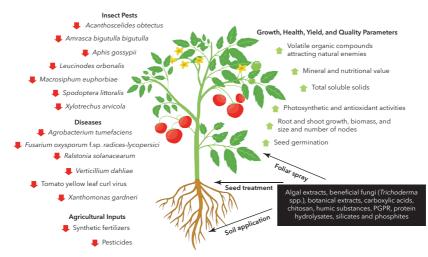


Figure 3 Tomato as a model crop for improving growth, health, yield, and parameters, and suppressing pests and diseases based on the various studies reviewed in this chapter.

It helps growers and crop care professionals to have a thorough knowledge of various biostimulants, their modes of action, interactions with other biotic and abiotic inputs and farming practices, and their indirect effect on pest and disease management. Biostimulants are part of a proactive strategy for maintaining plant health, while pesticides are curative measures to address specific issues. In other words, biostimulants are comparable to probiotics and vitamins, while pesticides are comparable to the medicines that humans take. Since biostimulants help improve nutrient uptake and trigger plant defenses against stressors, they contribute to the reduction of fertilizer and pesticide use and provide additional benefits beyond improved plant growth and yields. Modern-day agriculture can significantly benefit from incorporating biostimulant use as a good agricultural practice.

## 11 Challenges and future needs

Unlike the immediate effects of some fertilizer or pesticide inputs, the positive impact of biostimulants may not be as obvious and could easily be attributed to other crop production and protection practices. Additionally, the biological nature of several biostimulants also adds to the general misconception that biologicals are slow and less effective than synthetic agricultural inputs. The higher cost of some biostimulants and a lack of convincing applied research data are critical hurdles for the increased adaptation of biostimulants in many cropping systems. Based on the new IPM model and its emphasis on research, outreach, information management, and communication (Dara, 2019b), the

future needs for integrating biostimulants as a critical part of crop production and IPM are as follows:

- 1 Develop collaborations among researchers, biostimulant manufacturers, and growers for conducting applied research and continuously generating data from various crop and field situations.
- 2 Understand various interactions among biostimulants, other agricultural inputs, agricultural practices, plants, arthropods, and microorganisms, and develop appropriate storage, handling, and use strategies.
- 3 Incorporate biostimulants as a part of cultural control of IPM.
- 4 Effectively disseminate the research findings through outreach activities and encourage improved communication among key players for building confidence and refining use strategies.
- 5 Invest in research to improve their formulations and their efficacy while optimizing their cost.

## **12** Conclusion

This chapter, probably the first of its kind, provided an overview of various biostimulant categories and how they can be used in improving plant growth, health, and yields, especially as a part of an IPM program (Fig. 2). Depending on the strain, source, form, formulation, application rates, and other factors, some of the biostimulants might be commercially available as biopesticides or biofertilizers. While biostimulatory and biopesticidal properties overlap for some biostimulant active ingredients, the main objective was to present the biostimulatory properties and the indirect effect of various active ingredients on arthropod pests, plant pathogens, and in some cases, plant-parasitic nematodes. With a growing interest in biostimulants and their potential, as well as the continued demand for sustainable food production, biostimulant use is expected to increase further in the near future.

## 13 Where to look for further information

#### 13.1 Further reading

- Integrated pest and disease management in greenhouse crops (Springer) by Gullino et al. (2020).
- Biostimulants for sustainable crop production (Burleigh Dodds Science Publishing) by Youssef Rouphael et al. (2020).
- Biopesticides in organic farming: recent advances (Taylor and Francis) by Awasthi (2021).

#### 13.2 Key journals/conferences

- eJournal of Entomology and Biologicals (https://ucanr.edu/JEB).
- Frontiers in Sustainable Food Systems (https://www.frontiersin.org/journal s/sustainable-food-systems).
- Journal of Economic Entomology (https://academic.oup.com/JEE).
- Entomological Society of America annual meetings (https://entsoc.org).

## **14 References**

- Aguín, O., Mansilla, P., Vilariño, A. and Sainz, M. J. (2004). Effects of mycorrhizal inoculation on root morphology and nursery production of three grapevine rootstocks. *Am. J. Enol. Vitic.* 55: 108–111.
- Alfy, H., Ghareeb, R. Y., Soltan, E. and Farag, D. A. (2020). Impact of chitosan nanoparticles as insecticide and nematicide against *Spodoptera littoralis*, *Locusta migratoria*, and *Meloidogyne incognita*. *Plant Cell Biotehnol. Mol. Biol.* 21: 126-140.
- Alhousari, F. and Greger, M. (2018). Silicon and mechanisms of plant resistance to insect pests. *Plants (Basel)* 7(2): 33.
- Ali, A., Noh, N. M. and Mustafa, M. A. (2015). Antimicrobial activity of chitosan enriched with lemongrass oil against anthracnose of bell pepper. *Food Pack. Shelf Life* 3: 56-61.
- Ali, M., Cheng, Z.-H., Hayat, S., Ahmad, H., Ghani, M. I. and Liu, T. (2019). Foliar spraying of aqueous garlic bulb extract stimulates growth and antioxidant enzyme activity in eggplant (*Solanum melongena* L.). J. Integr. Agric. 18(5): 1001-1013.
- Amira, M., Lopez, D., Triki Mohamed, A., Khouaja, A., Chaar, H., Fumanal, B., Gousset-Dupont, A., Bonhomme, L., Label, P., Goupil, P., Ribeiro, S., Pujade-Renaud, V., Julien, J., Auguin, D. and Venisse, J. (2017). Beneficial effect of *Trichoderma harzianum* strain Ths97 in biocontrolling *Fusarium solani* causal agent of root rot disease in olive trees. *Biological Control* 110: 70-78.
- Araujo, I. B., Peruch, L. A. M. and Stadnik, M. J. (2012). Effect of seaweed extract and silicate clay on Alternaria leaf spot and on the yield of green onion (*Allium fistulosum* L.). *Trop. Plant Pathol.* 37(5): 363–367.
- Awasthi, L. P. (Ed) 2021. Biopesticides in organic farming: recent advances. CRC Press, 404 pages. Available at: https://www.google.com/books/edition/Biopesticides\_in \_Organic\_Farming/1Y8kEAAAQBAJ?hl=en&gbpv=0&kptab=overview.
- Aziz, A., Trotel-Aziz, P., Dhuicq, L., Jeandet, P., Couderchet, M. and Vernet, G. (2006). Chitosan oligomers and copper sulfate induce grapevine defense reactions and resistance to gray mold and downy mildew. *Phytopathology* 96(11): 1188-1194.
- Backer, R., Rokem, J. S., Ilangumaran, G., Lamont, J., Praslickova, D., Picci, E., Subramanian, S. and Smith, D. L. (2018). Plant growth-promoting rhizobacteria: context, mechanisms of action, and roadmap to commercialization of biostimulants for sustainable agriculture. *Front. Plant Sci.* 9: 1473.
- Baglieri, A., Cadili, V., Monterumici, C. M., Gennari, M., Tabasso, S., Montoneri, E., Nardi, S. and Negre, M. (2014). Fertilization of bean plants with tomato plants hydrolysates. Effect on biomass production, chlorophyll content and N assimilation. *Sci. Hort*. 176: 194-199.

- Bakhat, H. F., Bibi, N., Zia, Z., Abbas, S., Hammd, H. M., Fahad, S., Ashraf, M. R., Shah, G. M., Rabbani, F. and Saeed, S. (2018). Silicon mitigates biotic stresses in crop plants: a review. Crop Protec. 104: 21-34.
- Bakker, M. G., Manter, D. K., Sheflin, A. M., Weir, T. L. and Vivanco, J. M. (2012). Harnessing the rhizosphere microbiome through plant breeding and agricultural management. *Plant Soil* 360(1-2): 1-13.
- Barone, V., Baglieri, A., Stevanato, P., Broccanello, C., Bertoldo, G., Bertaggia, M., Cagnin, M., Pizzeghello, D., Moliterni, V. M. C., Mandolino, G., Fornasier, F., Squartini, A., Nardi, S. and Concheri, G. (2018). Root morphological and molecular responses induced by microalgae extracts in sugar beet (*Beta vulgaris* L.). *J. Appl. Phycol.* 30: 1061-1071.
- Battacharyya, B., Babgohari, M. Z., Rathor, P. and Prithiviraj, B. (2015). Seaweed extracts as biostimulants in horticulture. *Sci. Hort.* 196: 39-48.
- Bell, K., Narajno-Guevara, N., dos Santos, R. C. D., Meadow, R. and Bento, J. M. S. (2020). Predatory earwigs are attracted by herbivore-induced plant volatiles linked with plant growth-promoting rhizobacteria. *Insects* 11(5): 271.
- Bertsch, F., Ramírez, F. and Henríquez, C. 2009. Evaluación del fosfito como fuentefertilizante de fósforo vía radical y foliar. *Agron. Costarricense* 33: 249-265.
- Cameron, D. D., Neal, A. L., van Wees, S. C. M. and Ton, J. (2013). Mycorrhiza-induced resistance: more than the sum of its parts? *Tren. Pl. Sci.* 18(10): 539-545.
- Campos, E. V. R., Proença, P. L. F., Oliveira, J. L., Pereira, A. E. S., Ribeiro, L. N. M., Fernandes,
  F. O., Gonçalves, K. C., Polanczyk, R. A., Pasquoto-Stigliani, T., Lima, R., Melville,
  C. C., Della Vechia, J. F., Andrade, D. J. and Fraceto, L. F. (2018). Carvacrol and
  linalool co-loaded in β-cyclodextrin-grafted chitosan nanoparticles as sustainable
  biopesticide aiming pest control. *Sci. Rep.* 8(1): 7623.
- Canellas, L. P., Olivares, F. L., Aguiar, N. O., Jones, D. L., Nebbioso, A., Mazzei, P. and Piccolo, A. (2015). Humic and fulvic acids as biostimulants in horticulture. *Sci. Hort*. 196: 15-27.
- Cappelletti, M., Perazzolli, M., Nesler, A., Giovannini, O. and Pertot, I. (2017). The effect of hydrolysis and protein source on the efficacy of protein hydrolysates as plant resistant inducers against powdery mildew. *J. Bioprocess. Biotech.* 7: 5.
- Chaoui, H., Edwards, C. A., Brickner, A., Lee, S. S. and Arancon, N. Q. (2002). Suppression of the plant diseases, *Pythium* (damping-off), *Rhizoctonia* (root rot) and *Verticillium* (wilt) by vermicomposts. *Brighton Crop Prot. Conf. Pests and Dis.* 2: 711-716.
- Colla, G., Cardarelli, M., Bonini, P. and Rouphael, Y. (2017). Foliar applications of protein hydrolysate, plant and seaweed extracts increase yield but differentially modulate fruit quality of greenhouse tomato. *HortScience* 52: 1214–1220.
- Colla, G., Nardi, S., Cardarelli, M., Ertani, A., Lucini, L., Canaguier, R. and Rouphale, Y. (2015). Protein hydrolysates as biostimulants in horticulture. *Sci. Hort*. 196: 28-38.
- Coppola, M., Cascone, P., Chiusano, M. L., Colantuono, C., Lorito, M., Pennacchio, F., Rao, R., Woo, S. L., Guerrieri, E. and Digilio, M. C. (2017). *Trichoderma harzianum* enhances tomato indirect defense against aphids. *Insect Sci.* 24(6): 1025-1033.
- Coppola, M., Cascone, P., Lelio, I. D., Woo, S. L., Lorito, M., Rao, R., Pennacchio, F., Guerrieri, E. and Digilio, M. C. (2019). *Trichoderma atroviride* P1 colonization of tomato plants enhances both direct and indirect defense barriers against insects. *Front. Physiol.* 10: 813.
- Dara, S. K. (2017). Utilizing recycled food waste-based liquid compost: sustainable nutrient management option in strawberry. *Veg. West* 21: 16-17.

- Dara, S. K. (2019a). Non-entomopathogenic roles of entomopathogenic fungi in promoting plant health and growth. *Insects* 10(9): 277.
- Dara, S. K. (2019b). The new integrated pest management paradigm for the modern age. J. IPM 10(1): 1-9.
- Dara, S. K. (2019c). Effect of microbial and botanical biostimulants with nutrients on tomato yield. *CAPCA Adv.* 22: 40-45.
- Dara, S. K. (2019d). Evaluating the efficacy of anti-stress supplements on strawberry yield and quality. UCANR eJ. Entomol. Biol. Available at: https://ucanr.edu/blogs/blogco re/postdetail.cfm?postnum=31044.
- Dara, S. K. (2020). Improving strawberry yields with biostimulants and nutrient supplements: a 2019-2020 study. UCANR eJ. Entomol. Biol. Available at: https://uc anr.edu/blogs/blogcore/postdetail.cfm?postnum=43631.
- Dara, S. K. (2021). Impact of microbial, botanical, and organic acid-based biostimulants on strawberry health and yield. *CAPCA Adv.* 24: 30-35.
- Dara, S. K. and Lewis, E. (2019). Evaluating biostimulant and nutrient inputs to improve tomato yields and crop health. *Prog. Crop Consult.* 4: 38-42.
- Dara, S. K. and Peck, D. (2018). Evaluation of additive, soil amendment, and biostimulant products in Santa Maria strawberry. *CAPCA Adv.* 21: 44-50.
- De Assis, F. A., Morae, J. C., Silveira, L. C. P., Francoso, J., Nascimento, A. M. and Antunes, C. S. (2012). Inducers of resistance in potato and its effects on defoliators and predatory insects. *Rev. Colomb. Entomol.* 38: 30-34.
- De Oliveira Pedro, R., Takaki, M., Gorayeb, T. C., Del Bianchi, V. L., Thomeo, J. C., Tiera, M. J. and de Oliveira Tiera, V. A. (2013). Synthesis, characterization and antifungal activity of quaternary derivatives of chitosan on *Aspergillus flavus*. *Microbiol. Res.* 168(1): 50-55.
- De Pascale, S., Rouphael, Y. and Colla, G. (2017). Plant biostimulants: innovative tool for enhancing plant nutrition in organic farming. *Eur. J. Hortic. Sci.* 82: 277–285.
- Del Rosario Cappellari, L., Chiappero, J., Palermo, T. B., Giordano, W. and Banchio, E. (2020). Impact of soil rhizobacteria inoculation and leaf-chewing insect herbivory on *Mentha piperita* leaf secondary metabolites. J. Chem. Ecol. 46(7): 619-630.
- Deniau, M. G., Bonafos, R., Chovelon, M., Parvaud, C.-E., Furet, A., Bertrand, C. and Marchand, P. A. (2019). Willow extract (*Salix* cortex), a basic substance of agronomical interests. *Int. J. Bio-Res. Stress Manag.* 10(4): 408–418.
- Disi, J., Simmons, J. and Zebelo, S. (2019). Plant growth-promoting rhizobacteriainduced defense against insect herbivores. In: Maheshwari, D. and Dheeman, S. (Eds) Field Crops: Sustainable Management by PGPR. Sustainable Development and Biodiversity (vol. 123): 385-410, Springer, Cham.
- Elarroussi, H., Elmernissi, N., Benhima, R., El Kadmiri, I. M., Bendaou, N., Smouni, A. and Wahby, I. (2016). Microalgae polysaccharides a promising plant growth biostimulant. *J. Algal Biomass Utln* 7: 55-63.
- El Hadrami, A., Adam, L. R., El Hadrami, I. and Daayf, F. (2010). Chitosan in plant protection. *Mar. Drugs* 8(4): 968-987.
- El Modafar, C., Elgadda, M., El Boutachfaiti, R., Abouraicha, E., Zehhar, N., Petit, E., El Aloui-Talibi, Z., Courtois, B. and Coutois, J. (2012). Induction of natural defence accompanied by salicyclic acid-dependant systemic acquired resistance in tomato seedlings in response to bioelicitors isolated from green algae. *Sci. Hort.* 138: 55-63.
- Esitken, A., Karlidag, H., Ercisli, S. and Sahin, F. (2002). Effects of foliar application of Bacillus subtilis Osu-142 on the yield, growth and control of shot-hole disease (Coryneum blight) of apricot. Gartenbauwiss 67: 139-142.

- Estrada-Ortiz, E., Trejo-Téllez, L. I., Gómez-Merino, F. C., Núñez-Escobar, R. and Sandoval-Villa, M. (2012). Phosphite on growth and fruit quality in strawberry. *Acta Hortic.* 947(947): 277-282.
- Fan, Z., Qin, Y., Liu, S., Xing, R., Yu, H. and Li, P. (2020). Chitosan oligosaccharide fluorinated derivative control root-knot nematode (*meloidogyne incognita*) disease based on the multi-efficacy strategy. *Mar. Drugs* 18(5): 273.
- Featonby-Smith, B. C. and van Staden, J. (1987). Effect of seaweed concentrate on yield and seed quality of *Arachis hypogaea*. S. Afr. J. Bot. 53(3): 190-193.
- Fiorentino, N., Ventorino, V., Woo, S. L., Pepe, O., De Rosa, A., Gioia, L., Romano, I., Lombardi, N., Napolitano, M., Colla, G. and Rouphael, Y. (2018). *Trichoderma*based biostimulants modulate rhizosphere microbial populations and improve N uptake efficiency, yield, and nutritional quality of leafy vegetables. *Front. Plant Sci.* 9: 743.
- Fornes, F., Sánchez-Perales, M. and Guardiola, J. L. (2002). Effect of a seaweed extract on the productivity of 'de Nules' clementine mandarin and Navelina orange. *Botanica Marina* 45(5): 486-489.
- Föster, H., Adaskaveg, J. E., Kim, D. H. and Stanghellini, M. E. (1998). Effect of phosphite on tomato and pepper plants and on susceptibility of pepper to Phytophthora root and crown rot in hydroponic culture. *Plant Dis.* 82(10): 1165–1170.
- Gange, A. C., Brown, V. K. and Aplin, D. M. (2003). Multitrophic links between arbuscular mycorrhizal fungi and insect parasitoids. *Ecol. Lett.* 6(12): 1051-1055.
- Gemin, L. G., Mógor, Á. F., Amatussi, J. D. O. and Mógor, G. (2019). Microalgae associated to humic acid as a novel biostimulant improving onion growth and yield. *Sci. Hort.* 256: 108560.
- Ghosh, S. K. and Pal, S. (2016). Entomopathogenic potential of *Trichoderma longibrachiatum* and its comparative evaluation with Malathion against the insect pest *Leucinodes Orbonalis*. *Environ. Monit. Assess.* 188(1): 37.
- Glick, B. R. (2012). Plant growth-promoting bacteria: mechanisms and applications. *Scintif* 2012: 963401.
- Glinicki, R., Sas-Paszt, L. and Jadczuk-Tobjasz, E. (2010). The effect of plant stimulant/ fertilizer Resistim on growth and development of strawberry plants. *J. Fruit Ornam Plant Res.* 18: 111-124.
- Godlewska, K., Biesiada, A., Michalak, I. and Pcyga, P. (2019). The effect of plant-derived biostimulants on white head cabbage seedlings grown under controlled conditions. *Sustainability* 11(19): 5317.
- Gómez-Merino, F. C. and Trejo-Téllez, L. I. (2015). Biostimulant activity of phosphite in horticulture. *Sci. Hort*. 196: 82–90.
- González-Castro, A. L., Muñoz-Ochoa, M., Hernández-Carmona, G. and López-Vivas, J.
   M. (2019). Evaluation of seaweed extracts for the control of the Asian citrus psyllid Diaphorina citri. J. Appl. Phycol. 31(6): 3815-3821.
- Gullino, M. L., Albajes, R., & Nicot, P. C. (Eds.). (2020). Integrated Pest and Disease Management in Greenhouse Crops (vol. 9). Springer Nature.
- Gurusamy, D., Mogilicherla, K. and Palli, S. R. (2020). Chitosan nanoparticles help doublestranded RNA escape from endosomes and improve RNA interference in the fall armyworm, *Spodoptera frugiperda. Arch. Insect Biochem. Physiol.* 104(4): e21677.
- He, Z. Q., He, C. X., Zhang, Z. B., Zou, Z. R. and Wang, H. S. (2007). Changes in antioxidative enzymes and cell membrane osmosis in tomato colonized by arbuscular mycorrhizae under NaCl stress. *Colloids Surf. B. Bioint.* 59: 128-133.

- Hena, S., Gutierrez, L. and Croué, J. P. (2021). Removal of pharmaceutical and personal care products (PPCPs) from wastewater using microalgae: a review. J. Hazard. Mater. 403: 124041.
- Hernandez, O. L., Calderín, A., Huelva, R., Martínez-Balmori, D., Guridi, F., Aguiar, N. O., Olivares, F. L. and Canellas, L. P. (2015). Humic substances from vermicompost enhance urban lettuce production. *Agron. Sustain. Dev.* 35(1): 225-232.
- Huang, X. Q., You, Z. Y., Luo, Y., Yang, C. J., Ren, J., Liu, Y. L., Wei, G. J., Dong, P. and Ren, M. Z. (2021). Antifungal activity of chitosan against *Phytophthora infestans*, the pathogen of potato late blight. *Int. J. Biol. Macromol.* 166: 1365–1376.
- Islam, A. K. M. M. and Widhalm, J. R. (2020). Agricultural uses of juglone: opportunities and challenges. *Agronomy* 10(10): 1500.
- Jail, N. Gd, Luiz, C., Rocha Neto, A. Cd and Di Piero, R. M. (2014). High-density chitosan reduces the severity of bacterial spot and activates the defense mechanisms of tomato plants. *Trop. Plant Pathol.* 39(6): 434-441.
- Janvier, C., Villenueuve, F., Alabouvette, C., Edel-Hermann, V., Mateille, T. and Steinberg, C. (2007). Soil health through soil disease suppression: which strategy from descriptors to indicators? *Soil Biol. Biochem.* 39(1): 1-23.
- Jayaraj, J., Wan, A., Rahman, M. and Punja, Z. K. (2008). Seaweed extract reduces foliar fungal diseases on carrot. *Crop Prot.* 27(10): 1360-1366.
- Ji, S., Liu, Z., Liu, B., Wang, Y. and Wang, J. (2020). The effect of *Trichoderma* biofertilizer on the quality of flowering Chinese cabbage and the soil environment. *Sci. Hort*. 262: 109069.
- Jindo, K., Olivares, F. L., Malcher, D. J. D. P., Sánchez-Monedero, M. A., Kempenaar, C. and Canellas, L. P. (2020). From lab to field: role of humic substances under open-field and greenhouse conditions as biostimulant and biocontrol agent. *Front. Plant Sci.* 11: 426.
- Joshi, R., Singh, J. and Vig, A. P. 2015. Vermicompost as an effective organic fertilizer and biocontrol agent: effect on growth, yield and quality of plants. *Rev. Environ. Sci. Biotechnol.* 14: 137-159.
- Jung, S. C., Martinez-Medina, A., Lopez-Raez, J. A. and Pozo, M. J. (2012). Mycorrhizainduced resistance and priming of plant defenses. J. Chem. Ecol. 38(6): 651-664.
- Kamenidou, S., Cavins, T. J. and Marek, S. (2008). Silicon supplements affect horticultural traits of greenhouse-produced ornamental sunflowers. *Hort. Sci.* 43(1): 236-239.
- Kamenidou, S., Cavins, T. J. and Marek, S. (2010). Silicon supplements affect floricultural quality traits and elemental nutrient concentrations of greenhouse produced gerbera. *Sci. Hort.* 123(3): 390–394.
- Kiki, C., Rashid, A., Wang, Y., Li, Y., Zeng, Q., Yu, C. P. and Sun, Q. (2019). Dissipation of antibiotics by microalgae: kinetics, identification of transformation products and pathways. *J. Hazard. Mater.* 387: 121985.
- Kim, M. J., Radhakrishnan, R., Kang, S. M., You, Y. H., Jeong, E. J., Kim, J. G. and Lee, I. J. (2017). Plant growth promoting effect of *Bacillus amyloliquefaciens* H-2-5 on crop plants and influence on physiological changes in soybean under soil salinity. *Physiol. Mol. Biol. Plants* 23(3): 571–580.
- Klittich, D. and Parrella, M. (2014). Silicon supplementation: effects on chrysanthemum growth, leafminer populations, and parasitism. Proceedings of the IOBC/WPRS Working Group "Integrated Control in Protected Crops, Temperate Climate", Gent, Belgium, September 14-18, 2014 102: 117-119.
- Kumawat, K. C., Sharma, P., Nagpal, S., Gupta, R. K., Sirari, A., Nair, R. M., Bindumadhava, H. and Singh, S. (2021). Dual microbial inoculation, a game changer?–bacterial

biostimulants with multifunctional growth promoting traits to mitigate salinity stress in spring mungbean. *Front. Microbiol.* 11: 3491.

- Lachhab, N., Sanzani, S. M., Adrian, M., Chiltz, A., Balacey, S., Boselli, M., Ippolito, A. and Poinssot, B. (2014). Soybean and casein hydrolysates induce grapevine immune responses and resistance against *Plasmopara viticola*. Front. Plant Sci. 5: 716.
- Lafontaine, P. J. and Benhanou, N. (1996). Chitosan treatment: an emerging strategy for enhancing resistance of greenhouse tomato plants to infection by *Fusarium oxysporum* f. sp. *radices-lycopersici. Biocon. Sci. Technol.* 6(1): 111-124.
- Laing, M. D., Gatarayiha, M. C. and Adandonon, A. (2006). Silicon use for pest control in agriculture: a review. *Proc. South Afr. Sugar Tech.* 80: 278-286.
- Lal, R. (2016). Soil health and carbon management. Food Ener. Sec. 5(4): 212-222.
- Larkin, R. P. (2008). Relative effects of biological amendments and crop rotations on soil microbial communities and soilborne diseases of potato. *Soil Biol. Biochem.* 40(6): 1341-1351.
- Li, G. B., Wang, J. and Kong, X. P. (2020). Coprecipitation-based synchronous pesticide encapsulation with chitosan for controlled Spinosad release. *Carbohydr. Polym.* 249: 116865.
- Li, H., Ding, X., Wang, C., Ke, H., Wu, Z., WAng, Y., Liu, H. and Guo, J. (2016). Control of tomato yellow leaf curl virus disease by *Enterobacter asburiae* BQ9 as a result of priming plant resistance in tomatoes. *Turk. J. Biol.* 40: 150-159.
- Li, Y. C., Bi, Y., Ge, Y. H., Sun, X. J. and Wang, Y. (2009). Antifungal activity of sodium silicate on *Fusarium sulphureum* and its effect on dry rot of potato tubers. *J. Food Sci.* 74(5): M213-M218.
- Liang, Y. C., Sun, W. C., Si, J. and Römheld, V. (2005). Effects of foliar- and root-applied silicon on the enhancement of induced resistance to powdery mildew in *Cucumis* sativus. Plant Pathol. 54(5): 678-685.
- Lisiecka, J., Knaflewski, M., Spizewski, T., Fraszczak, B., Kaluzewicz, A. and Krazesinski, W. (2011). The effect of animal protein hydrolysate on quantity and quality of strawberry daughter plants cv. 'Elsanta'. *Acta Sci. Pol. Hort. Cult.* 10: 31-40.
- Liu, B., Davies, K. and Hall, A. (2020a). Silicon builds resilience in strawberry plants against both strawberry powdery mildew *Podosphaera aphanis* and two-spotted spider mites *Tetranychus urticae*. *PLoS ONE* 15(12): e0241151.
- Liu, H., Brettel, L. E., Qiu, Z. and Singh, B. K. (2020b). Microbiome-mediated stress resistance in plants. *Trends Pl. Sci.* 25(8): 733-743.
- Lucini, L., Rouphael, Y., Cardarelli, M., Canaguier, R., Kumar, P. and Colla, G. (2015). The effect of a plant-derived biostimulant on metabolic profiling and crop performance of lettuce grown under saline conditions. *Sci. Hort.* 182: 124–133.
- Maluin, F. N. and Hussein, M. Z. (2020). Chitosan-based agronanochemicals as a sustainable alternative in crop protection. *Molecules* 25(7): 1611.
- Meggio, F., Trevisan, S., Manoli, A., Ruperti, B. and Quaggiotti, S. (2020). Systematic investigation of the effects of a novel protein hydrolysate on the growth, physiological parameters, fruit development and yield of grapevine (*Vitis vinifera* L., cv Sauvignon Blanc) under water stress conditions. *Agronomy* 10(11): 1785.
- Meng, D., Garba, B., Ren, Y., Yao, M., Xia, X., Li, M.. and Wang, Y. (2020). Antifungal activity of chitosan against Aspergillus ochraceus and its possible mechanisms of action. Int. J. Biol. Macromol. 158: 1063-1070.
- Molla, A. H., Manjurul Haque, M., Amdadul Haque, M. and Ilias, G. N. M. (2012). *Trichoderma*-enriched biofertilizer enhances production and nutritional quality of

tomato (*Lycopersicon esculentum* Mill.) and minimizes NPK fertilizer use. *Agri. Res.* 1(3): 265-272.

- Moorthy, A. V., Shanthi, M., Chinnaiah, C. and Senthil, K. (2021). Synthesis of colloidal chitosan from sea waste and its antifeedant effect against maize fall armyworm *Spodoptera frugiperda* (J. E. Smith). *J. Entomol. Zool. Stud.* 9(1): 418-421.
- Moraes, J. C., Assis, F. A. and Assis, G. A. (2019). Use of silicon as resistance factor for plants against insect pests. In: Souza, B., Vázquez, L. and Marucci, R. (Eds) Natural Enemies of Insect Pests in Neotropical Agroecosystems: 497-508, Springer, Cham.
- Mulugeta, T., Mulatu, B., Tekie, H., Yesuf, M., Andreasson, E. and Alexandersson, E. (2018). Phosphite alters the behaviroal response of potato tuber moth (*Phthorimaea* operculella) to field-grown potato. *Pest Manag. Sci.* 75(3): 616-621.
- Murugan, K., Wang, L., Anitha, J., Dinesh, D., Amruthavalli, P., Vasanthakumaran, M., Paulpandi, M. and Hwang, J.-S. (2021). Insecticidal effect of chitosan reduced silver nanocrystals against filarial vector, *Culex quinquefasciatus* and cotton bollworm, *Helicoverpa armigera*. Adv. Nano-Fert. Nano-Pest Agric. 2021: 469-486.
- Nadhirawaty, R. and Titah, H. S. (2019). Simultaneous bioaugmentation and biostimulation to remediate soil contaminated by ship dismantling in Bangkalan District, Indonesia. *J. Health Pollut.* 9(24): 191212.
- Nandeeshkumar, P., Sudisha, J., Ramachandra, K. K., Prakash, H. S., Niranjana, S. R. and Shekar, S. H. (2008). Chitosan induced resistance to downy mildew in sunflower caused by *Plasmopara halstedii*. *Physiol. Mol. Plant Pathol.* 72(4-6): 188-194.
- Nardi, S., Pizzeghello, D., Muscolo, A. and Vianello, A. (2002). Physiological effects of humic substances on higher plants. *Soil Biol. Biochem.* 34(11): 1527-1536.
- Nardi, S., Pizzeghello, D., Schiavon, M. and Ertani, A. (2016). Plant biostimulants: physiological responses induced by protein hydrolyzed-based products and humic substances in plant metabolism. *Sci. Agric.* 73(1): 18-23.
- Nawaz, A., Gogi, M. D., Naveed, M., Arshad, M., Sufyan, M., Binyameen, M., Islam, S. U., Waseem, M., Ayyub, M. B., Arif, M. J. and Ali, H. (2020). In vivo and in vitro assessment of *Trichoderma* species and *Bacillus thuringiensis* integration to mitigate insect pests of brinjal (*Solanum melongena* L.). *Egypt. J. Biol. Pest Control* 30(1): 60.
- Nolla, A., Korndörfer, G. H. and Coelho, L. (2006). Efficiency of calcium silicate and carbonate in soybean disease control. *J. Plant Nutr.* 29(11): 2049-2061.
- Olivares, F. L., Busato, J. G., de Paula, A. M., da Silva Lima, L., Aguiar, N. O. and Canellas, L. P. (2017). Plant growth promoting bacteria and humic substances: crop promotion and mechanisms of action. *Chem. Biol. Technol. Agric.* 4(1): 30.
- Omara, A. E.-D., Hauka, F., Afify, A., El-Din, M. N. and Kassem, M. (2017). The role of some PGPR strains to biocontrol *Rhizoctonia solani* in soybean and enhancement the growth dynamics and seed yield. *Env. Biodiver. Soil Sec.* 1: 47-59.
- Pangesti, N., Weldegergis, B. T., Langendorf, B., van Loon, J. J., Dicke, M. and Pineda, A. (2015). Rhizobacterial colonization of roots modulates plant volatile emission and enhances the attraction of a parasitoid wasp to host-infested plants. *Oecologia* 178(4): 1169-1180.
- Pastor, V., Balmer, A., Gamir, J., Flors, V. and Mauch-Mani, B. (2014). Preparing to fight back: generation and storage of priming compounds. *Front. Plant Sci.* 5: 295.
- Patterson, M. and Alyokhin, A. (2014). Survival and development of Colorado potato beetles on potatoes treated with phosphite. *Crop Protec*. 61: 38-42.

- Paula Júnior, T. J., F. Vieira, R., Teixeira, H. and Eustáquio S. Carneiro, J. (2009). Foliar application of calcium chloride and calcium silicate decreases white mold intensity on dry beans. *Trop. Plant Pathol.* 34(3): 171-174.
- Pedraza, R. O., Motok, J., Salazar, S. M., Ragout, A. L., Mentel, M. I., Tortora, M. L., Guerrero-Molina, M. F., Winik, B. C. and Díaz-Ricci, J. C. (2010). Growth-promotion of strawberry plants inoculated with Azospirillum brasilense. World J. Microbiol. Biotechnol. 26(2): 265-272.
- Perazzolli, M., Dagostin, S., Ferrari, A., Elad, Y. and Pertot, I. (2008). Induction of systemic resistance against *Plasmopara viticola* in grapevine by *Trichoderma harzianum* T39 and benzothiadiazole. *Biological Control* 47(2): 228-234.
- Perazzolli, M., Morett, M., Fontana, P., Ferrarini, A., Velasco, R., Moser, C., Delledonne, M. and Pertot, I. (2012). Downy mildew resistance induced by *Trichoderma harzianum* T39 in susceptible grapevines partially mimics transcriptional changes of resistant genotypes. *BMC Genom.* 13: 660.
- Peterson, R. K. D., Higley, L. G. and Pedigo, L. P. (2018). Whatever happened to IPM? Am. Entomol. 64(3): 146-150.
- Pichyangkura, R. and Chadchawan, S. (2015). Biostimulant activity of chitosan in horticulture. *Sci. Hort.* 196: 49-65.
- Pindi, P. K. and Satyanarayana, S. D. V. (2012). Liquid microbial consortium-a potential tool for sustainable soil health. J. Biofert Biopest 3: 4.
- Poveda, J., Hermosa, R., Monte, E. and Nicolás, C. (2019). *Trichoderma harzianum* favours the access of arbuscular mycorrhizal fungi to non-host Brassicaceae roots and increase plant productivity. *Sci. Rep.* 9(1): 11650.
- Rabea, E. I., El Badawy, M. E., Rogge, T. M., Stevens, C. V., Höfte, M., Steurbaut, W. and Smagghe, G. (2005). Insecticidal and fungicidal activity of new synthesized chitosan derivatives. *Pest Manag. Sci.* 61(10): 951–960.
- Raghavendra, V. B., Lokesh, S. and Prakash, H. S. (2007). Dravya, a product of seaweed extract (*Sargassum wightii*), induces resistance in cotton against *Xanthomonas campestris* pv. *malvacearum*. *Phytoparasitica* 35(5): 442-449.
- Recep, K., Fikrettin, S., Erkol, D. and Cafer, E. (2009). Biological control of the potato dry rot caused by *Fusarium* species using PGPR strains. *Biological Control* 50(2): 194-198.
- Ren, J., Tong, J., Li, P., Huang, X., Dong, P. and Ren, M. (2021). Chitosan is an effective inhibitor against potato dry rot caused by *Fusarium oxysporum*. *Physiol. Mol. Plant Pathol.* 113: 101601.
- Rengasamy, K. R. R., Kulkarni, M. G., Pendota, S. C. and Van Staden, J. (2016). Enhancing growth, phytochemical constitutents and aphid resistance capacity in cabbage with foliar application of eckol–a biologically active phenolic molecule from brown seaweed. *New Biotechnol.* 33(2): 273-279.
- Rickard, D. A. 2000. Review of phosphorus acid and its salts as fertilizer materials. *J. Plant Nutr.* 23: 161–180.
- Rodríguez-González, Á., Casquero, P. A., Suárez-Villanueva, V., Carro-Huerga, G., Álvarez-García, S., Mayo-Prieto, S., Lorenzana, A., Cardoza, R. E. and Gutiérrez, S. (2018). Effect of trichodiene production by *Trichoderma harzianum* on *Acanthoscelides* obtectus. J. Stored Prod. Res. 77: 231-239.
- Rodríguez-González, Á., Mayo, S., González-López, Ó., Reinoso, B., Gutierrez, S. and Caquero, P. A. (2017). Inhibitory activity of *Beuaveria bassiana* and *Trichoderma* spp. on the insect pests *Xylotrechus Arvicola* (Coleoptera: Cerambycidae) and

Acanthoscelides obtectus (Coleoptera: Chrisomelidae: Bruchinae). Env. Mon. Assess. 189: 12.

- Ronga, D., Biazzi, E., Parati, K., Carminati, D., Carminati, E. and Tava, A. (2019). Microalgal biostimulants and biofertilisers in crop productions. *Agronomy* 9(4): 192.
- Rose, M. T., Patti, A. F., Little, K. R., Brown, A. L., Jackson, W. R. and Cavagnaro, T. R. (2014). A meta-analysis and review of plant-growth response to humi substances: practical implications for agriculture. *Adv. Agron.* 124: 37–89.
- Rouphael, Y., du Jardin, P., Brown, P., De Pascale, S. and Colla, G. (Eds) 2020. *Biostimulants for Sustainable Crop Production*. Burleigh Dodds Science Publishing. Available at: https://shop.bdspublishing.com/store/bds/detail/workgroup/3-190-84222.
- Rouphael, Y., Franken, P., Schneider, C., Schwarz, D., Giovannetti, M., Agnolucci, M., De Pascale, S., Bonini, P. and Colla, G. (2015). Arbuscular mycorrhizal fungi act as biostimulants in horticultural crops. *Sci. Hort.* 196: 91-108.
- Saberi, M., Sarpeleh, A., Askary, H. and Rafiei, F. (2013). Effects of wood vinegar and vermicompost combination in the control of *Verticillium dahliae* the causal agent of Verticillium wilt of greenhouse cucumber. *Appl. Entomol. Phytopathol.* 81: 0-0.
- Sahebzadeh, N., Ghaffari-Moghaddam, M. and Sabagh, S. K. (2017). Toxicity of N-alkyl derivatives of chitosan obtained from adult of *Chrotogonus trachypterus* (Orthoptera, Acrididae) against the wheat, cabbage and oleander aphid (Hemiptera: Aphididae) species. *Jordan J. Biol. Sci.* 10: 49–55.
- Salah, I. B., Aghrouss, S., Douira, A., Aissam, S., El Alaoui-Talibi, Z., Filali-Maltouf, A. and El Modafar, C. (2018). Seaweed polysaccharides as bio-elicitors of natural defenses in olive trees against Verticillium wilt of olive. J. Plant Int. 13: 248-255.
- Saravanakumar, D., Lavanya, N., Muthumeena, B., Raguchander, T., Suresh, S. and Samiyappan, R. (2008). *Pseudomonas fluorescens* enhances resistance and natural enemy population in rice plants against leaffolder pest. *J. Appl. Entomol.* 132(6): 469-479.
- Savvas, D. and Ntatsi, G. (2015). Biostimulant activity of silicon in horticulture. *Sci. Hort.* 196: 66-81.
- Seenivasan, N. and Senthilnathan, S. (2018). Effect of humic acid on *Meloidogyne* incognita (Kofoid & White) Chitwood infecting banana (*Musa* spp.). Int. J. Pest Manag. 64(2): 110-118.
- Selvaraj, A. and Thangavel, K. (2021). Arbuscular mycorrhizal fungi: potential plant protective agent against herbivorous insect and its importance in sustainable agriculture. In: Shrivastava, N., Mahajan, S. and Varma, A. (Eds) Symbiotic Soil Microorganisms. Soil Biology (vol. 60): 319-337, Springer, Cham.
- Seyedbagheri, M.-M. (2010). Influence of humic products on soil health and potato production. *Potato Res.* 53(4): 341-349.
- Shabbir, I., Samad, M. Y. A., Othman, R., Wong, M.-Y., Sulaiman, Z., Jaafar, N. M. and Bukhari, S. A. H. (2021). Evaluation of bioformulation of *Enterobacter* sp. UPMSSB7 and mycorrhizae with silicon for white root rot disease suppression and growth promotion of rubber seedlings inoculated with *Rigidoporus microporus*. *Biological Control* 152: 104467.
- Shahid, M., Duma, C., Silvestre, J. and Pinelli, E. (2012). Effect of fulvic acids on lead-induced oxidative stress to metal sensitive Vicia faba L. plant. Biol. Fertil. Soils 48(6): 689-697.
- Sharma, H. S. S., Fleming, C., Selby, C., Rao, J. R. and Martin, T. (2014). Plant biostimulants: a review on the processing of macroalgae and use of extracts for crop management to reduce abiotic and biotic stresses. *J. Appl. Phycol.* 26(1): 465-490.

- Shearer, B. L. and Fairman, R. G. (2007). Application of phosphite in a high-volume foliar spray delays and reduces the rate of mortality of four *Banksia* species infected with *Phytophthora cinnamomic*. *Austral*. *Plant Pathol*. 36(4): 358-368.
- Shukla, P. S., Borza, T., Critchley, A. T. and Prithiviraj, B. (2021). Seaweed-based compounds and products for sustainable protection against plant pathogens. *Mar. Drugs* 19(2): 59.
- Silva, O. C., Santos, H. A. A., Dalla Pria, M. and May-De Mio, L. L. (2011). Potassium phosphite for control of downy mildew of soybean. *Crop Prot.* 30(6): 598-604.
- Sousa, A. J. S., Silva, C. F. B., Sousa, J. S., Júnior, J. E. M., Freire, J. E. C., Sousa, B. L., Lobo, M. D. P., Monteiro-Moreira, A. C. O. and Grangeiro, T. B. (2019). A thermostable chitinase from the antagonistic *Chromobacterium violaceum* that inhibits the development of phytopathogenic fungi. *Enz. Microb. Technol.* 126: 50-61.
- Spinelli, F., Fiori, G., Noferini, M., Srocatti, M. and Costa, G. (2009). Perspectives on the use of a seaweed extract to moderate the negative effects of alternate bearing in apple trees. *J. Hortic. Sci. Biotechnol.* 84(6): 131-137.
- Stephenson, W. M. (1966). The effect of hydrolysed seaweed on certain plant pests and diseases. Proceedings of the Fifth Inter Seaweed Symposium, Halifax, Pergamon, August 25-28, 1965: 405-415.
- Supraja, K. V., Behera, B. and Balasubramanian, P. (2020). Efficacy of microalgal extracts as biostimulants through seed treatment and foliar spray for tomato cultivation. *Ind. Crops Prod.* 151: 112453.
- Szczeponek, A., Mazur, S. and Nawrocki, J. (2006). The usage of chitosan in protection of some peppermint and lemon balm pathogens. *Pol. Chitin Soc. Monogr.* 11: 193-200.
- Tambascio, C., Covacevich, F., Lobato, M. C., de Lasa, C., Caldiz, D., Dosio, G. and Andreu, A. (2014). The application of K phosphites to seed tubers enhanced emergence, early growth and mycorrhizal colonization in potato (*Solanum tuberosum*). *Am. J. Plant Sci.* 5: 132-137.
- Tanwar, A., Aggarwal, A. and Prakash, V. (2014). Effect of bioinoculants and superphosphate fertilizer on the growth and yield of broccoli (*Brassica oleracea* L. var. *italica* Plenck). *Nea Zea J. Crop Hort. Sci.* 42(4): 288–302.
- Trejo-Téllez, L. I. and Gómez-Merino, F. C. (2018). Phosphite as an inductor of adaptive responses to stress and stimulator of better plant performance. In: Vats, S. (Ed) *Biotic and Abiotic Stress Tolerance in Plants*: 203-238, Springer, Singapore.
- Uppal, A. K., El Hadrami, A., Adam, L. R., Tenuta, M. and Daayf, F. (2008). Biological control of potato *Verticillium* wilt under controlled and field conditions using selected bacterial antagonists and plant extracts. *Biological Control* 44(1): 90-100.
- Van Hemelrijck, W., Hauke, K., Creemers, P., Mery, A. and Joubert, J.-M. (2013). Efficacy of a new oligosaccharide active against scab on apple. *Acta Hortic*. 1009: 45-52.
- Vera, J., Castro, J., Gonzalez, A. and Moenne, A. (2011). Seaweed polysaccharides and derived oligosaccharides stimulate defense responses and protection against pathogens in plants. *Mar. Drugs* 9(12): 2514–2525.
- Visconi, F., de Paz, J. M., Bonet, L., Jordà, M., Quiñones, A. and Intrigliolo, D. S. (2015). Effects of a commercial calcium protein hydrolysate on the salt tolerance of *Diospyros kaki* L. cv. "Rojo Brillante" grafted on *Diospyros lotus* L. Sci. Hort. 185: 129-138.
- Wang, C., Li, X., Zhou, J., Wang, G. and Dong, Y. (2008). Effects of arbuscular mycorrhizal fungi on growth and yield of cucumber plants. *Commun. Soil Sci. Plant Anal.* 39(3-4): 499-509.

- Wei, Z., Yang, X., Yin, S., Shen, Q., Ran, W. and Xu, Y. (2011). Efficacy of *Bacillus*-fortified organic fertilizer in controlling bacterial wilt of tomato in the field. *Appl. Soil Ecol.* 48(2): 152-159.
- Wu, Q. S. and Xia, R. X. (2006). Arbuscular mycorrhizal fungi influence growth, osmotic adjustment and photosynthesis of citrus under well-watered and water stress conditions. J. Plant Physiol. 163(4): 417-425.
- Wu, Q.-S., Zou, Y.-N. and He, X.-H. (2010). Contributions of arbuscular mycorrhizal fungi to growth, photosynthesis, root morphology and ionic balance of citrus seedlings under salt stress. *Acta Physiol. Plant.* 32(2): 297-304.
- Xi, Y., Han, X., Zhang, Z., Joshi, J., Borza, T., Aqa, M. M., Zhang, B., Yuan, H. and Wang-Pruski, G. (2020). Exogenous phosphite application alleviates the adverse effects of heat stress and improves thermotolerance of potato (*Solanum tuberosum* L.) seedlings. *Ecotoxicol. Environ. Saf.* 190: 110048.
- Xiong, J.-Q., Kurade, M. B. and Jeon, H.-H. (2017). Can microalgae remove pharmaceutical contaminants from water? *Tren Biotech*. 36: 30-44.
- Yoshioka, Y., Ichikawa, H., Naznin, H. A., Kogure, A. and Hyakumachi, M. (2012). Systemic resistance induced in *Arabidopsis thaliana* by *Trichoderma asperellum* SKT-1, a microbial pesticide of seedborne diseases of rice. *Pest Manag. Sci.* 68(1): 60-66.
- Zebelo, S., Song, Y., Kloepper, J. W. and Fadamiro, H. (2016). Rhizobacteria activates (+)-δ-cadinene synthase genes andinduces systemic resistance in cotton against beet armyworm (*Spodoptera exigua*). *Plant Cell Environ*. 39(4): 935-943.
- Zehnder, G., Kloepper, J., Tuzun, S., Yao, C., Wei, G., Chambliss, O. and Shelby, R. (1997). Insect feeding on cucumber mediated by rhizobacteria-induced plant resistance. *Entomol. Exp. Appl.* 83(1): 81-85.
- Zehnder, G. W., Murphy, J. F., Sikora, E. J. and Kloepper, J. W. (2001). Application of rhizobacteria for induced resistance. *Eur. J. Plant Pathol.* 107(1): 39-50.
- Zhao, H., Sun, R., Albrecht, U., Padmanabhan, C., Wang, A., Coffey, M. D., Girke, T., Wang,
   Z., Close, T. J., Roose, M., Yokomi, R. K., Folimonova, S., Vidalakis, G., Rouse, R.,
   Bowman, K. D. and Jin, H. (2013). Small RNA profiling reveals phosphorus deficiency
   as a contributing factor in symptom expression for citrus huanglongbing disease.
   *Mol. Plant* 6(2): 301-310.
- Zhang, M., Liang, Y. and Chu, G. (2017). Applying silicate fertilizer increases both yield and quality of table grape (*Vitis vinifera* L.) grown on calcareous grey desert soil. *Sci. Hort*. 225: 757–763.
- Zhang, M., Tan, T., Yuan, H. and Rui, C. (2003). Insecticidal and fungicidal activites of chitosan and oligo-chitosan. *J. Bioact. Compat. Poly* 18(5): 391-400.
- Zhou, J., Feng, Z., Liu, S., Wei, F., Shi, Y., Zhao, L., Huang, W., Zhou, Y., Feng, H. and Zhu, H. (2021). CGTase, a novel antimicrobial protein from *Bacillus cereus* YUPP-10, suppresses *Verticillium dahliae* and mediates plant defence responses. *Mol. Plant Pathol.* 22(1): 130-144.