

# THE EFFECTS OF ANTAGONISTIC BACTERIA AGAINST WHITE MOLD DISEASE AGENT [*SCLEROTINIA SCLEROTIORUM* (LIB.) DE BARY] IN CUCUMBER

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**Abstract.** This study was carried out to determine the effects of antagonistic bacteria, which were obtained from the soil rhizosphere and have the strong abilities to dissolve phosphorus and reduce nitrogen, on the white mold disease agent *Sclerotinia sclerotiorum* (Lib.) de Bary. In the study, ten antagonistic bacteria of the genus *Pseudomonas* were used. In the *in vitro* study, the tested antagonistic bacteria suppressed the mycelial growth of *S. sclerotiorum* by 65.49-88.25%. Isolates with a high inhibition rate on mycelial growth were also effective on the viability of the sclerotia of the pathogenic fungus. Five bacterial isolates caused complete sclerotia viability loss for *S. sclerotiorum*. According to the result of the *in vitro* studies, antagonistic bacterial isolates with strong effects on the pathogenic fungus were selected and assayed in pot studies under greenhouse conditions. In pot studies, single treatments with *Pseudomonas chlororaphis* (R-7) and the dual treatments with *Pseudomonas brassicacearum* + *Pseudomonas chlororaphis* (R-3 + R-7) were the most effective against white mold disease. The results of this study show that these antagonistic bacteria can be used in the biological control of *S. sclerotiorum*. In addition to the *in vitro* and *in vivo* studies, it is necessary to determine the performances of antagonistic bacterial isolates, which had the highest effect on white mold disease, in greenhouses and field conditions.

**Keywords:** *biological control, Sclerotinia sclerotiorum, Pseudomonas brassicacearum, Pseudomonas chlororaphis*

## Introduction

Species in *Sclerotinia* genus are soil-borne pathogens that cause significant losses to host plants. *Sclerotinia sclerotiorum* (Lib.) de Bary, causes white mold disease in field crops and vegetables all over the world especially in temperate regions (Ferreira and Boley, 2002). *S. sclerotiorum* is a reported host range of over 500 plant species from 75 families (Saharan and Mehta, 2008). In Turkey, white mold diseases has been reported in various proportions on lettuce (*Lactuca sativa* L.) in Izmir, Manisa and Aydın (Yıldız, 1970), on sunflower (*Helianthus annuus* L.) in Thrace Region and Cukurova Region (Yucer, 1980; Cinar and Bicici, 1982), on cucumber (*Cucumis sativus* L.), tomato (*Solanum lycopersicum* L.) and eggplants (*Solanum melongena* L.) in the Eastern Mediterranean Region (Aksay et al., 1991), on cucumber in Tokat and Amasya (Onaran and Yanar, 2004), on lettuce in Canakkale (Turk and Dogu, 2004), on greenhouse tomatoes in the provinces of Hatay, Adana, Mersin and Antalya (Tok and Kurt, 2007), in the sunflower fields of the Pasinler Plain of Erzurum (Tozlu and Demirci, 2008), on cucumbers in the greenhouses of Antalya (Onaran, 2009) and on potato in Hatay province (Kurt et al., 2017).

*Sclerotinia sclerotiorum* produces lots of mycelia and these mycelia aggregate to form a hard, dark brown or black sclerotia, which are of different sizes and irregular

shapes. This agent is easily recognized from this sclerotium, which is resistant to unfavorable environmental conditions. *S. sclerotiorum* usually attacks the roots or stem of the host that are in close proximity to the soil. Lesions develop on the stem and gradually surround the roots, and the host plant withers and dies. This fungus can cause complete rotting of seedlings, especially in humid environments (Anonymous, 2008).

This pathogen overwinters as sclerotium in soil, contaminated plant debris or as mycelia. In the spring, sclerotia germinate and forms apothecia in which asci and ascospores are produced. Ascospores are released into the air from the apothecia and infection begins when they land and germinate on host plant (Agrios, 1997). Studies have stated that the sclerotia of this fungus can survive in the soil for more than 5 years (Adams and Ayers, 1979) and that environmental conditions are important in the development of the disease.

Cultural measures, physical control, fumigation of soil and green parts are used in the management of white mold disease (Anonymous, 2008), however, difficulties are experienced in managing the disease as some varieties are susceptible to the pathogen, sclerotia of this fungal pathogen can remain viable for many years and not all of them germinate at the same time and more importantly the pathogen is becoming more resistant to currently available fungicides. Accordingly, different alternative methods to chemical control such as the use of antagonistic bacteria and plant growth promoting rhizobacteria (PGPR) are in demand. These bacteria are densely located around the soil rhizosphere of plant roots and they carry out physiochemical activities in the soil. They are in close relationship with plant roots and directly or indirectly have positive effect on the development of the plants. These bacteria can be found in several genera but those in the genus *Pseudomonas* are highly efficacious and can multiply rapidly in the soil rhizosphere and spread to the root. In addition, by protecting the plant from stress, they promote plant root and shoot development, improve yield and protect plants against diseases (Imriz et al., 2014).

Several studies have been conducted by various researchers on the biological control of *Sclerotinia sclerotiorum* (Zizzerini and Tosi, 1985; Bogdanova et al., 1986; Turhan and Grossmann, 1986; Inbar et al., 1996; Aksay et al., 1991; Tuncer and Damdere, 1997; Thaning et al., 2001; Al-Masri et al., 2002; Kamensky et al., 2003; Mansour et al., 2008; El-Kafrawy, 2008; Onaran and Yanar, 2011; Tozlu and Demirci, 2011; Rostami et al., 2013; Tozlu et al., 2016; Abdeljalil et al., 2016; Helmy, 2016; Mun et al., 2019). In the search for alternatives, a number of mycoparasitic fungi grow on sclerotia (Adams, 1990) and *Coniothyrium minitans* was the most investigated fungi on *S. sclerotiorum* (Whipps and Gerlagh, 1992; Budge et al., 1995; McLaren et al., 1996; Whipps et al., 2008) and has become available as a commercial product for sclerotia (Gerlagh et al., 1999). Also, *Gliocladium virens* and *Trichoderma harzianum* have been reported to reduce growth of mycelium and apothecial production of *S. sclerotiorum* (Phillips, 1986; Whipps and Budge, 1990; Srinivasan et al., 2001; Mehta et al., 2012). Among the antagonistic bacteria species tested, *Bacillus subtilis* and *Bacillus cereus* have been reported to reduce germination of sclerotia and cause hyphal death (Zizzerini et al., 1987; Kamal et al., 2015; Sun et al., 2017). Similarly, it has been determined that *Pseudomonas putida* and *P. fluorescens* species reduce the damage caused by *S. sclerotiorum* (Expert and Digat, 1995), and that *Bacillus* spp. and *Pseudomonas* spp. prevent disease development by 75.3% (Soylu et al., 2005). Also, *Serratia plymuthia* was highly effective in inducing complete suppression of apothecial formation and

strongly inhibited the germination of ascospores (Thaning et al., 2001). Under in vivo conditions, *Burkholderia cepacia*, and *Serratia plymuthica* species (Onaran and Yanar, 2011), *B. subtilis*, *Serratia* sp., and *P. fluorescens* (Helmy, 2016; El-Kafrawy, 2008), *P. chlororaphis* PA-23 (Savchuck, 2002) species have described to significantly reduce disease development. Studies conducted at the same time have also determined that *B. subtilis*, *B. thuringiensis*, *B. amyloliquefaciens*, *Enterobacter cloacae*, and *P. fluorescens* isolates positively affect plant growth (El-Kafrawy, 2008; Abdeljalil et al., 2016).

The aim of this study was to determine the effects of antagonistic bacteria isolates obtained from tomato production areas in Tokat province on *Sclerotinia sclerotiorum*, which causes white mold disease in cucumbers.

## Materials and methods

In the study, 10 antagonistic bacterial isolates with high phosphate-reducing and nitrogen-binding properties, previously isolated from tomato production areas in Tokat province in Turkey and described with biochemical tests and MALDI-TOF technique, were used (Table 1). *Sclerotinia sclerotiorum* isolate used in the study was isolated from infected tissues of cucumber plant and identified using morphologically and molecular analysis (PCR).

Tests were carried out in Tokat province of Turkey in 2019. In vitro and in vivo tests were carried out as laboratory test and pot studies, respectively.

**Table 1.** Scientific names and codes of antagonistic bacteria used in this study

Isolate code	Scientific name of the isolate
R-1	<i>Pseudomonas kilonensis</i>
R-2	<i>Pseudomonas monteilii</i>
R-3	<i>Pseudomonas brassicacearum</i>
R-4	<i>Pseudomonas thivervalensis</i>
R-5	<i>Pseudomonas koreensis</i>
R-6	<i>Pseudomonas thivervalensis</i>
R-7	<i>Pseudomonas chlororaphis</i>
R-8	<i>Pseudomonas</i> sp.
R-9	<i>Pseudomonas</i> sp.
R-10	<i>Pseudomonas thivervalensis</i>

## In vitro tests

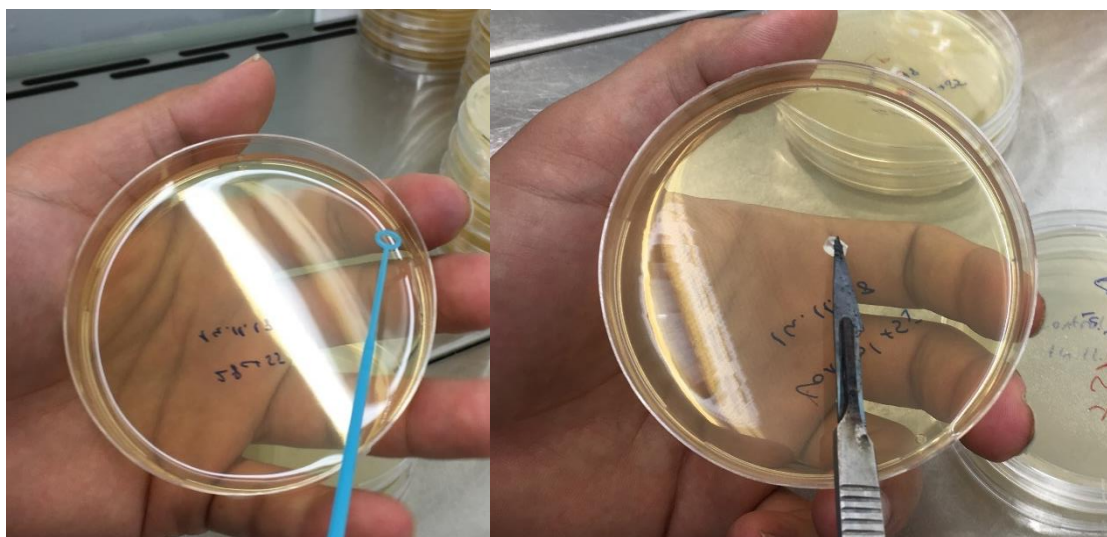
### Determination of the effects of antagonistic bacteria on the mycelial development of *Sclerotinia sclerotiorum*

Tryptic soy agar (TSA) was used throughout this study. The antagonistic bacterial isolates grown TSA for 24 h were inoculated in a ring form on the edges of the medium in 90 mm diameter petri dishes. Then, a 5 mm diameter mycelium disc was taken from *S. sclerotiorum* culture and placed in the center of the medium (Fig. 1). In the control group, only the pathogen fungus was plated onto the medium without bacteria. Petri dishes were wrapped with Parafilm and left to incubation at 25 °C. The

experiments were ended when the fungus in the control group completely covered the petri dish and the radial growth of fungi in the control group and petri dishes treated with antagonist bacteria were measured (Tozlu et al., 2016; Kotan, 2017). Inhibition rates (%) were calculated by comparing the measured mycelium diameters. The experiment was repeated twice with five replicates per treatment (Aeron et al., 2011).

$$I (\%) = [(C - T) / (C - C_0)] \times 100$$

I: inhibition rate (%); C: mycelial diameter of the pathogen in control; T: mycelial diameter of the pathogen in bacterial treatment; C<sub>0</sub>: the diameter of the test fungus agar discs (5 mm).



**Figure 1.** Inoculation of antagonistic bacteria in a ring form on the edges of the medium-position of mycelium disc from *S. sclerotiorum* culture in the center of the medium

#### *Determination of the effects of antagonistic bacteria on germination of Sclerotinia sclerotiorum sclerotia*

First the sclerotia of *S. sclerotiorum*, which were developed on Potato Dextrose Agar (PDA) medium for 10 days, were subjected to surface sterilization by immersing in 2% sodium hypochlorite and passing through sterile distilled water three times. The surface-sterilized sclerotia were placed in bacterial suspensions with 10<sup>8</sup> cells/ml density prepared with 10 ml of Luria Bertani Broth (LB) medium. Five sclerotia were used for each antagonistic bacteria isolate, and the treated erlenmeyers were incubated in a rotary incubator at 175 rpm for 24 h. The sclerotia treated with antagonistic bacteria were dissected into two with a scalpel and transferred to the PDA medium and left to incubate at 25 °C for five days. The sclerotia that germinated at the end of the incubation period were considered to be alive and the diameter of mycelium were measured and recorded. For the control group, five sclerotia were placed in LB medium without bacterial isolates. Each treatment in the experiment had three replicates and the study was repeated twice. The effects of the antagonistic bacterial isolates on the survival rates and mycelium growth of sclerotia was determined and compared with the control group (Abdeljalil et al., 2016).

### ***In vivo test***

According to the results obtain in the in vitro test, pot experiments were conducted using single, double and triple combinations of the three antagonistic bacteria isolates which had the highest effect on the pathogen. Dörtel F1 cucumber variety, which is sensitive to pathogen fungus, was used in the study which was conducted under controlled greenhouse conditions. Bacterial isolates that were streaked on King B media and incubated at 28 °C for 48 h. Grown bacterial isolates were suspended in saline (0.85% NaCl), and the density was adjusted to 10<sup>8</sup> cells/ml at 600 nm using spectrophotometer (PG Instrument T60U UV/VIS). Cucumber seedlings were dipped into the prepared suspension and left for 1 h for bacteria colonization and then planted in pots with sterile soil, peat and perlite (1:1:0.5). Seven days after the seedlings were transferred to the pots, a wound with a diameter of 5 mm was opened on the plant stem 4 cm above the soil, then 0.5 ml (10<sup>8</sup> cells/ml) bacterial suspension pipetted was on the wounds. Immediately afterwards, a 5-mm mycelium disc of *S. sclerotiorum* was placed on the opened wound. The inoculated spot was covered with moist cotton and wrapped with plastic film. Positive control group had plants inoculated only with *S. sclerotiorum* (Tozlu et al., 2016). In the negative control group, only water was applied to the plants, and for the chemical control group, Metalaxyl-M + Fludioxonil was applied to the plants at the dose recommended by the producer. Trial coincidence plots were set up two times with 10 replicates according to the trial pattern. Seven days after the application, dead and living plants in the control group and the treatments were counted.

### ***Statistical analyses***

The data obtained in the study were analyzed using variance analysis in SPSS statistics 25 package program and the differences between the means were determined with Duncan multiple comparison test at the  $p \leq 0.05$  significance level.

## **Results**

### ***In vitro tests***

#### ***Effects of antagonistic bacteria on the mycelial development of Sclerotinia sclerotiorum***

In in vitro studies, ten of the antagonistic bacteria isolates tested showed varying effects ranging between 65.49 and 88.25% on the pathogenic fungus, *S. sclerotiorum*. As can be seen in *Table 2*, it was determined that the effect of all bacterial isolates on the fungus were statistically significant compared to the control group. R-6 coded *Pseudomonas thivervalensis* isolate had the highest (88.25%) effect on fungal pathogen (*Fig. 2*). All of the bacterial isolates tested had an inhibitory effect on the mycelium growth of the fungal pathogen greater than 60% (*Table 2*).

#### ***The effect of antagonistic bacteria on the germination of Sclerotinia sclerotiorum sclerotia***

In this part of the study, the effect of the bacterial isolates on sclerotia viability was examined and according to the results, sclerotia treated with five isolates (R-1, R-2, R-4, R-5, R8) remained viable while the sclerotia treated with other five isolates (R-3, R-6, R-7, R-9, R-10) were dead. Isolates which were highly efficacious on mycelium

development were also effective on sclerotia. In addition, R2, R4 and R5 coded antagonists significantly suppressed mycelial growth of sclerotia that did not lose their viability. This situation reveals that these antagonists have fungistatic effects (Table 3).

**Table 2.** The effects of antagonistic bacteria on the mycelial development of *Sclerotinia sclerotiorum*

Codes	Treatments	Diameter of mycelium (mm)	Average effect rate (%)
K	Control	90 a*	0 f
R-1	<i>Pseudomonas kilonensis</i>	23.81 cd	73.54 cd
R-2	<i>Pseudomonas monteilii</i>	13.46 ef	85.04 ab
R-3	<i>Pseudomonas brassicacearum</i>	10.85 f	87.93 a
R-4	<i>Pseudomonas thivervalensis</i>	12.40 ef	86.31 ab
R-5	<i>Pseudomonas koreensis</i>	15.27 e	83.03 b
R-6	<i>Pseudomonas thivervalensis</i>	10.56 f	88.25 a
R-7	<i>Pseudomonas chlororaphis</i>	14.71 ef	83.64 ab
R-8	<i>Pseudomonas</i> sp.	31.05 b	65.49 e
R-9	<i>Pseudomonas</i> sp.	25.86 c	71.26 d
R-10	<i>Pseudomonas thivervalensis</i>	20.03 d	77.73 c

\*The same letters next to averages in the same column indicate that difference between applications are not significant. (Duncan multiple comparison test,  $p \leq 0.05$ )

**Table 3.** The effect of antagonistic bacteria on *Sclerotinia sclerotiorum* sclerotia viability

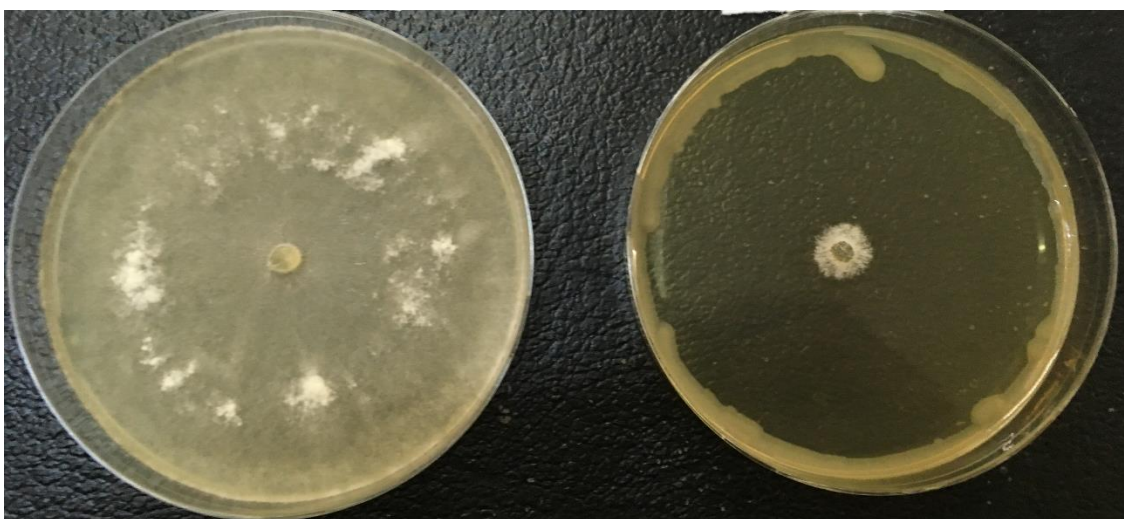
Codes	Treatments	Sclerotia viability	Diameter of mycelium (mm)
K	Control	Viable	60 a*
R-1	<i>Pseudomonas kilonensis</i>	Viable	53.09 b
R-2	<i>Pseudomonas monteilii</i>	Viable	21.25 d
R-3	<i>Pseudomonas brassicacearum</i>	Dead	0 f
R-4	<i>Pseudomonas thivervalensis</i>	Viable	14.27 e
R-5	<i>Pseudomonas koreensis</i>	Viable	26.38 cd
R-6	<i>Pseudomonas thivervalensis</i>	Dead	0 f
R-7	<i>Pseudomonas chlororaphis</i>	Dead	0 f
R-8	<i>Pseudomonas</i> sp.	Viable	32.39 c
R-9	<i>Pseudomonas</i> sp.	Dead	0 f
R-10	<i>Pseudomonas thivervalensis</i>	Dead	0 f

\*The same letters next to averages in the same column indicate that difference between applications are not significant. (Duncan multiple comparison test,  $p \leq 0.05$ )

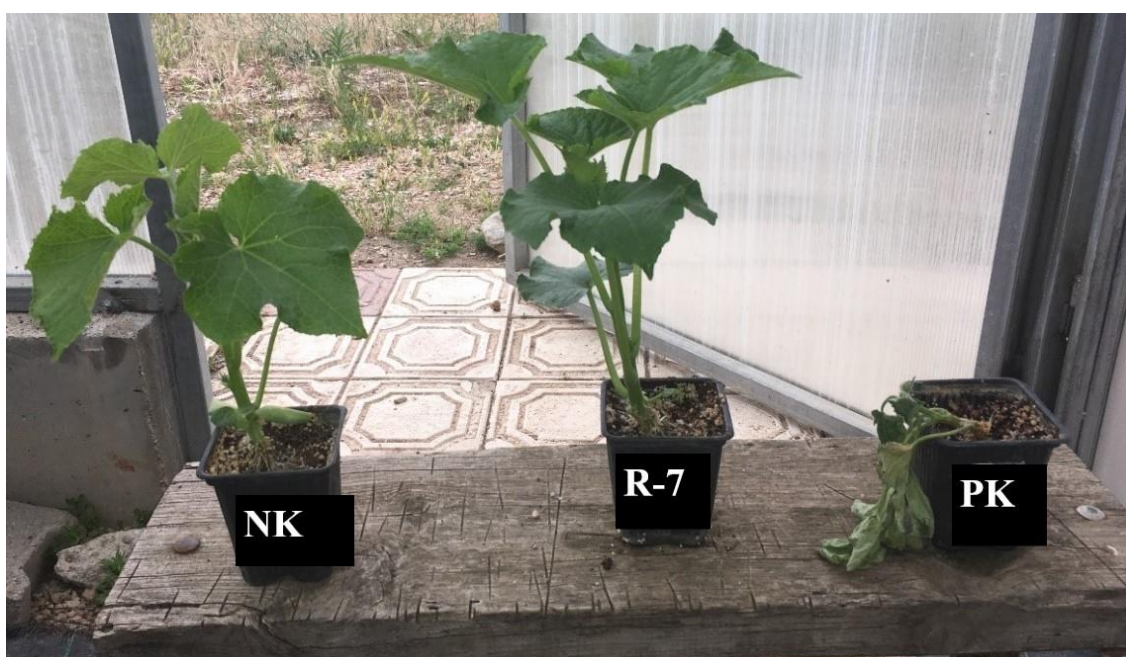
### ***In vivo* test**

*In vivo* tests were conducted with three isolates (R-3, R-6 and R-7) which had the highest effects on the mycelial growth and sclerotia viability of *S. sclerotiorum* in the *in vitro* studies. Single, double and triple combinations of these three isolates were used. In the evaluation of the applications in the pot study conducted under greenhouse conditions,

the dead/alive plants were counted. According to the results of the study, 100% death occurred in positive control where only the fungal pathogen was applied, whereas no plant death occurred in negative control and pesticide control applications. In R-3 single application and R-3 + R-6 + R-7 triple combination application, death (50%) occurred in 5 out of 10 plants. In R-6 single application, and R-3 + R-6 and R-6 + R-7 double combination applications, death occurred in 6 out of 10 plants (60%). Among the applications, the highest effect was seen in R-7 single application (Fig. 3) and R-3 + R-7 double application. In both applications, while death occurred in 2 out of 10 plants, it was observed that 8 of them were alive and there were no signs of disease or death. It was found that the disease was suppressed by 80% in both applications (Table 4).



**Figure 2.** Effect of R-6 coded *Pseudomonas thivervalensis* isolate on the mycelial development of *S. sclerotiorum*



**Figure 3.** Differences in plant growth in NK, R-7 and PK treatments

**Table 4.** The effect of antagonistic bacteria on white rot disease in cucumber plant

Codes	Treatments	Number of dead plants	Number of live plants	Inhibition rate (%)
PK	<i>Sclerotinia sclerotiorum</i> treatment	10	0	0
NK	Treatment with pure water	0	10	100
IK	Metalaxyl-M + Fludioxonil	0	10	100
R-3	<i>Pseudomonas brassicacearum</i>	5	5	50
R-6	<i>Pseudomonas thivervalensis</i>	6	4	40
R-7	<i>Pseudomonas chlororaphis</i>	2	8	80
R-3 + R-6	<i>Pseudomonas brassicacearum</i> + <i>Pseudomonas thivervalensis</i>	6	4	40
R-3 + R-7	<i>Pseudomonas brassicacearum</i> + <i>Pseudomonas chlororaphis</i>	2	8	80
R-6 + R-7	<i>Pseudomonas thivervalensis</i> + <i>Pseudomonas chlororaphis</i>	6	4	40
R-3 + R-6 + R-7	<i>Pseudomonas brassicacearum</i> + <i>Pseudomonas thivervalensis</i> + <i>Pseudomonas chlororaphis</i>	5	5	50

## Discussion

Today, in order to increase the quality and yield in plant production and reduce the use of pesticides, the use of antagonistic bacteria against plant diseases and/or plant growth regulating rhizobacteria within the scope of biological control has been an important research topic. Studies with these bacteria show that they are promising in terms of suppressing plant diseases and ensuring healthy plant growth. Although cultural measures, physical and chemical control are used in the management of white mold disease, there are difficulties in managing the disease due to fungicide resistance problem and the survival of sclerotia for many years.

In this study, the effects of antagonistic bacteria which were obtained from the soil rhizosphere and have the high ability to dissolve phosphorus and reduce nitrogen, was determined on the white mold disease agent *Sclerotinia sclerotiorum* (Lib.) de Bary in vitro and in vivo conditions. Our in vitro studies, antagonistic bacteria isolates tested showed varying effects ranging between 65.49-88.25% on *S. sclerotiorum*. And, five bacterial isolates caused complete sclerotia viability loss for *S. sclerotiorum*. Numerous studies similar to our in vitro study have been carried out by various researchers. El-Kafrawy (2008) reported that *Pseudomonas fluorescens* had an antagonistic effect of 69.26% on the radial growth of *S. sclerotiorum* and showed a significant suppressive effect on formation of sclerotia. A study by Helmy (2016) established that *Streptomyces* sp., *Pseudomonas fluorescens* and *Bacillus subtilis* (Bs1) reduced mycelium growth of *S. sclerotiorum* by 72.2%, 68.0, 62.22 respectively. Soylu et al. (2005) examined the antagonistic potential of 113 bacterial isolates against *S. sclerotiorum* in dual petri dishes and reported that *Pseudomonas* spp. AFP104 isolate was 83.3% effective on fungal pathogen. Similarly, study conducted by Onaran and Yanar (2011) showed that 12 of the 23 tested isolates especially *P. putida* and *P. fluorescens* from the genus *Pseudomonas*, *Paenibacillus macerans*, and *Bacillus pumilis* were highly effective and caused sclerotia death.

In our pot studies, *Pseudomonas chlororaphis* (R-7) and *Pseudomonas brassicacearum* + *Pseudomonas chlororaphis* (R-3 + R-7) were the most effective against white mold disease. It was found that the disease was suppressed by 80% in both applications. Previous studies conducted on the biological control of white mold disease



have determined that several antagonistic bacteria partially or completely inhibits this fungal agent. Similar to our study, Onaran and Yanar's (2011) study under greenhouse conditions showed that the bacterial isolates tested were effective at 43.20-89.22% on white mold disease with *Burkholderia cepacia* being the most effective. Likewise, Helmy (2016) reported that dipping seedlings into bacterial suspensions of *Bacillus subtilis*, *B. thuringiensis* and *Streptomyces* sp. prevented the disease severity caused by fungal pathogen by 100% after one week. A greenhouse study determined that *Bacillus subtilis* and *Pseudomonas fluorescens* species prevented white mold disease by 95-100% and 85-90%, respectively when cucumber seedlings were planted in soils treated with these antagonistic bacteria (El-Kafrawy, 2008).

Besides cucumber plants, a study conducted under greenhouse and field conditions for the biological control of *S. sclerotiorum* in canola plant, showed that *P. chlororaphis* (PA-23), *Pseudomonas* sp. (DF-41) significantly suppressed the disease compared to *B. amyloliquifaciens* (BS6) fungicide application. While the rate of stem mold in the canola plants varied between 20-75%, this rate was 5.0-29.6% in *P. chlororaphis* application (Fernando et al., 2007). In addition, Savchuck and Fernando (2004) stressed that the application of *P. chlororaphis* and *P. brassicacearum* at a density of  $10^4$ - $10^8$  cfu/ml was highly effective on ascospore germination.

Antagonistic bacteria against *S. sclerotiorum* affect the systemic resistance of the host plant. *Pseudomonas* spp., produces numerous antimicrobial compounds such as pyoluteorin, pyrrolnitrin, phenazines, siderophores, cyanide, 2,4-diacetylphloroglucinol (Compant et al., 2005) and enzymes such as cellulose, chitinase, proteases, beta-glucanase (Hernandez-Leon et al., 2015). Similarly, bacterial isolates of *Pseudomonas* spp. DF200 and DF209 which inhibit *S. sclerotiorum* under in vitro conditions produce benzothiazole, cyclohexanol, n-decanal, dimethyl trisulfide, 2-ethyl 1-hexanol, nonanal antifungal organic volatile compounds. In the field, bacteria prevent carpogenic germination of sclerotia and emergence of ascospores (Fernando et al., 2004). It has been reported that *Pseudomonas brassicacearum*, used in the present study, uses hydrogen cyanide (HCN), protease, alginate, sclerocin and lipopeptide molecules, 2,4-diacetylphloroglucinol and antifungal compounds such as cyanide in suppressing *S. sclerotiorum* (Berry et al., 2010, 2014; Ortet et al., 2011; Loewen et al., 2014).

*Pseudomonas chlororaphis*, also used in this study, is an important biocontrol agent that suppresses *S. sclerotiorum* by producing antibiotics such as phenazine-1-carboxylic acid, 2-hydroxyphenazine and pyrrolnitrin (PRN) which causes hyphal lysis, vacuolation and protoplast leakage, hereby preventing sclerotia and spore germination (Savchuck, 2002; Fernando et al., 2007; Zhang et al., 2004a, b; Selin et al., 2010).

## Conclusion

In recent years, negatives effects of pesticides used extensively in agricultural areas on both the environment and human health have been reported in many studies. Antagonistic bacteria or plant growth promoting rhizobacteria have the potential to protect against soil-borne pathogens. Biopreparat or biofertilizers made from such microorganisms can be widely used in areas where sustainable organic agriculture. Our study revealed that antagonistic bacterial isolates have the potential to be used as a biological control agent against white rot disease. In this context, the efficacy of the R-7 coded isolate and the R-3 + R-7 coded double combination, which were effective on the agent in both the in vitro and in vivo studies, should also be evaluated under greenhouse and field conditions.

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