
Induction of Tomato Plant Resistance to the Whitefly *Bemisia tabaci* Using the Entomopathogenic Fungus *Beauveria bassiana*

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Abstract

Whitefly (*Bemisia tabaci*) is a major pest on tomato plants. Whiteflies not only attack plants directly but also serve as vectors for viruses that spread diseases such as geminiviruses, causing plants to turn yellow. One alternative for controlling this pest is to use entomopathogenic fungi, such as *Beauveria bassiana*. This study aimed to isolate a *B. bassiana* strain that enhances tomato plant resistance to *B. tabaci*. The experiment was arranged in a Completely Randomized Design (CRD) with six treatments and five replications. The treatments included five isolates of *B. bassiana*, namely BbWs, TD312, PA221, PD114, and PB21 plus a control. The *B. bassiana* concentration used was 10^8 conidia/mL. The fungus was applied by soaking tomato seed lots. Results showed that seed soaking with *B. bassiana* significantly reduced egg, nymph, and adult populations across all isolates compared to the control. Notably, PA221 and TD312 isolates showed significant nymph suppression. Five weeks after planting, the TD312 isolate resulted in the lowest populations: 20.40 eggs per plant, 7.25 nymphs per plant, and 2.30 adults per plant. Additionally, *B. bassiana* application affected plant morphology by increasing trichome density; plants treated with TD312 had a trichome density of 616.73 trichomes/cm², higher than the control (295.73 trichomes/cm²). Although statistically comparable to several other isolates on some parameters, TD312 consistently exhibited the lowest pest counts and the highest trichome density, indicating its potential to enhance tomato resistance to *B. tabaci* by suppressing pest populations and reinforcing trichome density.

Keywords: Egg, endophytes, fungi, Gemini virus, nymph.

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INTRODUCTION

The whitefly (*Bemisia tabaci* Gennadius, 1889) is a significant pest of tomato plants. *B. tabaci* damages plants by piercing tissue and sucking phloem fluid from tomato leaves, causing abnormal plant growth, wilting, and decreased productivity (Agastya et al., 2020). Whiteflies not only attack plants directly but also act as vectors for viruses that transmit diseases such as Geminivirus, which causes yellowing (Soumia et al., 2021). Their role as pests and disease vectors in plants can cause significant yield losses, reaching 90-100% (Hidayat et al., 2020).

Standard methods of controlling whitefly pests used by farmers include adjusting planting times, planting resistant varieties, using barrier plants, crop rotation, and applying synthetic insecticides (Inayati and Marwoto, 2015). The most common method of controlling whiteflies is using insecticides. However, these insecticides are considered ineffective because the insect's body is coated with wax, making it difficult for the active ingredient to penetrate. Whiteflies typically live under the surface of leaves, and their adults fly, making it difficult for insecticide sprays to reach their target. Furthermore, the pest is suspected to belong to a lineage resistant to many insecticides (Sugiyama, 2005; Mota-Sanchez and

Wise, 2019; Horowitz et al., 2020). Besides that, the intensive use of synthetic insecticides can lead to resistance, pest resurgence, the elimination of natural enemies, the accumulation of harmful residues on agricultural products, environmental pollution, and a risk to human health. One of the more environmentally friendly pest control efforts that also suppresses the development of pest resistance or resurgence is the use of biological agents such as the fungus *Beauveria bassiana* (Erdiansyah et al., 2023; Hendra et al., 2026).

B. bassiana is an entomopathogenic fungus widely used as an effective biological control agent against various insect pest species, including the cabbage pest *Crociodolomia pavonana* (Trizelia and Nurdin, 2010), *Spodoptera exigua* (Razak et al., 2016), *S. litura* (Trizelia et al., 2016), *Nezara viridula* (Siahaan et al., 2021), *Bemisia tabaci* (Flawerina, 2021), *Eurydema pulchrum* (Trizelia et al., 2019), *Nilaparvata lugens* (Hendra et al., 2022), and Thrips sp. on chili plants (Sofwah and Prastowo, 2023)

As knowledge advances, *B. bassiana* is not only an entomopathogen capable of directly infecting and killing insects. Still, it is also reported to live as an endophyte in various plant species and to colonize plant tissues. The colonization of endophytic fungi in plant hosts can negatively affect insects, especially those that feed on the host and become pests, thereby enhancing plant resistance to pest attacks. The plant's ability to inhibit pest development is a complex process regulated by various compounds, including salicylic acid, jasmonic acid, and ethylene (Kessler & Baldwin, 2002). The presence of *B. bassiana* in plant tissues can occur naturally (as a natural endophyte) or be artificially introduced through seed inoculation, foliar application, seedling and vegetative propagation material soaking, soil drenching, and injection (Vega, 2018; Bamisile et al., 2018; Saragih, 2019).

The ability of *B. bassiana* to live endophytically in plants and enhance plant resistance to pest attacks has been reported by several researchers. Gautam et al. (2016) found that *B. bassiana* can live endophytically in cauliflower plants, reducing the number of *Plutella xylostella* eggs and causing up to 100% larval mortality. This mortality results from *B. bassiana*'s ability to reside in plant tissues and produce secondary metabolites toxic to larvae. Batool et al. (2020) reported that *B. bassiana* can live endophytically in corn plants. Its presence reduces *Ostrinia furnacalis* (Guenee) larvae by up to 85%, decreasing the number and length of larval tunnels. Larval mortality is attributed to secondary metabolites such as proline and polyphenol oxidase found in corn plants, which are toxic to larvae. Trizelia et al. (2020) reported that chili plants inoculated with *B. bassiana* can suppress the development of *Myzus persicae* populations. Hendra et al. (2023) found that applying *B. bassiana* to rice plants through seed soaking affects the oviposition preferences of adult brown planthoppers (*Nilaparvata lugens*). The number of eggs laid by adult planthoppers was lower in *B. bassiana*-treated rice plants. The presence of *B. bassiana* in rice tissues reduces the percentage of planthopper eggs that hatch by 62.8% and increases nymph mortality by 46%. It also affects adult planthopper mortality. The presence of *B. bassiana* in rice stems can increase secondary metabolite levels, such as salicylic acid and oxalic acid, and decrease primary metabolite levels, such as sucrose (Hendra, 2022).

The ability of *B. bassiana* to colonize plant tissues is affected by application methods, plant age (Posada and Vega, 2005), and the specific fungal strain used (Zhang, 2014). The presence of endophytic fungi within plant tissues can promote the production of various toxic or antifeedant compounds (Gao et al., 2010; McCormick et al., 2016). The relationship between applying *B. bassiana* to tomato plants and the development of aphid populations warrants further study. This research aims to isolate a *B. bassiana* strain that effectively reduces *B. tabaci* populations.

MATERIALS AND METHODS

Place and Time

This study was conducted at the Biological Control Laboratory, Department of Plant Protection, Faculty of Agriculture, Universitas Andalas, and at the Experimental Farm of Universitas Andalas in Padang from July to October 2024.

Materials and Tools

The materials used in this research were an isolate of the fungus *Beauveria bassiana*, whiteflies, Sabouraud Dextrose Agar supplemented with Yeast Extract (SDAY), tomato seeds of the Selena variety, soil, 5-kg polybags, heat-resistant plastic, 70% alcohol, distilled water, Tween 80, tissue paper, plastic wrap, millimeter paper, labels, and stationery.

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The tools used in this study included Petri dishes (9 cm in diameter), test tubes, a test tube rack, Schott bottles, a Bunsen burner, an autoclave, a laminar air flow cabinet, an oven, an analytical balance, drop pipettes, a micropipette and tips, a haemocytometer, a spatula, object glasses, cover glasses, a vortex mixer, a cork borer, pot trays, scissors, a fine brush, tweezers, an Olympus CX23 microscope, and a smartphone camera.

Method

This study employed an experimental method using a Completely Randomized Design (CRD) with six treatments and five replications. The treatments comprised five isolates of *Beauveria bassiana* and a control treatment. The treatments were as follows:

A = Control (immersion using sterile distilled water)

B = Isolate PA221 (chili root endophyte, Parabek, Agam)

C = Isolate BbWs (*Leptocorisa oratorius*, Duku, Padang Pariaman)

D = Isolate PB211 (chili stem endophyte, Parabek, Agam)

E = Isolate PD114 (chili leaf endophyte, Parabek, Agam)

F = Isolate TD312 (wheat stem endophyte, Koto Laweh, Tanah Datar)

Propagation of *Beauveria bassiana*

The *B. bassiana* fungal isolate used was obtained from the Biological Control Laboratory at the Faculty of Agriculture, Andalas University. *B. bassiana* fungi were propagated using Sabouraud Dextrose Agar Yeast (SDAY) media. *B. bassiana* fungal propagation was carried out by transferring pure fungal cultures with a 0.7 cm diameter cork borer to a petri dish containing SDAY media and incubated at 28 °C for 21 days until fungal conidia were formed and ready for use.

Preparation of *Beauveria bassiana* Suspension

Conidia of a 21-day-old *B. bassiana* fungal isolate were collected by adding 10 ml of sterile distilled water and three drops of Tween 80 solution as a leveling agent to a petri dish. All ingredients were mixed using a soft brush to loosen the conidia, then transferred to a test tube and homogenized using a vortex. Next, serial dilutions were performed up to 3 times, and the final conidial concentration was 10^8 conidia/mL. The density of fungal conidials was calculated using a hemocytometer (Improved Neubauer) (González-Guzmán et al., 2021; Ibrahim et al., 2022).

Preparation of Planting Media

The planting medium consisted of a 2:1 (v/v) mixture of sterile soil and manure. The soil and manure mixture was sterilized using the Tyndallization method, which involves heating the mixture to 100°C for 1 hour in a steamer, followed by a 24-hour incubation, then reheating the mixture three times at 100°C. The sterilized soil was placed in 5 kg polybags and was ready for use as a planting medium (Gullino et al., 2022)

Treatment of Tomato Seeds with *B. bassiana* Fungus

The tomato seeds used in this experiment were of the Selena variety. Before treatment, the seeds were surface-sterilized by sequential soaking: first in sterile distilled water for 1 minute, then in 70% ethanol for 1 minute, and finally rinsed again in sterile distilled water for 1 minute. After sterilization, the seeds were air-dried for 10 minutes inside a Laminar Air Flow Cabinet. Once dry, the seeds were immersed in a 10 mL suspension of *B. bassiana* at 10^8 conidia/mL for 6 hours, as described by Trizelia (2020). After soaking, the seeds were placed in petri dishes lined with sterile filter paper and allowed to dry for one hour. The fully dried seeds were then sown into the prepared planting medium.

Seed Sowing and Seedling Planting

Tomato seeds soaked in a suspension of the fungus *B. bassiana* are planted in plastic seed trays filled with a 2:1 soil and manure mixture, which has been kept moist. The planting medium is gently hollowed out with small pieces of wood, 0.5-1 cm deep, and two tomato seeds are placed in each hole. Then, the medium is evenly covered again. Watering is performed daily under moist soil conditions (Pandey et al., 2023).

The 21-day-old tomato seedlings were transferred into polybags filled with a mixture of manure and sterilized soil. One tomato seedling was planted in each polybag. The polybags were then arranged according to the tomato planting distance, 40 cm x 60 cm (Singh & Shrama, 2022; Melo et al., 2024).

Tomato Plant Maintenance

Tomato plants are watered once a day, either in the morning or the evening. They are staked at 3 months of age. The stakes are made from 1-meter-long bamboo strips placed vertically beside the tomato plants. Fertilization is done by applying NPK fertilizer around the plants at a distance of 10 cm and a depth of 5 cm. This fertilization is performed twice: once a week after planting and again four weeks later. Weeds are removed manually every two weeks.

Observation of the *B. tabaci* Population in Tomato Plants

Observations of the *B. tabaci* population were made by counting adults, nymphs, and eggs in each treatment. Pest population data were collected every 7 days, during morning hours. Observations started when plants were 2 weeks old and continued until 5 weeks after planting. Adults were counted directly on the plants. Nymph and egg populations were estimated by sampling three leaves from the top, middle, and bottom of each plant. The numbers of nymphs and eggs were then counted in the laboratory using a microscope (Kumar et al., 2023; Smith et al., 2024).

Counting the Number of Trichomes on Leaves

Trichome counts were performed by collecting three leaves (top, middle, and bottom) from each plant for each treatment. The leaves were cut into 1 cm² pieces, and trichomes were counted under a microscope (Xu et al., 2023; Earsakul et al, 2025).

Data Analysis

The data obtained were presented in tables, and the standard error (SE) was calculated. The data were analyzed using analysis of variance (ANOVA) in Statistix 8, and when significant differences were detected, a post hoc test was performed at the 5% significance level.

RESULTS AND DISCUSSION

Population of *B. tabaci* egg

Applying *B. bassiana* to tomato plants through seed soaking significantly reduced the number of *B. tabaci* eggs across all tested fungal isolates. Analysis of variance showed a notable difference between treatments with *B. bassiana* isolates and the control group. The average number of *B. tabaci* eggs per plant at five weeks after planting is detailed in Table 1.

Table 1. Average Number of *Bemisia Tabaci* Eggs on Tomato Plants Treated with *Beauveria Bassiana* Fungus at 5 Weeks After Planting

Treatment	Egg number/plant ± SE
Control	81.10 ± 3.32 a
PA221	37.40 ± 10.69 b
PB211	35.20 ± 4.23 bc
PD114	33.65 ± 3.38 bc
BBWs	25.70 ± 4.30 bc
TD312	20.40 ± 4.84 c

Note: Means within the same column followed by the same lowercase letter are not significantly different according to LSD at the 5% level.

As shown in Table 1, all treatments with *B. bassiana* isolates significantly reduced the number of eggs laid by *B. tabaci* compared to the control. The PA221 isolate had the highest average number of eggs (37.40 per plant). All *B. bassiana* treatments significantly lowered *B. tabaci* egg populations compared to the control (Table 1). Although TD312 had the lowest egg count (20.40 per plant), it was not statistically different from BBWs (25.70), PD114 (33.65), or PB211 (35.20). Only PA221 (37.40) had a significantly higher egg count than TD312. These results show that most isolates, especially TD312, BBWs, PD114, and PB211, are similarly effective in reducing oviposition.

Kogan and Ortman (1978) noted that resistant plants can produce allelochemicals with repellent properties **to deter pests. Similarly, Mawan et al. (2013) demonstrated that treating rice**

seeds with *Nigrospora* sp. reduced both egg-laying and egg-hatching rates of the white-backed planthopper. Hendra et al. (2023) also reported that *B. bassiana* applied to rice seeds decreased the number of eggs laid by brown planthopper females. Furthermore, Prayogo et al. (2023) observed that colonization of *B. bassiana* in shallot tissues influenced oviposition preference in *S. exigua*, resulting in fewer eggs laid and lower hatch rates compared to the control (Figure 1).



Figure 1. *Bemisia tabaci* eggs on tomato plant leaves

***B. tabaci* Nymph Population**

The seed-soaking application of *B. bassiana* to tomato plants significantly affected the number of *B. tabaci* nymphs. Analysis of variance confirmed that treatments with *B. bassiana* isolates resulted in a statistically significant decrease in nymph counts compared to the control. The mean number of *B. tabaci* nymphs per plant at five weeks after planting is shown in Table 2.

Table 2. Average Number Of *Bemisia Tabaci* Nymphs On Tomato Plants Treated With *Beauveria Bassiana* Isolates At 5 Weeks After Planting

Treatment	Nymph/plant) \pm SE
Control	22.15 \pm 7.69 a
PD114	14.65 \pm 4.50 ab
BBWs	12.50 \pm 2.19 ab
PB211	12.25 \pm 3.00 ab
PA221	9.10 \pm 2.74 b
TD312	7.25 \pm 1.31 b

Note: Means within the same column followed by the same lowercase letter are not significantly different according to LSD at the 5% level.

All *B. bassiana* treatments reduced nymph populations compared to the control, though not all reductions were statistically significant (Table 2). PD114 had the highest nymph count among treated plants (14.65 per plant), while TD312 had the lowest (7.25 per plant).

Statistical analysis showed that PD114, BBWs (12.50), and PB211 (12.25) were not significantly different from the control. In contrast, PA221 (9.10) and TD312 (7.25) were significantly lower than the control. However, PA221 and TD312 were statistically similar to PD114, BBWs, and PB211.

These findings indicate that while all isolates suppressed nymph populations to some extent, only PA221 and TD312 demonstrated significant suppression compared to the control, with TD312 showing the numerically lowest nymph count.



Figure 2. *Bemisia* nymph on a tomato plant leaf

The lower population of *B. tabaci* nymphs on *B. bassiana*-treated tomato plants can be attributed to a reduction in egg laying by adult females, as well as a decreased egg hatching rate. The application of *B. bassiana* likely contributes to both factors by producing repellent compounds within colonized plant tissues, which deter oviposition and interfere with embryonic development. According to Flowerina (2021), *B. bassiana* can establish endophytic colonization in tomato plants, inhabiting roots, stems, and leaves, with the highest colonization observed in leaf tissues. This internal presence of *B. bassiana* has been shown to reduce pest populations and enhance plant resistance to insect pests under field conditions (Saragih et al., 2018). In support of this, Abdullah et al. (2020) reported that rice plants treated with *B. bassiana* via seed soaking experienced mortality of *Nephotettix virescens*, further demonstrating the fungus's potential to suppress pest populations through endophytic activity.

Population of *B. tabaci* adult

The application of the fungus *B. bassiana* reduced the number of *B. tabaci* adults on tomato plants compared to control plants. Analysis of variance results showed that all tested isolates were significantly different from the control. The population of *B. tabaci* adults on tomato plants, aged 5 weeks after planting (WAP), is shown in Table 3.

Table 3. Average number of *Bemisia tabaci* adults on tomato plants to which the application of *Beauveria bassiana* fungus at the age of 5 WAP

Treatment	Adult/plant) \pm SE
Control	10.00 \pm 1.90 a
PA221	5.35 \pm 1.96 b
BBWs	3.75 \pm 0.53 b
PB211	3.70 \pm 0.50 b
PD114	3.45 \pm 0.76 b
TD312	2.30 \pm 0.37 b

Note: Numbers followed by the same lowercase letter in the same column are not significantly different according to LSD at the 5% level.

All *B. bassiana* treatments significantly reduced *B. tabaci* adult populations compared to the control (Table 3). Although TD312 showed the numerically lowest adult count (2.30 per plant), it was statistically similar to PA221 (5.35), BBWs (3.75), PB211 (3.70), and PD114 (3.45). These results indicate that all isolates are equally effective at suppressing adult *B. tabaci* populations. The morphology of *B. tabaci* adults is shown in Figure 3.



Figure 3. Adults of *Bemisia tabaci* on tomato leaves

The study results indicate that treating tomato seeds with *B. bassiana* enhanced plant resistance against *B. tabaci*. This was demonstrated by lower nymph and adult populations observed across the four *B. bassiana* isolate treatments compared to the control. The reduced nymph population is attributed to decreased egg deposition, while the adult population was also suppressed in *B. bassiana*-treated plants. Seed treatment with *B. bassiana* enhanced tomato resistance against *B. tabaci*. All isolates significantly reduced egg and adult populations, while only PA221 and TD312 significantly suppressed nymphs. TD312 consistently showed the lowest numerical pest counts across all stages, though statistically similar to several other isolates. These findings support previous studies (Trizelia et al., 2020) on the effectiveness of *B. bassiana*, particularly isolate TD312, in suppressing pest populations through endophytic colonization and induced resistance. Hendra et al. (2023) observed that *B. bassiana* application to rice seeds reduced the percentage of brown planthopper adults. Additional studies support these results: Akello et al. (2008) found that soaking banana roots in *B. bassiana* reduced *Cosmopolites sordidus* populations by up to 88.9% and plant damage by 86.7%, and Saragih (2024) reported that *B. bassiana* applications suppressed the population growth and attack intensity of *B. tabaci*, *M. persicae*, and *Aphis gossypii* on chili plants. Overall, *B. bassiana* colonizing plant tissues can indirectly induce resistance to insect pests.

The protective mechanism of endophytic fungi against pests may involve qualitative and quantitative changes in plant nutrition—such as alterations in carbohydrate, nitrogen, and phytosterol content (Grabka et al., 2022)—as well as morphological and physiological modifications in the plant (Gao et al., 2010). Inoculation with *B. bassiana* can elevate the levels of various defense-related compounds, including ethylene, chitinase, phytoalexins, alkaloids, jasmonic acid, and salicylic acid, which are critical in pest development (Vlot et al., 2009). For instance, Hendra (2022) reported that *B. bassiana* treatment in rice increased salicylic acid and oxalic acid content while reducing sucrose levels in stems. Salicylic acid is essential for plant defense against pests (Moran & Thompson, 2001), and oxalic acid inhibits phloem feeding by planthoppers and is toxic to insects at high concentrations (Korth et al., 2006). According to Yoshihara et al. (1980), oxalic acid is a key factor in rice resistance to planthoppers. Endophytic *B. bassiana* can also be directly toxic to insects by producing feeding deterrents or antibiotics (Bing and Lewis, 1992; Vega et al., 2008).

Antibiosis is one mode of action in which ingested plant tissues or fluids adversely affect insects, leading to mortality, growth reduction, developmental failure, or behavioral abnormalities (Fatahuddin et al., 2003). *B. bassiana* exhibits antibiosis by producing compounds that inhibit pest growth and development, often through the induction of plant secondary metabolites such as alkaloids, flavonoids, terpenoids, phenolics, and toxins (Schulz et al., 1999). Its secondary metabolites can act as bioinsecticides, disrupting insect metabolism and immune responses (Ávila-Hernández et al., 2020). Azevedo et al. (2000) linked pest control effects to the toxin-producing capacity of endophytes, and Calhoun et al. (1992) demonstrated that endophytic fungi synthesize toxins that inhibit *Choristoneura fumifera* growth. Additionally, endophytes can indirectly enhance resistance by stimulating the production of defense-signaling compounds, such as salicylic acid, jasmonic acid, and ethylene (Gao et al., 2010).

Trichome Density in Tomato Plants

Application of *B. bassiana* through seed soaking significantly influenced trichome density on tomato leaves (Table 4). TD312 showed the numerically highest trichome density (616.73 per cm²), followed by PA221 (597.66), BbWs (507.13), PD114 (439.26), and PB211 (392.40), while the control had the lowest density (295.73).

Table 4. Number of trichomes on tomato plant leaves

Treatment	Trichome Density (cm ²) ± SE
TD312	616.73 ± 81.71 a
PA221	597.66 ± 60.63 a
BbWs	507.13 ± 82.58 ab
PD114	439.26 ± 56.26 abc
PB211	392.40 ± 28.43 bc
Control	295.73 ± 33.79 c

Note: Means within the same column followed by the same lowercase letter are not significantly different according to LSD at the 5% level.

B. bassiana application significantly affected trichome density on tomato leaves (Table 4). TD312, PA221, BbWs, and PD114 showed significantly higher trichome density than the control, with TD312 and PA221 having the highest numerical values. These four isolates were statistically similar to each other. PB211 had moderate trichome density, statistically similar to both PD114 and the control. Enhanced trichome density, particularly in TD312, PA221, BbWs, and PD114, may strengthen plant physical barriers against *B. tabaci*

The influence of *B. bassiana* on plant morphology, including increased trichome density, is likely due to its endophytic colonization, which may stimulate the production of plant growth regulators such as auxin, cytokinin, and gibberellin. According to Halo et al. (2023), endophytic fungi can increase trichome density on tomato leaves by secreting gibberellins, thereby enhancing drought resistance. Similarly, Qin et al. (2021) reported that *B. bassiana* colonization in tobacco plants promotes growth by enhancing photosynthetic rate, chlorophyll content, and trichome and stomatal density. High trichome density is a key physical trait associated with pest resistance, as dense trichomes can impede *B. tabaci* females from penetrating the leaf epidermis and inserting their stylets to feed. Consequently, leaves with dense trichomes are less preferred for oviposition and feeding. In this study, tomato plants treated with *B. bassiana* isolates showed higher trichome density and lower adult *B. tabaci* populations compared to the control. This aligns with findings by Flowerina (2021), who reported that increased trichome density in tomato plants enhances resistance to *B. tabaci* infestations.

CONCLUSION

Seed treatment of tomato plants with *Beauveria bassiana* effectively suppressed *Bemisia tabaci* populations. Isolate TD312 resulted in the lowest populations of eggs (20.40 per plant), nymphs (7.25 per plant), and **adults (2.30 per plant) compared to the control (81.10, 22.15, and 10.00 per plant, respectively). Additionally, application of *B. bassiana* affected plant morphology by increasing trichome density. TD312 isolates increased trichome density to 616.73/cm², which was higher than the control (295.73/cm²), potentially enhancing antixenosis (non-preference) mechanisms against *B. tabaci*. Although statistically comparable to several other isolates on some parameters, TD312 consistently exhibited the lowest pest counts and the highest trichome density, indicating its potential to enhance tomato resistance to *B. tabaci* by suppressing pest populations and reinforcing trichome density. Further studies are needed to investigate the secondary metabolites involved in induced resistance.**

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