Co-Inoculation of *Sechium edule* (Jacq.) Sw. Plants with *Rhizophagus intraradices* and *Azospirillum brasilense* to Reduce *Phytophthora capsici* Damage

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1. Introduction

Agricultural systems use large amounts of external inputs, such as agrochemicals, to achieve their production goals. High energy consumption, especially of a fossil nature, is used in this process [1]. This agricultural production model tends to be unsustainable, and consequently, alternative inputs are required even more so when considering that the demand for food produced without agrochemicals has increased [2,3], which requires the use of natural components to control growth, yield, and diseases in plants. In this context, special interest has been focused on improving crop nutrition and protection through the inclusion of microorganisms capable of establishing symbiosis with the root system of the host plant and favoring its functioning, especially under stress conditions.

Beneficial microorganisms have been considered for use as biological fertilization agents or biofertilizers, which the Ministry of Agriculture, Livestock, Rural Development, Fisheries and Food (SAGARPA) [4] has promoted in Mexico due to evidence that they increase annual crop yields and reduce carbon dioxide emissions. The biofertilization of various microorganisms favors the nutrition of the host plant through various mechanisms of action and, concomitantly, reduces the severity of the attack by radical pathogens, as.
reported with *Phytophthora capsici* in horticultural plants when endomycorrhizal fungi are applied [5,6].

In Mexico, the production of chayote (*Sechium edule* (Jacq.) Sw.) (Cucurbitaceae), a vegetable for export, ranges from 54 [7], to 130–136 t ha⁻¹ [8], and generates more than 27,700 local jobs for every 35 ha yr⁻¹, which increases its importance as a social crop [9].

In general, the planting density in technified orchards of *S. edule* in mesophilic forest conditions (1200–1400 m) ranges from 100 to 130 plants ha⁻¹, and one plant produces up to 1.0 t ha⁻¹ in a period of 6 to 8 months [9].

The planted area in Mexico is 2953 ha [7], and in all regions, *P. capsici* has been recorded as one of the fungi responsible for stem rot (vital node) and chayote root rot (*S. edule* (Jacq.) Sw.), thus showing, in addition, different degrees of aggressiveness [10,11].

With this background, the effectiveness of co-inoculation of *Sechium edule* plants (chayote) with *Rhizophagus intraradices* and *Azospirillum brasilense* to reduce *P. capsici* damage was evaluated. We considered the hypothesis that co-inoculation can increase the nutrition of *S. edule* plants and reduce the damage to or loss of plants caused by *P. capsici*, attributed to the effect of competition for infection sites, by acting as a biocontrol.

### 2. Materials and Methods

The present research was carried out in the greenhouse of the Rosario Izapa Experimental Field of the National Institute for Forestry, Agricultural, and Livestock Research (INIFAP) in Tuxtla Chico, Chiapas (14°58′28.89″ LN, 92°09′19.64″ LO and 435 m altitude). The climate is generally warm and humid, with the influence of summer rains and monsoon. The average annual rainfall is 4720 mm and the average temperature, considered isothermal, is 25.4 °C.

#### 2.1. Plant Material

Seeds of Costa Rica-type chayote (*S. edule var. virens levis*), provided by the Interdisciplinary Research Group of *Sechium edule* in Mexico, from the area of conservation and genetic improvement, were used (19°08′48″ N, 97°57′00″ W, at 1340 m). The fruits were cut between 36 and 38 days after anthesis—that is, 18–20 days after horticultural maturity [12]—and transported to the experimental site one day later. Due to the recalcitrant and endocarpic nature of the seed, the fruit of *S. edule* containing the seed is normally planted. Sowing was carried out two days after cutting.

#### 2.2. Experimental Setup, Substrate, and Microorganisms

One chayote seed (one fruit) was planted per bag. The substrate used was a mixture of andosol molic soil and washed river sand in a 1:1 (v/v) ratio. With this, 6.0 kg polyethylene bags, perforated at the bottom to promote drainage, were filled, and randomly distributed on metal tables at a height of 1.0 m from the ground. Physicochemically, the substrate had the following components and characteristics: sand, 82.76%; silt, 11%; clay, 6.24%; crumbly sand texture; organic matter, 2.85%; pH, 5.50; N, 0.13%; P, 7 mg kg⁻¹; K, 13 mg kg⁻¹; Na, 16 mg kg⁻¹; Mg, 12 mg kg⁻¹; Ca, 96 mg kg⁻¹; and electrical conductivity (EC), 0.055 dS m⁻¹.

#### 2.3. Microorganisms

Co-inoculation can enhance plant growth through mechanisms attributed to the positive effects of *Azospirillum*, which are mainly attributed to improved root development and mycorrhiza-improved nutrient status of the host plant, mainly with the transportation of P and the bioprotective effect of mycorrhization against soil-borne fungal pathogens. The co-inoculation of *S. edule* plants was justified by the expected advantages of improving plant nutrition under stress conditions due to *P. capsici* infection in addition to inducing competition for infection sites in the roots. In this regard, Ju et al. [13], mention the benefits of co-inoculation with rhizobacteria and *rhizobium* in alfalfa plants, observing a reduction
in oxidative damage and an improvement in the antioxidation capacity of the plant in addition to observing less microbial diversity in the rhizosphere.

*Rhizophagus intraradices* (Schenck & Sm.) Walker & Schuessler (RI) was acquired at the INIFAP Campo Experimental Rosario Izapa, Chiapas, with 40 spores g$^{-1}$ of soil plus mycelium and rootlets as sources of inoculum.

*Azospirillum brasilense* Tarrand, Krieg, & Döbereiner (AB) was obtained at the Benemérita Universidad Autónoma de Puebla (BUAP) with $9 \times 10^6$ bacteria g$^{-1}$ of peat (information contained in the products). The *P. capsici* (PC) strain was provided by the Colegio de Postgraduados, Campus Montecillo, and cultured from a purified hyphal tip in V8 agar culture medium.

### 2.4. Application of Microorganisms

At the time of sowing, 20 g of the biofertilizer of each of the microorganisms was applied in the cavity of the corresponding treatment. The base of the fruit with the seed was in direct contact with the biofertilizers. Subsequently, two-thirds of the fruit was covered with the same substrate to avoid burning. The plants were irrigated with distilled water.

The application of *P. capsici* was inoculated at the base of the stem 14 days after planting (DAP). For this purpose, the fully sporulated strain with sporangium formation was separated from the medium with sterile distilled water and was gauged at 150 mL and refrigerated for 15 min to induce the release of zoospores. Subsequently, it was allowed to rest at room temperature for 2 min, and from this preparation, 2 mL plant$^{-1}$ was applied at the base of the stem (vital node).

### 2.5. Treatments, Repetitions, and Experimental Design

The treatments included the inoculation of the microorganisms alone or combined, plus the control, with and without the application of *P. capsici*. A total of eight treatments were distributed completely at random with four repetitions. Irrigation was carried out every third day at the rate of one liter of water per plant (Table 1).

**Table 1.** Description of the treatments and abbreviations used in the study.

| S. No | Treatments | Abbreviation |
|-------|------------|--------------|
| 1     | Control    | CL           |
| 2     | *P. capsici* | PC         |
| 3     | *R. intraradices* | RI          |
| 4     | *A. brasilense* | AB          |
| 5     | *R. intraradices* + *A. brasilense* | RI + AB     |
| 6     | *R. intraradices* + *P. capsici* | RI + PC    |
| 7     | *A. brasilense* + *P. capsici* | AB + PC    |
| 8     | *R. intraradices* + *A. brasilense* + *P. capsici* | RI + AB + PC |

### 2.6. Morphological and Physiological Variables

The morphological variables, including the number of leaves at 28, 56, and 84 DAP, the mean number of vines and the average length of four vines at 84 DAP, were recorded. The aerial and root parts of the plant were harvested at 84 days, and they were weighed on a semi-analytical balance (Ohaus Adventurer Pro, California USA) after drying in a forced air oven at 60–75 °C to a constant weight. The leaf area (cm$^2$) was measured using a leaf area integrator (LI-COR, LI 3000, LI-COR Biosciences, Lincoln, NE, USA).

### 2.7. Histological Sections and Microscopy

Histological sections were made on a rotary microtome (Spencer, American Optical Company model 820, NY USA) and cut transversely to a thickness of 15 µm. The cut sections were placed in a flotation bath with hot water (43–45 °C) and gelatin. Finally, they were mounted on glass slides sequentially. They were deparaffinized by melting the paraffin and eliminating the excess contained in the tissues by means of three changes of
xylol at intervals of 3 min each, and they were hydrated with two changes of ethyl alcohol, both absolute and at 96%, for 3 min in both cases. At the end of this procedure, the sections were placed in safranin (SIGMA-ALDRICH, St. Louis, MO, USA) for 24 h [14].

The sections remained for 24 h in safranin, after which they were rinsed three times with running water for 25 min in each rinse. Next, they were washed with picric acid-ethanol 96%, ammonia-ethanol 96% and absolute ethanol. Fast green was added for five seconds, followed by clove oil with a short exposure time. To remove excess dye (fast green), a mixture of clove oil–xylol–absolute alcohol was used; the sections were rinsed in xylol for 3 min and finally mounted in synthetic resin for observation [10]. Scanning electron microscopy was performed according to the protocol of Bozzola and Russell [15].

2.8. Statistical Analysis

The results were evaluated using the program SAS (Software as a Service) version 8.1 (SAS 1999-2000) [16] based on a completely randomized analysis. Differences between treatment means were compared using Tukey’s test ($p \leq 0.05$).

3. Results

3.1. Morphological Components

The initial number of leaves (28 DAP) increased 35% more compared to the control when inoculating $S. edule$ with $R. intraradices$ or $A. brasilense$ together with $P. capsici$. However, when co-inoculated with $R. intraradices + A. brasilense$ and $P. capsici$, the number of leaves was similar to that of the control, without statistical difference (Figure 1). On the other hand, at 56 DAP, the co-inoculation of the two beneficial microorganisms in interaction with $P. capsici$ induced an increase of 24% in the number of leaves compared to the control.

At 84 DAP, the most notable increase in the number of leaves was presented in the biofertilized treatments with $R. intraradices$, counting 52 leaves; in $A. brasilense$, counting 50 leaves; and in the double biofertilization ($R. intraradices + A. brasilense$), counting 40 leaves, and they were statistically different ($p \leq 0.05$) to the other treatments. The lowest number of leaves occurred in the control and in the treatments where $P. capsici$ was inoculated (Figure 1).

The length of the vines was shorter in the controls with and without $P. capsici$, at an average of 17 cm. In the case of the treatments biofertilized with $R. intraradices$ and/or $A. brasilense$ alone plus the inoculation of $P. capsici$, a reduction in the length of the vines of up to 70% was also recorded, compared to the same treatments without $P. capsici$. In the case of double biofertilization, the average length of the vines was 56 cm and when $P. capsici$ was inoculated, it was 15 cm (Figure 2). It should be noted that introducing $P. capsici$ in the treatments colonized by $R. intraradices$ led to the development of one more vine compared to treatment with $A. brasilense$ and the double symbiosis.
3.2. Physiological Components

The allocation of dry matter to different components of the plants showed changes over time between treatments. The application of *P. capsici* on plants biofertilized with the microorganisms, *R. intraradices* and *A. brasilense*, decreased the biomass of *S. edule* by 8%
compared to the same treatments without the application of the pathogen. This may be the effect of the increase observed in the necrotic root system of plants inoculated with P. capsici. However, compared to the average biomass of the controls with and without P. capsici, the biomass of biofertilized plants inoculated with P. capsici was 230% higher (Table 2).

Table 2. Comparisons of means of dry weight of root, stem, leaf blade, petiole, and leaf area of S. edule var. virens levis Costa Rica Type, biofertilized with A. brasilense and R. intraradices and inoculated with P. capsici in an andosol-molic soil from Soconusco, Chiapas.

| Days | Treatment       | Root (g Plant) | Stem and Vine | Leaf Lamina | Petiole | Leaf Area (cm² Plant) |
|------|----------------|----------------|---------------|-------------|---------|----------------------|
| 28   | CL             | 1.55 ± 0.09 ab | 2.75 ± 0.15 a | 2.52 ± 0.31 ab | 0.32 ± 0.02 a | 868 ± 182 a |
|      | PC             | 0.85 ± 0.13 c  | 1.22 ± 0.11 cd | 1.25 ± 0.32 b | 0.14 ± 0.05 a | 357 ± 23 c  |
|      | RI             | 1.70 ± 0.05 ab | 1.77 ± 0.08 bc | 2.57 ± 0.10 ab | 0.32 ± 0.02 a | 821 ± 86 a  |
|      | AB             | 1.30 ± 0.1 abc | 2.75 ± 0.14 a | 2.67 ± 0.36 a | 0.35 ± 0.02 a | 756 ± 69 ab |
|      | RI + AB        | 1.05 ± 0.10 bc | 2.0 ± 0.20 ab  | 2.70 ± 0.30 a | 0.37 ± 0.06 a | 982 ± 66 a  |
|      | RI + PC        | 1.85 ± 0.38 a  | 0.83 ± 0.06 d  | 1.42 ± 0.19 ab | 0.22 ± 0.04 a | 350 ± 65 d  |
|      | AB + PC        | 0.95 ± 0.05 bc | 0.97 ± 0.14 d  | 1.25 ± 0.33 b | 0.27 ± 0.02 a | 305 ± 38 c  |
|      | RI + AB + PC   | 1.53 ± 0.10 abc| 2.40 ± 0.29 ab | 2.10 ± 0.30 ab | 0.30 ± 0.04 a | 490 ± 48 bc |
| CV%  |                | 23.9           | 17.9          | 28.1        | 31.2     | 21.2                 |
| 56   | CL             | 0.55 ± 0.06 c  | 0.85 ± 0.13 a | 2.57 ± 0.15 ab | 0.37 ± 0.04 bc | 734 ± 195 bc |
|      | PC             | 0.42 ± 0.02 c  | 1.35 ± 0.15 a | 2.10 ± 0.21 b | 0.30 ± 0.04 bc | 560 ± 78 c  |
|      | RI             | 1.52 ± 0.17 a  | 0.97 ± 0.04 a | 2.75 ± 0.08 ab | 0.47 ± 0.11 ab | 922 ± 148 ab |
|      | AB             | 0.92 ± 0.04 bc | 1.20 ± 0.15 a | 2.80 ± 0.23 ab | 0.40 ± 0.04 bc | 655 ± 72 ab |
|      | RI + AB        | 1.72 ± 0.27 a  | 2.02 ± 0.26 a | 3.40 ± 0.42 a | 0.55 ± 0.05 ab | 1060 ± 55 a |
|      | RI + PC        | 0.62 ± 0.07 c  | 1.00 ± 0.07 a | 1.77 ± 0.30 b | 0.20 ± 0.04 c | 333 ± 17 c  |
|      | AB + PC        | 1.47 ± 0.06 ab | 2.10 ± 0.57 a | 3.35 ± 0.32 a | 0.67 ± 0.02 a | 775 ± 82 bc |
|      | RI + AB + PC   | 0.75 ± 0.02 c  | 2.12 ± 0.41 a | 2.65 ± 0.15 ab | 0.35 ± 0.02 bc | 583 ± 91 c  |
| CV%  |                | 24.5           | 39.1          | 19.3        | 27.0     | 13.4                 |
| 84   | CL             | 0.82 ± 0.13 b  | 1.55 ± 0.43 bc | 3.97 ± 0.48 bc | 0.60 ± 0.14 bc | 1303 ± 534 bc |
|      | PC             | 0.60 ± 0.20 b  | 1.00 ± 0.14 c | 3.30 ± 0.09 c | 0.32 ± 0.04 c | 819 ± 142 c |
|      | RI             | 1.97 ± 0.04 a  | 5.75 ± 0.42 a | 8.82 ± 1.09 ab | 1.02 ± 0.06 ab | 3405 ± 391 a |
|      | AB             | 1.92 ± 0.08 a  | 6.75 ± 0.23 a | 9.35 ± 0.47 a | 0.97 ± 0.10 ab | 3207 ± 349 a |
|      | RI + AB        | 1.67 ± 0.25 a  | 4.42 ± 0.39 a | 9.70 ± 1.07 a | 1.25 ± 0.05 a | 3456 ± 183 a |
|      | RI + PC        | 0.85 ± 0.15 b  | 4.07 ± 1.13 ab | 7.22 ± 2.01 abc | 0.77 ± 0.07 abc | 1335 ± 257 bc |
|      | AB + PC        | 0.55 ± 0.06 b  | 6.80 ± 0.39 a | 8.30 ± 1.23 abc | 1.02 ± 0.06 ab | 1403 ± 9 b |
|      | RI + AB + PC   | 0.75 ± 0.12 b  | 4.90 ± 0.42 a | 5.77 ± 0.31 abc | 0.75 ± 0.08 abc | 122 ± 117 bc |
| CV%  |                | 26.0           | 27.5          | 29.8        | 24.9     | 11.8                 |

Values with the same letter within each factor and column are equal according to Tukey’s test at p ≤ 0.05. CV: coefficient of variation (%).

The initial growth of S. edule assigned an average of 25, 32, 37, and 6% photosynthates to the root, stem plus vines, leaf blade, and petiole, respectively. In the second sampling (56 DAP), there was an increase in the allocation of biomass in the physiological components of yield, expressed in the leaf blade, representing about 50% of the total plant, which decreased the biomass in the root system. This growth in S. edule was also observed at 84 DAP, observing that the yield components with the greatest changes were the leaf blade and the root system. When P. capsici was applied, root growth decreased in all treatments during the three samplings and was even more restricted in the controls with P. capsici. In this treatment, the number of roots with external necrosis increased. The total number of plants killed by necrosis was three for the CL treatment and one that was biofertilized with AB + PC (Figures 3 and 4).
Figure 3. Sequence of damage in the basal vine (vital node) of *S. edule var. virens levis* Costa Rica Type. (A–C) 22 days after inoculation with *P. capsici*. (D) Transversal histological section of the stem, with coenocytic mycelial hyphae of *P. capsici* in the vascular bundle 30 days after inoculation. (E) Total necrosis of the adult plant basal guide taken in the field.

Figure 4. *Phytophthora capsici* on seed root tissue of *S. edule var. virens levis* Costa Rica Type. (A, B) Sporangia and zoospores. (C) In germinated seed root tissue. Scanning electron microscope micrographs. 500×, 3000×, 2500×, respectively.
4. Discussion

Dead plants were recorded in the CL treatment, as well as one dead plant in the AB + PC treatment, even though it was expected that in PC, all the plants would die; nevertheless, lower values were observed in the length of the vines (Figure 2). The AB, RI, and RI + AB treatments stood out, demonstrating their possible effect on plant nutrition. However, the RI + PC, AB + PC and RI + AB + PC plants showed survival and medium to low values of vine length, without registering plant loss.

Similar behavior was observed for the number of leaves (Figure 1), where the plants inoculated with RI, AB, and RI + AB, were outstanding compared to those inoculated with PC.

The foregoing is relevant, as successful cultivation of *S. edule* is identified by the formation of a canopy of vines that protects the youngest and most productive vines, since it is grown in a horizontal structure. *S. edule* produces one to two fruits per node longitudinally; therefore, the longer the vines are, the greater the number of fruits will be [8]. In terms of plant survival in the field, RI + PC treatment, as well as AB + PC, is relevant for its possible protective effect.

In other tropical perennial crops, such as *Theobroma cacao* L. and *Coffea arabica* L., without application of *P. capsici*, increases in the number of leaves are cited when endomycorrhizal fungi or nitrogen-fixing bacteria such as *A. brasilense* are applied [17]. In relation to the number of vines per plant, an increase was recorded in the treatments without the application of *P. capsici*, but the effect was outstanding with the individual biofertilization of *R. intraradices* and *A. brasilense* compared to when the two microorganisms were applied together.

The aforementioned response is attributed to the improvement in the sanitary condition of plants biofertilized with endomycorrhizal fungi by reducing the root tissue necrotized by *P. capsici* [18]. Endomycorrhizal fungi, when interacting with radical pathogens, trigger a host plant defense response by stimulating phytoalexins, chitinases, arginine, isoflavonoids, and glucanolytic enzymes [19].

In the case of root dry weight, at 28 DAP, an increase of 42% was observed in the *R. intraradices + P. capsici* treatment; however, the most contrasting increase was recorded with the individual or combined biofertilization with microbial biofertilizers at 56 and 84 DAP.

Stem dry weight presented the most contrasting changes in biomass at 84 DAP; in this case, all treatments with biofertilizers, alone or in combination, and with *P. capsici* presented the highest values and were statistically different from the controls (p ≤ 0.05). In other tropical plants such as *T. cacao* and *C. arabica*, increased stem growth was observed when biofertilizing with *R. intraradices* [20].

The effect of increased growth of biofertilized plants is attributed, on the one hand, to the ability of *A. brasilense* to induce the growth of the root system, or to the external growth of the mycelium due to *R. intraradices*, which serves as an extension of the root system, thus contributing to the absorption of nutrients, such as phosphorus [21]. On the other hand, it is attributed to the bioprotection and fortification of the cell wall by the increased production of polysaccharides for lignification [22], which hinders the penetration of the pathogen [23]. A precursor of lignification in vegetables is β-glucosidase, which was identified in *S. edule* [24], and is involved in several functions in plants, forming part of the cell wall [25,26].

The above effects may be related to the benefits that occur in plant–microorganism symbiosis. In the case of *Azospirillum*, the root system of plants is increased through the production of growth-regulating substances [27], and the increase in root biomass favors the ability to explore and transport nutrients to the host plant [28]. Authors such as Akhtar and Siddiqui [29], observed a delay in the development of *Phytophthora* with mycorrhizal colonization and added that the pathogen did not penetrate the arbuscle. This structure is responsible for the release of nutrients into the cell [30], being unaltered, the flow of nutrients is maintained, which results in an increase in the biomass of biofertilized plants.
In *Ocimum basilicum*, Copetta et al. [31], cited an increase in phenols in the tissue when colonized by endomycorrhizal fungi, as well as an increase in the production of root hairs. In *Carica papaya* L. cv. Sunrise inoculated with *Glomus mossea*, a decrease was observed in the effect of *Fusarium*, *Pythium*, *Phytophthora*, *Rhizoctonia*, *Verticillium*, and root-knot nematodes (*Meloidogyne* spp.) or lesion nematodes (*Pratylenchus* sp.) [5], and the severity of infection in tomato with *Glomus macrocarpum* or *Glomus fasciculatum* after 20 days of being infected by *Fusarium oxysporum* f. *sp. Iycopersici* decreased between 75% and 78% [32].

Reyes et al. [33], warned about the differential effects of *Phytophthora* spp. by endomycorrhizal fungi isolates on *Capsicum annuum* L. In general, the defense mechanisms responsible for disease inhibition in mycorrhizal plants could also be explained by improved nutrition of the host plant due to the transportation of phosphorus, other nutrients, and water as well as competition with the pathogen for root colonization space [32].

In general, the biomass increased with the individual and combined application of *R. intraradices* and *A. brasilense*, compared to the controls with and without *P. capsici* (Table 2).

Similar results have been reported in other crops when biofertilized with endomycorrhizal fungi and/or nitrogen-fixing bacteria, such as in *C. arabica* [34], and forest species such as *Tabebuia donnell-smithii* [35], and *Cedrela odorata* [36].

The petiole was the plant structure that did not show contrasting changes in its dry weight at 28 DAP (without statistical differences); however, differences were observed at 56 and 84 DAP. The biofertilized treatment with AB + PC increased petiole biomass at 56 DAP (*p* ≤ 0.05), and at 84 DAP, the highest values in petiole biomass were in the double symbiosis treatment without *P. capsici*, but all treatments with the biofertilizers were clustered in the first statistical group and surpassed the controls (*p* ≤ 0.05). This plant structure was the most sensitive in the allocation of dry matter to biomass.

The initial leaf blade biomass was lower with the application of *P. capsici* (*p* ≤ 0.05). At 84 DAP, the highest dry matter allocation to the leaf blade of chayote was observed with the biofertilizer microorganisms and without *P. capsici*, and the lowest was observed with the controls. The leaf area was strongly affected by the inoculation with *P. capsici* from the beginning of the research. The biofertilized plants increased the leaf area and were statistically different (*p* ≤ 0.05). The effect on these increases in plants inoculated with RI, and/or AB has also been observed in *C. arabica* [17] and *Coffee canephora* [37].

In *O. basilicum* biofertilized with endomycorrhizal fungi [31], reported an increase in the number of glandular trichomes, which, in addition to being resistance barriers, are sites of accumulation of secondary metabolites [38], that block the action of pathogens and herbivores, while also regulating the biological activity of cytokinins [39,40]. The inoculation of *S. edule* plants with the microorganisms was beneficial, since even when inoculated with PC, the biomass reduction was only 27%, which is relevant, since, under field conditions, infected plants die (Figure 3E), as demonstrated by Siddiqui and Akhtar in their tomato research [41].

Alarcón et al. [42], mentioned that when inoculating *Solanum lycopersicum* L. with *Glomus mosseae*, it favored growth indicators and reduced infestation by the nematode *Meloidogyne* spp. Other authors such as Perez-Moncada [43], reported that inoculation with three types of *Glomus macrocarpum* and one of *Rhizoglomus intraradices* decreased the absorption of cadmium in Theobroma cacao L. plants, registering the above as benefits to reduce stress in the plant. Evaluations carried out [44], with *Rhizophagus aggregatus* and a consortium of *Glomus claroides*, *Rhizophagus diaphanous*, and *Paraglomus albidum* on seedlings of varieties of *Coffea arabica* L. showed that inoculation with the consortium registered the highest values in seedling growth, phosphorus assimilation, and health.

Studies carried out with *R. intraradices* [45], showed that the phosphorus concentration, water potential in leaves, photosynthetic rate, transpiration, stomatal conductance, and efficiency in the use of water increased significantly in barley plants. The results of combined inoculation with *R. intraradices* and *Paenibacillus mucilaginosus* showed that the inoculated *Poncirus trifoliat* seedlings registered higher levels of antioxidant enzymes than the non-inoculated ones did [46]. Tiwari et al. [47], working with *O. basilicum* plants
that were invaded by the nematode M. incognita, applied bioinoculants based on *Bacillus megaterium* and *B. subtilis*, showing improvements in plant growth and oil production and an outstanding reduction in infestation of the nematode. The foregoing is relevant since it presents a friendly alternative for the environment and human health to avoid the use of synthetic nematicides. This shows the advantages of the symbiotic association between plants and microorganisms, as shown in the present case with *S. edule* and *P. capsici*.

In relation to the possible greenhouse effects, the plants grew in conditions with a permanent temperature of 26–28 °C. The above based on [48], who reported that *S. edule* registers an ambient temperature of 26 °C for the exchange of gases and water relations, for the best rate of assimilation of CO₂ and 28 °C for the beginning of stomatal closure. Relative humidity was controlled at 85% to maintain adequate turgor that allows water potential without permanent wilting, which impacts stomatal conductance.

### 5. Conclusions

Plants of *S. edule* var. *virens levis* of the Costa Rica type inoculated with *P. capsici* and biofertilized with *R. intraradices* and *A. brasilense* showed a decrease of only 27% in their biomass. The petiole biomass and leaf area were the plant components with the lowest growth during the three evaluation periods in the presence of *P. capsici*. As with root biomass, other components responded differently with time, and at the end of the evaluation, the components in the stem indicated more decreases. The use of co-inoculation with symbiotic microorganisms is advantageous to improve the nutrition and bioprotection of *S. edule* plants in the presence of *P. capsici*.

### Author Contributions

Conceptualization, J.C.-I.; Data curation, M.I.A.-L.; Formal analysis, J.F.A.-M.; Investigation, G.O.-H.; Software, J.F.A.-C.; Writing—original draft, J.C.-I. All authors have read and agreed to the published version of the manuscript.

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### Conflicts of Interest

The authors declare no conflict of interest.

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