



Enhancement of clover growth by inoculation of P-solubilizing fungi and arbuscular mycorrhizal fungi

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ABSTRACT

This study evaluated the synergism between several P-solubilizing fungi isolates and arbuscular mycorrhizal fungi to improve clover (*Trifolium pratense*) growth in the presence of Araxá apatite. Clover was sown directly in plastic pots with 300g of sterilized washed sand, vermiculite and sepiolite 1:1:1 (v:v:v) as substrate, and grown in a controlled environment chamber. The substrate was fertilized with 3 g L⁻¹ of Araxá apatite. A completely randomized design, in 8 × 2 factorial scheme (eight P-solubilizing fungi treatments with or without arbuscular mycorrhizal fungi) and four replicates were used. The P-solubilizing fungi treatments consisted of five Brazilian P-solubilizing fungi isolates (PSF 7, 9, 20, 21 and 22), two Spanish isolates (*Aspergillus niger* and the yeast *Yarrowia lipolytica*) and control (non-inoculated treatment). The greatest clover growth rate was recorded when *Aspergillus niger* and PSF 21 were co-inoculated with arbuscular mycorrhizal fungi. *Aspergillus niger*, PSF 7 and PSF 21 were the most effective isolates on increasing clover growth in the presence of arbuscular mycorrhizal fungi. Greater mycorrhizal colonization resulted in greater clover growth rate in most PSF treatments. PSF 7 was the best isolate to improve the establishment of mycorrhizal and rhizobia symbiosis.

Key words: Araxá apatite, phosphate solubilization, co-inoculation, plant growth promotion, rhizosphere.

INTRODUCTION

Soil microorganisms play an important role in supplying nutrients to plants. Phosphorus (P) is the least mobile and available to plants in most soil conditions, in comparison to other major nutrients (El-Azouni 2008). Nevertheless, some microorganisms are able of mineralizing and solubilizing P from the organic and inorganic soil pools (Richardson 2001). Souchie et al. (2006) stated that P is the most limiting nutrient in tropical soils; in contrast, P-solubilizing microorganisms (PSM) and the arbuscular mycorrhizal fungi (AMF) are among the most relevant soil microorganisms.

The inoculation of PSM is a promising technique in order to increase P availability in soils fertilized with rock phosphates (Reyes et al. 2002). Several authors reported yield increases on maize (Reyes et al. 2002), sorghum (Kumar et al. 1999), and wheat (Whitelaw et al. 1997, Wahid and Mehana 2000) by inoculation of P-solubilizing fungi (PSF). Rodríguez et al. (1999) reported a greatest alfalfa growth in soil fertilized with rock phosphate and sugar beet waste through inoculation of *Aspergillus niger* (an isolate with high P solubilization potential) and *Glomus deserticola*. The synergism between these microorganisms also promoted considerable increases in P absorption and growth in clover (*Trifolium repens*) (Vassileva et al. 1998). Similarly, the

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co-inoculation of *Yarrowia lipolytica* (a P-solubilizing yeast) increased mycorrhizal colonization by *Glomus deserticola* in tomato roots (Vassilev et al. 2001). Ravnkov et al. (1999) also observed mycorrhizal colonization increasing forage leguminous roots after the inoculation of *Saccharomyces cerevisiae*. Nowadays, it is widely accepted that several PSM, besides solubilizing P, also increase mycorrhizal colonization, produce vitamins, amino acids and phytohormones (Barea et al. 1997, Arshad and Frankenberger Jr. 1998). Therefore, more studies to establish the functional compatibility among these microorganisms on improving plant growth of several species are required.

The PSF efficiency changes according to isolate and phosphate type (Souchie et al. 2005). Therefore, studies about P solubilization of rock phosphates in systems involving plants inoculated with PSM and AMF should be done. This study evaluated the synergism between several PSF and AMF isolates in improving clover growth fertilized with Araxá apatite.

MATERIALS AND METHODS

The experiment was done at Estación Experimental del Zaidín, Granada, Spain, in a controlled environment chamber. Clover (*Trifolium pratense*) was sown in plastic pots containing 300 g of substrate (washed sand, vermiculite and sepiolite, 1:1:1, v:v:v). Each component was autoclaved (120°C, 20 min) separately. Araxá apatite (12 g of P kg⁻¹) was mixed to the substrate at 3 g L⁻¹.

A completely randomized design, in 8 × 2 factorial scheme (eight inoculation treatments and presence or absence of AMF) and four replicates were used. The inoculation treatments were: 1) control (non-inoculated); 2) PSF 7; 3) PSF 9; 4) PSF 20; 5) PSF 21; 6) PSF 22; 7) *Aspergillus niger* and 8) *Yarrowia lipolytica*. The isolates *Aspergillus niger* and *Yarrowia lipolytica* belong to the Estación Experimental del Zaidín collection and are considered to be isolates with high P-solubilization potential (Vassileva et al. 1998, 2000). PSF 7 and 9 were isolated from the rhizosphere of *Mimosa caesalpinifolia* grown in an Argisol collected at an Atlantic Forest area in Paraty, Rio de Janeiro, Brazil. PSF 20, 21 and 22 were obtained from the rhizosphere of *Acacia holosericea* grown in Planosol collected at Seropédica, Rio de Janeiro, Brazil.

The mycorrhizal inoculum consisted of soil containing spores, mycelia and root fragments colonized by *Glomus clarum* and *Glomus geosporum*. Equal proportions were prepared by mixing 300 mL of soil containing 8 and 20 spores per mL⁻¹ of soil of *G. clarum* and *G. geosporum*, respectively. Both AMF were multiplied in sorghum and maize trap plants and maintained as a stock culture. Prior to sowing, an amount of 5 mL of inoculum was added to each pot and homogenized with the substrate. In the control, 2 mL of an inoculum filtrate, obtained by mixing 1 mL of both *G. clarum* and *G. geosporum* inoculum filtrates without AMF propagules, were added to the plant basis in each pot.

Four days after emergence, seedlings were thinned to one plant per pot. Each plant was inoculated with 1 mL of *Rhizobium trifolii* (strain 2152). This strain was incubated in Ty liquid medium (Beringer 1974), to 10⁸ CFU mL⁻¹. Two days after *R. trifolii* inoculation, the plants were inoculated with PSF treatments. These PSF isolates were grown in Petri dishes (four days, 28°C, darkness), with GL solid medium (Sylvester-Bradley et al. 1982) covering the Petri dish basis. Spores were suspended in water + tween (1%) solution, quantified following the successive dilution technique till 10⁸ CFU mL⁻¹ and, subsequently, 1 mL of spore suspension was applied to each pot.

The experiment lasted 60 days in a controlled environment chamber with 50% relative humidity, day and night temperature of 27 and 18°C, respectively, and a photoperiod of 14 hours, with photosynthetic photon flux density of 503 μmol m² s⁻¹. Twice a week, 25 mL of a modified nutrient solution (Hewitt 1966) (without N, 10% of P and pH 5.5) were added to each pot.

Forty days after emergence, shoot dry matter was determined. Sixty days after emergence, plants were harvested and shoot dry matter, colonized root length (Tennant 1975, Giovannetti and Mosse 1980), number of nodules, N and P shoot contents were evaluated. Data were submitted to ANOVA and the averages compared by the F and Tukey tests ($p \leq 0.05$).

RESULTS

AMF and PSF inoculation increased shoot dry matter in most treatments, except for plants inoculated with PSF 7, PSF 22 and *Y. lipolytica* (Fig. 1). PSF 7 and PSF 21 stimulated the greatest shoot dry matter pro-

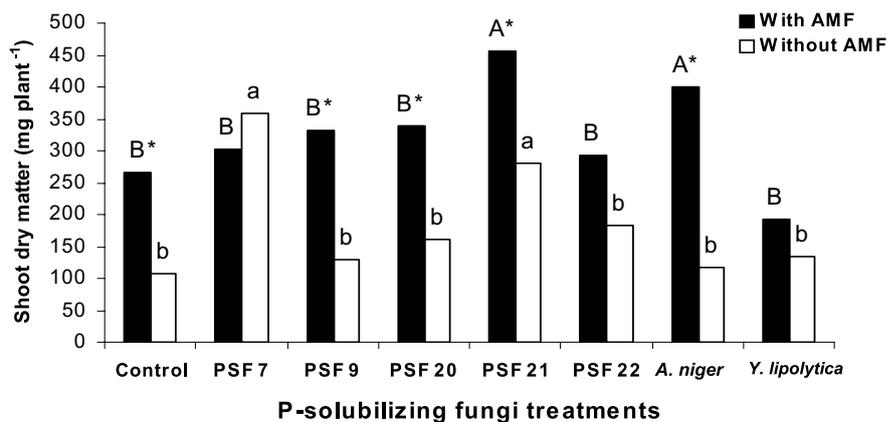


Fig. 1 – Shoot dry matter (two cuts) of clover inoculated with P-solubilizing fungi from Paraty, Rio de Janeiro, Brazil (PSF 7, 9, 20, 21 and 22) and from Granada, Spain (*A. niger* and *Y. lipolytica*), in the presence or absence of AMF. (Lower-case and upper-case letters compare PSF treatments in the absence and presence of AMF inoculation, respectively, by the Scott-Knott test ($p \leq 0.05$). *: indicates AMF effect by the F test ($p \leq 0.05$).

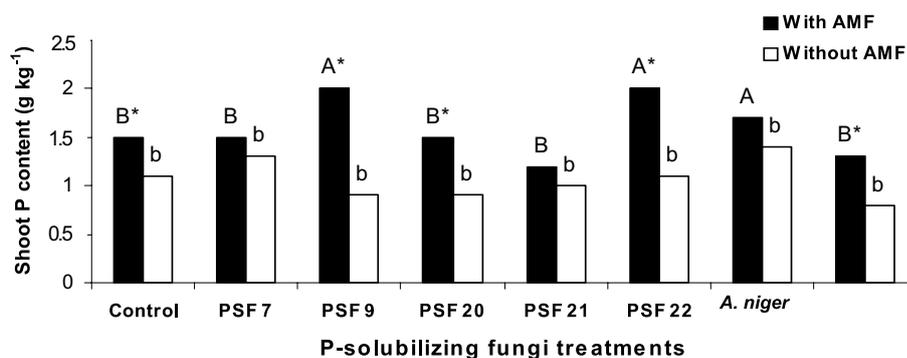


Fig. 2 – Shoot P content of clover inoculated with P-solubilizing fungi from Paraty, Rio de Janeiro, Brazil (PSF 7, 9, 20, 21 and 22) and from Granada, Spain (*A. niger* and *Y. lipolytica*), in the presence or absence of AMF. (Lower-case and upper-case letters compare PSF treatments in the absence and presence of AMF inoculation, respectively, by the Scott-Knott test ($p \leq 0.05$). *: indicates AMF effect by the F test ($p \leq 0.05$).

duction in the absence of AMF. *Aspergillus niger* and PSF 21 with AMF inoculation were the best treatments to increase shoot dry matter. In contrast, plants inoculated with *Y. lipolytica* showed little shoot dry matter production both in the presence and in the absence of AMF (Fig. 1).

No interactions were observed between AMF and PSF when N shoot content was evaluated since AMF inoculation did not affect this parameter. The PSF inoculation yielded the greatest shoot N content (3.2 g N kg^{-1}), which was similar to the control (3.1 g N kg^{-1}).

Similar values of shoot P content were found in plants inoculated or not with AMF in the presence of PSF 7, PSF 21 and *A. niger* (Fig. 2). AMF inoculation increased the shoot P content on the other treatments. No significant differences were found among plants inoculated with PSF, without AMF inoculation, and the control treatment (Fig. 2).

Although PSF 7 and PSF 21 increased the number of nodules in plants, AMF inoculation affected neither this parameter nor the interaction between PSF and AMF (Fig. 3). However, an enhancement on mycor-

rhizal colonization was observed in plants inoculated with AMF in the presence of PSF 7 and PSF 20 (Fig. 4). The other treatments were similar to the control.

DISCUSSION

Since plants were grown in an inert substrate where a nutritive solution with low P concentration was periodically amended, PSF contribution on growth promotion was probably due to an effective Araxá apatite solubilization (Souchie et al. 2005) and/or by phytohormone production (Barea et al. 1997), or even to the increase of rhizobia and AMF symbiosis (Omar 1998, Zaidi et al. 2004). In the case of PSF 21, growth promotion can be related to greater nodulation or release of stimulating substances since this isolate did not increase AMF colonization. Similarly, Cabello et al. (2005) reported that mycorrhizal colonization in *Mentha piperita* was not affected by inoculation with the PSF *Penicillium thomii*, despite this isolate increased other analyzed variables. Zaidi et al. (2004) reported some incompatibility when *Glomus fasciculatum*, *Bradyrhizobium* sp. and *Penicillium* sp. were inoculated in mung bean without any improvement on plant growth. Otherwise, Valdenegro et al. (2001) reported that the inoculation of plant growth promoting rhizobacteria was effective only when *Glomus mosseae* and a native strain of *Rhizobium meliloti* were also inoculated. The success of the inoculation technique is directly related to the functional compatibility among the chosen microorganisms. Therefore, more research is required to discover the best combinations among them in order to increase nutrition and growth of plant species.

Yarrowia lipolytica inoculation yielded similar results to the control treatment for almost all the analyzed variables. However, several authors (Vassileva et al. 2000, Vassilev et al. 2001, Medina et al. 2004) reported considerable improvement on plant growth with *Y. lipolytica* inoculation. Probably, in this study, the main limiting factor was the substrate since no source of organic matter was used. Consequently, the establishment of *Y. lipolytica* was negatively affected.

PSF 7 increased clover shoot dry matter in the absence of AMF similarly to plants inoculated with AMF (Fig. 1). Since this isolate is capable of solubilizing Araxá apatite in liquid medium, and aluminum phosphate in solid and liquid media (Souchie et al. 2005),

it has probably greater solubilization ability in comparison to the other isolates. Moreover, PSF 7 was the one that stimulated the greatest nodulation and mycorrhizal colonization (Figs. 3 and 4). Reyes et al. (2002) also reported that all *Penicillium rugulosum* P-solubilizing strains were able to stimulate maize growth as indicated by 3.6 to 28.6% increases in dry matter yield. Peix et al. (2001) observed a greater number of nodules in common bean plants inoculated with a P-solubilizing *Burkholderia cepacia* strain. Similarly, Fracchia et al. (2004) reported an increased mycorrhizal colonization in soybean after inoculation with the PSF *Aspergillus niger*. Since biological nitrogen fixation depends on the available forms of phosphorus, the combined inoculation of nitrogen fixers, P-solubilizing microorganisms and AMF may benefit plant growth more than any organism group alone.

In this study, although all PSF isolates belonged to the *Aspergillus* genus, it is possible that PSF 7 is a different species, explaining its different effect on clover growth. Also, Nguyen et al. (1992) reported that the P solubilization ability can vary among microorganisms within the same species. Silva Filho et al. (2002) reported that the most efficient P-solubilizing fungi belong to *Aspergillus* and *Penicillium* genera. Future research should be done to determine the organic acids released by these PSF, especially by PSF 7. Moreover, evaluation of the ability to improve plant growth and rhizobia and AMF symbiosis under non-sterilized substrate conditions should be done.

CONCLUSIONS

- 1 – *Aspergillus niger*, PSF 7 and PSF 21 were the best isolates to increase clover growth in the presence of AMF.
- 2 – High mycorrhizal colonization resulted in greater clover growth rate in most PSF treatments.
- 3 – The greatest clover growth rate was recorded when *Aspergillus niger* and PSF 21 were co-inoculated with AMF.
- 4 – The isolate PSF 7 was the best isolate to increase clover growth and to establish rhizobia and AMF symbiosis.

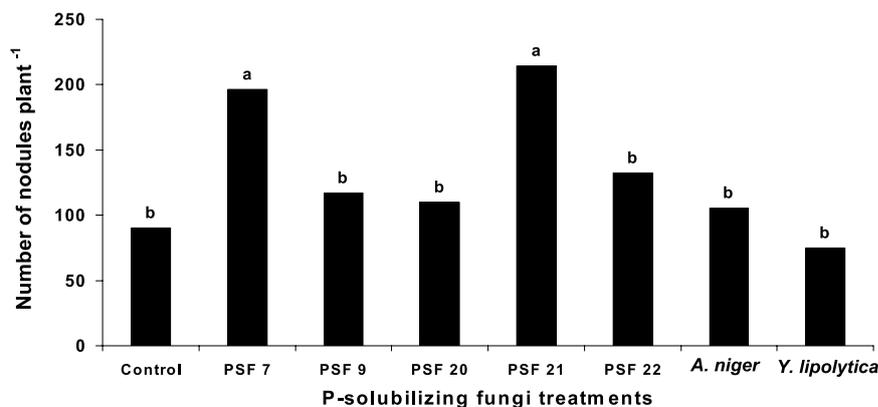


Fig. 3 – Number of *Rhizobium* nodules of clover inoculated with P-solubilizing fungi from Paraty, Rio de Janeiro, Brazil (PSF 7, 9, 20, 21 and 22) and from Granada, Spain (*A. niger* and *Y. lipolytica*). (Lower-case letters compare PSF treatments by the Scott-Knott test ($p \leq 0.05$)).

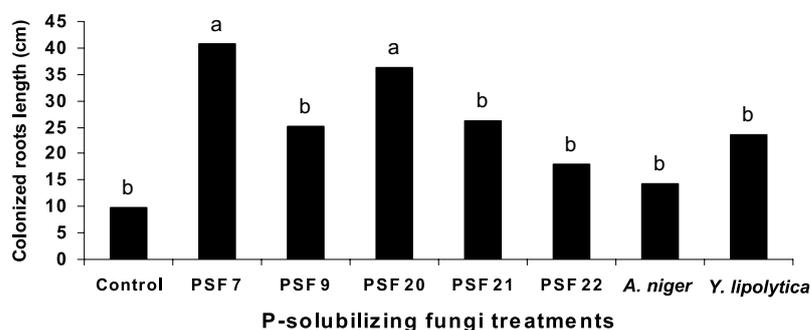


Fig. 4 – Mycorrhizal colonized root length of clover inoculated with P-solubilizing fungi from Paraty, Rio de Janeiro, Brazil (PSF 7, 9, 20, 21 and 22) and from Granada, Spain (*A. niger* and *Y. lipolytica*). (Lower-case letters compare PSF treatments by the Scott-Knott test ($p \leq 0.05$)).

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RESUMO

Este estudo avaliou o sinergismo entre diversos isolados de fungos solubilizadores de fósforo e micorrízicos arbusculares para beneficiar o crescimento de trevo (*Trifolium pratense*) na presença de apatita de Araxá. A cultura foi semeada diretamente em potes plásticos com 300 g de substrato esterilizado formado por areia lavada, vermiculita e sepiolita 1:1:1 (v:v:v) e cultivada em câmara climática. O substrato foi fertilizado com 3 g L⁻¹ de apatita de Araxá. O experimento foi instalado em delineamento completamente casualizado, esquema fatorial 8 × 2 (oito tratamentos de inoculação de fungos solubilizadores de fósforo com ou sem fungos micorrízicos arbus-

culares) e quatro repetições. Os tratamentos de fungos solubilizadores de fósforo consistiram em cinco isolados brasileiros de fungos solubilizadores de fósforo (FSF 7, 9, 20, 21 e 22), dois isolados procedentes da Espanha (*Aspergillus niger* e a levedura *Yarrowia lipolytica*) e o controle (tratamento não inoculado). A maior taxa de crescimento da cultura foi obtida quando *Aspergillus niger* e FSF 21 foram co-inoculados com fungos micorrízicos arbusculares. *Aspergillus niger*, FSF 7 e o FSF 21 foram os isolados mais efetivos para incrementar o crescimento de trevo na presença de fungos micorrízicos arbusculares. A maior taxa de colonização micorrízica resultou em alta taxa de crescimento de trevo na maioria dos tratamentos com fungos solubilizadores de fósforo. O isolado FSF 7 foi o melhor para favorecer o estabelecimento das simbioses com fungos micorrízicos e com rizóbio.

Palavras-chave: apatita de Araxá, solubilização de fósforo, co-inoculação, promoção do crescimento vegetal, rizosfera.

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