DOMINANCE OF _PARIS_-TYPE MORPHOLOGY ON MYCOTHALLUS OF _LUNULARIA CRUCIATA_ COLONISED BY _GLOMUS PROLIFERUM_

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ABSTRACT

Microscopic evidence confirms that _L. cruciata_ hosting _G. proliferum_ shows major anatomical traits (arbuscules, coils, arbusculate coils and vesicles) generally associated arbuscular mycorrhizal roots and the anatomical morphology of intra-thalli mycelium is predominantly of the _Paris_-type. Colonised _L. cruciata_ showed a reduction of biomass when compared with axenic plants suggesting a drain of resources towards the fungus and depletion of nutrients required for optimum plant growth. The behaviour of mycothalli regarding available _KH₂PO₄_ indicates that the nutritional stress threshold for phosphorus (P) is above the residual amount of P already present in Phytagel™ and in plant inoculum. These raise the possibility that in certain circumstances the relationship between _L. cruciata_ and _G. proliferum_ be parasitic rather than symbiotic and open the door for future studies to ascertain the nature of liverwort-AM fungi relationships.

Key-words: Arbuscular mycorrhizal fungi; Phosphorus; Arum-type; Liverwort, Monoxenic cultures

INTRODUCTION

Arbuscular mycorrhizas (AM) are ubiquitous underground symbiotic associations between plants and obligate fungi of the phylum Glomeromycota (21). From this symbiosis plants generally obtain higher yields as they improve their capacity to acquire low mobile soil nutrients and increase resistance to biotic and abiotic stresses. Concomitantly the fungi are able to access the host photosynthate carbon pools and so to complete their life cycle (1,15,23). Most available information on the physiology and anatomy of mycorrhizae is related to the sporophyte of Tracheophyta. Conversely for non-vascular plants the knowledge is still scarce. Within these the liverworts are an important and extremely successful group found in all continents and environments. Because liverworts are thought to be amongst the original colonisers of terrestrial habitats and appear to have remained relatively unchanged through time they probably hold the key to early terrestrial diversification of land plants (17,18). Some complex thalloid liverworts (Marchantiales) are known to form mycorrhiza-like associations with AM fungi. Mycothallus (3) develop structures that are analogous to those observed in AM roots, thus indicating possible functional similarities (8,10,11,13,17,19). In these plants, as in roots, the AM fungus grows and connects two distinct environments: within the cells of the host body as inter or intra-cellular mycelium; and externally in the soil or medium matrix, thus extending the host plant capacity to access well beyond their body limits into the soil matrix. In roots the internal mycelium has been shown to have one or a combination of two different morphological types (2,9,22): Arum and Paris. The Arum-type morphology, first described on _Arum maculatum_, shows intercellular hyphae mainly growing longitudinally between cells with arbuscules rising on short upright intra-cellar branches. The Paris-type, originally described on _Paris quadrifolia_, is defined by cell-to-cell intracellular growth with the formation of coils and arbusculate coils. Depending upon the host plant and fungus these morph types can occur isolate or simultaneously within the same plant forming a continuous mycorrhizal structure.

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(5). In liverworts the morphology of hyphae within mycothallus is not yet fully characterised, however most reports indicate that AM fungi generally progresses within the plant body with patterns resembling the Paris-type (6,11).

Phosphorus (P) is an important macronutrient involved in key structural and metabolic functions of all organisms. Moreover, it is widely accepted that in many plants AM fungi plays the main role in the resistance to biotic and abiotic stresses by improving the host capacity to uptake major nutrients from soil, ex., inorganic phosphate (23). Although this is true for Tracheophyta the available information regarding nonvascular plants and particular those cultured in vitro is scarce or nonexistent (7). The nutrient requirements of liverworts vary considerably from those of most vascular plants.

The present work aims to address (i) the physiological effect of phosphorus on biomass production of L. cruciata colonisation by AM fungi, (ii) To survey the morph types of G. proliferum mycelium within L. cruciata thallus.

MATERIALS AND METHODS

Biological material and growth conditions

*Glomus proliferum* Dalpé & Declerck (MUCL 41827), acquired from GINCO (Mycotheque de l’Université Catholique de Louvain, Laboratoire de Mycologie, Belgique) was multiplied and maintained in monoxenic cultures of *Lunularia cruciata* (L.) Dumortier ex. Lindberg. Plants and fungi were kept throughout the experiments at 25°C with a 10/14 hours light/dark photoperiod in a Sanyo MLR–350H chamber with light of an average intensity of 68.3 ± 6.4 μmol s⁻¹ m⁻² as described by Fonseca et al. (8).

Inocula preparation

Inocula were obtained from axenic and monoxenic thallus of *L. cruciata* cultured for 100 days on SRV (8) with 29.2 mM sucrose, monoxenic cultures used showed profusion production of external hyphae and spores of *G. proliferum* (Fig. 1).

Light microscopy

To study the pattern of colonisation of *L. cruciata* by *G. proliferum* 0.5 cm thallus segments were cultured for 49 days on SRV medium as described by Fonseca et al. (8). The growth length of 44 mycothallus apices were measured weekly along an imaginary line through the thallus midrib (Fig. 2a). Measured segments were then cropped, fixed in Bouin’s fluid and cleared with 10% KOH, at 80°C for 20 min. Samples were washed in distilled water, acidified in 1 N HCl before being dehydrated and embedded in paraffin wax. Sections of about 10 μm were cut with a microtome (Leitz model 1512), mounted on microscope slides and stained overnight in 0.05% trypan blue (16). Images were digitally acquired with a Carl Zeiss Axiocam HR apparatus.

Phosphorus experiment

Discs of thallus (10.7 ± 1.7 mm² made by Ø3.27 mm cork-borers) from axenic and monoxenic *L. cruciata* were cultured for 70 days on 30 ml of SRV (8) with 29.2 mM sucrose and three levels of added phosphorus. A 2×4 factorial design was setup with the fungal treatment consisted of the presence *G. proliferum* and absence of fungus. The phosphorus treatment consisted of SRV media with three levels of added KH₂PO₄ (123.0, 61.5 and 30.7 μg KH₂PO₄) and SRV medium without added KH₂PO₄. There were ten replicate Petri dishes per treatment.

Data collection

Thallus length was estimated by engraving the contour of each thallus apex, under stereoscope microscope, on the lower side of the plastic Petri dish. At the end of the experiment the lines drawn along the thallus midrib were measured. Plant biomass was estimated as dry weight after oven-drying (60°C) to a constant weight. Number of spores and hyphal length were measured under a stereomicroscope with a 6×6 square hairline.
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graticule of 20.25 mm² regularly placed at 0.5 cm intervals over
the surface of inverted Petri dishes and following the method
by McGonigle et al. (12). Data were evaluated for significance
(P<0.05) using multivariate analysis of variance (ANOVA/
MANOVA) and post hoc Least Significant Differences test with
Statistica software for Windows v4.5 (Statsoft Inc.).

RESULTS

Effect of phosphorus

The quantity of phosphorus present in Strullu-Romand (4) and SRV media is suitable for both axenic and monoxenic Ri T-
DNA transformed roots cultures colonised by AM fungi and
for mycothallic L. cruciata. However, when added P is reduced
by 50 or 75% from SRV levels no significant responses were
observed on mycothallic L. cruciata dry weight and AM fungus
hyphae and spore production (Table 1). Only when any KH₂PO₄
wacS added to the medium did plants show a decrease in biomass
and the fungus presented a reduction in spore production,
compared to the treatment with 123 μg KH₂PO₄ (Table 1), but
not for external hyphae length. Alternatively, if one analyses
the data in terms of presence/absence of AM fungus
colonisation the results shows that liverwort growth was
negatively affected (P<0.05) in monoxenic conditions. Colonised
plants showed a reduction of 54.7% in dry weight, compared
with axenic L. cruciata (Table 1).

Thallus growth rate

When cultured in 30 ml of SRV (with 123 μg of added
KH₂PO₄) mycothallus of L. cruciata showed during the first
35 days a steady growth rate of 0.382 mm·day⁻¹ (r² = 0.999).
From then on (the last 14 days of culture) the growth average
rate decreased till about 13.1% at the end of the experiment
(Fig. 2b) giving an overall average growth rate of 0.358 ± 0.025
mm·day⁻¹ for 49 days culture.

Anatomical characterisation of symbiosis

L. cruciata is a complex thalloid liverwort with an internal
differentiated anatomy (Fig. 2c). Intercellular hyphae were
conspicuous throughout the mycothalli with few exceptions:
hyphae were scarcely present among the small chlorophyllous
cells within the photosynthetic layer; no fungus was observed
in the meristematic zones on thallus apices; and, as reported by
Fonseca et al. (8), no hyphae colonised rhizoid cells. Conversely,
G. proliferum was profusely present within the highly vacuolated
parenchyma cells with special prevalence in the central midrib
area where trypan blue stain revealed a distinct fungal layer
(Fig. 2c). Within this layer a denser network of internal hyphae
could be seen connecting arbuscules, coils and arbusculate
coils, as well as, scattered vesicles. Hyphae and arbuscules
were also present in oil cells (Fig. 2d). The morphological type
of L. cruciata colonisation by G. proliferum was predominantly

![Figure 2. Lunularia cruciata colonised by Glomus proliferum](image-url)

grown for 49 days on SRV medium with 29.2 mM of sucrose. (a) Mycothallus apex showing midrib imagine line arrow used
for the last measurement of thallus length. (open arrows) Spore
clusters imbedded in the medium. (b) Mycothallic averages
and standard deviations (vertical bars) of growth length of 44
apices measured at 7 days intervals. Fitted line and equation
describes the lengthening pattern of mycothallus during the
first 35 days of growth. Equation: y, Length (mm); x, Time
(days); r², R-squared value. (c to f) Light microscopy of trypan
blue-stained samples: (c) Anatomic section of mycothallus
showing (1) photosynthetic layer under an upper epidermis;
(2, fungal layer) thallus’ midrib parenchyma with high
concentration of arbuscules, vesicles and oil cells; (3) the
lower epidermis with (4) rhizoids and (5) scales; (d) Arbuscules
within (6) oil cells located in the thallus’ midrib parenchyma.
These cells show their oily content only partially removed by
the method for microscopy. (e) Small section of mycothallus
midrib anatomy exemplifies a common morphological pattern
showing (7) hyphae crossing cell-to-cell in a pattern
characteristic of the Paris-type. (8) Arbusculate coils. (f) The
less frequent morphology of Arum-type was also present in
some cluster of cells within the thallus midrib showing several
overlapping arbuscules connected to (9) hyphae
progressing close to liverwort cell wall. Bars: (a) 5 mm; (c) 200
μm; (d) 50 μm; (e, f) 20 μm.
Table 1. Biomass, number of fungal spores and hyphae length production of *Lunularia cruciata*, cultured for 70 days in 30 ml of SRV medium with 29.2 mM sucrose, with and without *Glomus proliferum* and with different levels of added KH2PO4 plus without added phosphorus. Yes – plants colonised by *G. proliferum*; No – axenic *L. cruciata*.

| G. proliferum | KH2PO4 (μg) | Dry weight (g) | Number of spores | Hyphae length (mm) |
|---------------|-------------|----------------|------------------|-------------------|
| Yes           | 0.0         | 0.066±0.017 a  | 11038±9232 a     | 69805±41445 a     |
| Yes           | 30.8        | 0.082±0.010 ab | 12766±6966 a     | 75937±38726 a     |
| Yes           | 61.5        | 0.074±0.032 ab | 25940±15216 ab   | 72975±48888 a     |
| Yes           | 123.0       | 0.091±0.023 b  | 54749±34710 b    | 135278±73912 a    |
| No            | 0.0         | 0.131±0.030 c  | —                | —                 |
| No            | 30.8        | 0.159±0.025 c  | —                | —                 |
| No            | 61.5        | 0.129±0.016 c  | —                | —                 |
| No            | 123.0       | 0.153±0.027 c  | —                | —                 |

Means (± Standard deviation) followed by the same letter do not differ significantly (*P*<0.05).

of the *Paris*-type (9,22) (Fig. 2e), however in places the hyphae and arbuscules denoted a pattern closest related to the *Arum*-type morphology (Fig. 2f). Here the hyphae appear to grow close to the plant cell wall with arbuscules rising on short upright intra-cellular branches.

**DISCUSSION**

The cultures used as inoculum for the experiments had their origin in 2003 (8). Since then they have been regularly subcultured producing a reliable means to maintain and multiply *G. proliferum*. In addition, the use of *L. cruciata* as host for AM fungi has the advantage, over the Ri T-DNA transformed root systems as they allow easy crop and manipulation of the external mycelium. With these cultures the plants usually occupy less than half the Petri dish area and sometimes leaving undisturbed mycelium. With these cultures the plants usually occupy less than half the Petri dish area and sometimes leaving undisturbed mycelium. With these cultures the plants usually occupy less than half the Petri dish area and sometimes leaving undisturbed mycelium.

In vitro cultured *L. cruciata* with and without *G. proliferum* behaved indifferently to changes in medium added P. Both plant dry weight and AM fungi growth (number of spores and external hyphae length) could not resolve significant changes with these used levels of KH2PO4 that is in accordance with the capacity of liverworts to exhibit normal development on wide range of media (7). A different plant and fungus behaviour was observed when any KH2PO4 was added to the SRV medium. For these plants there were trace amounts of P derived from residual P brought by the initial plant inoculum (thallus discs) and by medium Phytagel™ component (14). Because differences in plant and fungal growth were observed between treatments where any and the maximal KH2PO4 was used, we may speculate that the limiting threshold for P to induce differences in plant and AM fungal behaviour lay above residual P present in the experiment. Moreover, in the present culture conditions, the plant overall growth pattern strongly suggests that the colonisation of *L. cruciata* by AM fungi is a heavy burden for the plant growth thus implying that the colonisation of *L. cruciata* by *G. proliferum* is not a mutualistic symbiosis but rather a parasitic one. However, the anatomical traits observed in this association are consistent with those present in mycorrhizae colonised by *G. proliferum*. The liverwort may gain from the association only if important nutrients, such as phosphorus, are directly unavailable to the plant or if available they are below the optimum threshold for maximal mycothallus growth.

Considerations about the morphotype of *G. proliferum* growth within *L. cruciata* thallus were already raised by Fonseca *et al.* (8). At the time it was suggested that the architecture *G. proliferum* was more consistent with the *Paris*-type. Our extensive survey of mycothallus of *L. cruciata* agrees with this assertion. The fungus colonises almost all parts of the thallus except areas close to the meristematic zones. In this survey any information was gathered on the presence of AM fungi in reproductive structures (sexual and asexual) because the photoperiod imposed to the cultures was conducive to the absence of those structures. Our study also confirms findings reported by Fonseca *et al.* (8) that *G. proliferum* hyphae and arbuscules were present in oil cells and absent from rhizoids. Although the converse was described for other liverworts (11,19) the persistent absence of rhizoids from in vitro studies of liverworts and hornworts (20) may indicate that the diffuse presence of light *in vitro* within the medium and all over the plant excludes the rhizoids from being the principal source of plant colonisation. Within the thallus midrib, the trypan blue staining revealed a layer of intense marked cells where colonisation by *G. proliferum* formed a zone rich in arbuscules, coils, arbusculate coils and other AM anatomical traits. The predominant morphotypes are consistent with the *Paris*-type as the hyphae were observed to progress from cell to cell and, particularly within the midrib fungal layer, numerous arbuscules.
and coils were present. However scattered within the AM fungal zone of the parenchymatous layer one occasionally observed clusters of cells where the fungi appear to progress in a mode more consistent with the *Arum*-type.

**CONCLUSIONS**

From the observations in this study and from those of an earlier study on mycothallus of *L. cruciata* (8) we propose that the colonisation by *G. proliferum* not only has the major mycorrhizal traits generally associated with the colonisation of roots by AM fungi (arbuscules, coils, arbuscule coils and vesicles), but it also shows that the internal hyphae colonises the thallus predominantly with a *Paris* morphotype. Furthermore, the colonisation of *L. cruciata* by *G. proliferum* resulted in a reduction of host biomass compared with axenic plants suggesting a bypass of resources towards the fungus. Hence the relationship between *L. cruciata* and *G. proliferum* may not always be, if ever, symbiotic in nature. Finally, the present study shows that, despite any response from *L. cruciata* to changes in medium above 30.7 µg KH₂PO₄, the P stress observed in mycothallus when any P was added indicates that the threshold for optimum growth is above the residual amounts of P brought by contaminants in Phytagel™ and P in the plant inoculum. This finding opens the door for future nutritional studies to ascertain the nature of liverwort-AM fungi relationships.

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