

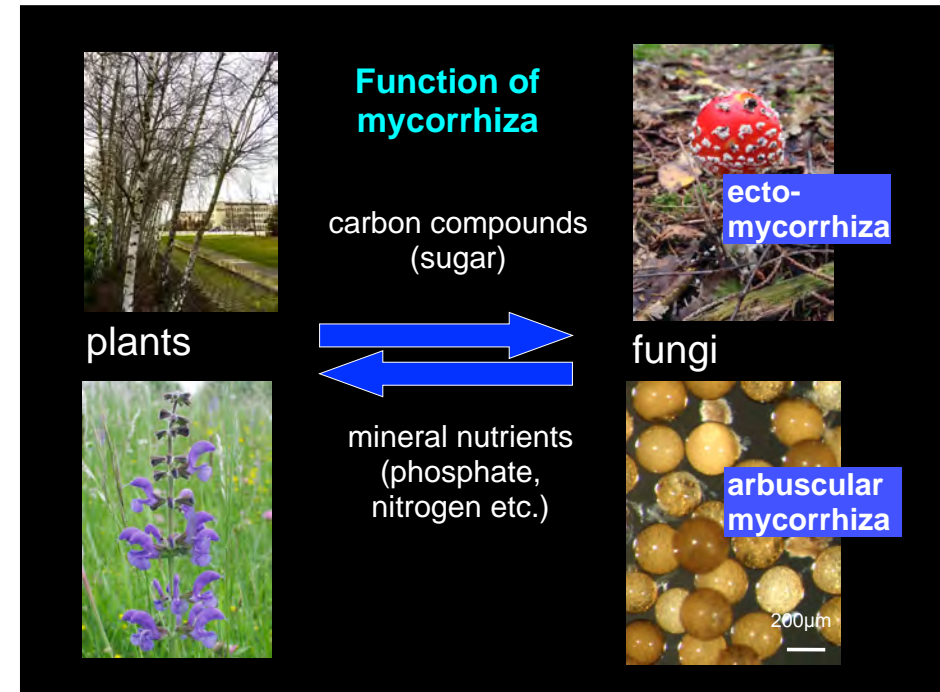
Arbuscular mycorrhiza and agroecology

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International Bank for the Glomeromycota

UMR 1347 Agroécologie (INRAE, UBFC, AgroSup, CNRS)



Symbiotic partners of arbuscular mycorrhiza

plants

-about 70-80% of vascular plants: mainly herbs, tropical trees

-hornworts, liverworts, horsetails, clubworts

(> 150 000 species)

fungi

phylum

Glomeromycota

(>300 described species)

(+Mucoromycota: ?? species)

The ubiquitous symbiosis for house, garden and agriculture

Fungi: Glomeromycota

Plant hosts:

leek, onion, tomatoes, potatoes, beans, peas, carrots, parsley, cereals, lettuce, vines, apple trees, grasses etc.

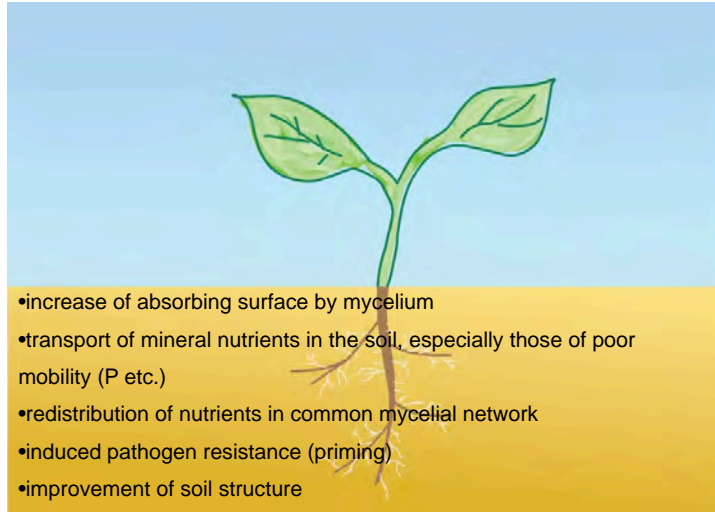


(Photo: V. Gianinazzi-Pearson)

Non-hosts:

cabbage, spinach, rutabaga

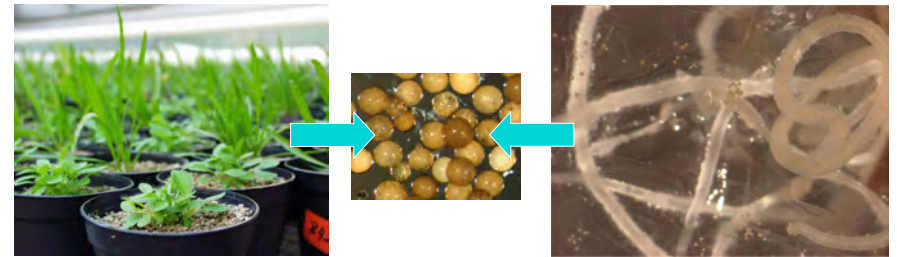
Common Mycorrhiza Functions:



- increase of absorbing surface by mycelium
- transport of mineral nutrients in the soil, especially those of poor mobility (P etc.)
- redistribution of nutrients in common mycelial network
- induced pathogen resistance (priming)
- improvement of soil structure

Culturing Glomeromycota

- obligate biotrophs
- spores contain numerous other microorganisms
- except few species cultivated on transformed roots (“in vitro culture”)
- host specificity in cultivated isolates very low, host preferences in seminatural systems



The International Bank for the Glomeromycota



Installations

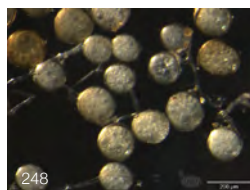


main collection:
3 climatized rooms with
10 custom-built growth cabinets



The isolates

- 43 isolates in 30 species (312 pot cultures) on their plant hosts
- 6 of these also available "in vitro" on transformed carrot roots
- origin:
 - Europe: 24 (France: 3)
 - N America: 9
 - S America: 3
 - Australia/New Zealand: 3
 - Asia: 4
- type strains (*ex-type*): 11



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Approaches for mycorrhizal biotechnology

- management: maintaining and supporting existing indigenous species
 - establishing conditions conducive to AMF diversity and abundance: e.g. no-till, low pesticides, low fertilisation, avoiding non-host plants in rotations
- inoculation: adding fungal propagules
 - competition problem, can be challenging due to inoculum quantity; most inoculation studies do not verify inoculation success and many use sterile soil as control
 - well-established for plants in sterile potting substrates and then outplanted
- (plant breeding: plant cultivars differ in their mycorrhizal response)



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Effects of agriculture on AMF diversity: Land use intensity and AM fungal diversity in soils (spore abundance) Oehl et al. AEM 2003

Land use	Fertilisation	Total number AM fungal species	Species diversity index (Shannon-Weaver)
Permanent grassland	No fertilisation (20 yrs)	24	2.29ab
		25	2.32ab
		20	2.34ab
Arable			
7 yr crop rotation (grass-clover)	Bio-organic Conventional	18	2.45a
		13	2.14b
Monocropping (maize)	Mineral & organic	10	1.70c
	Mineral	9	1.72c
	Mineral	8	1.32c

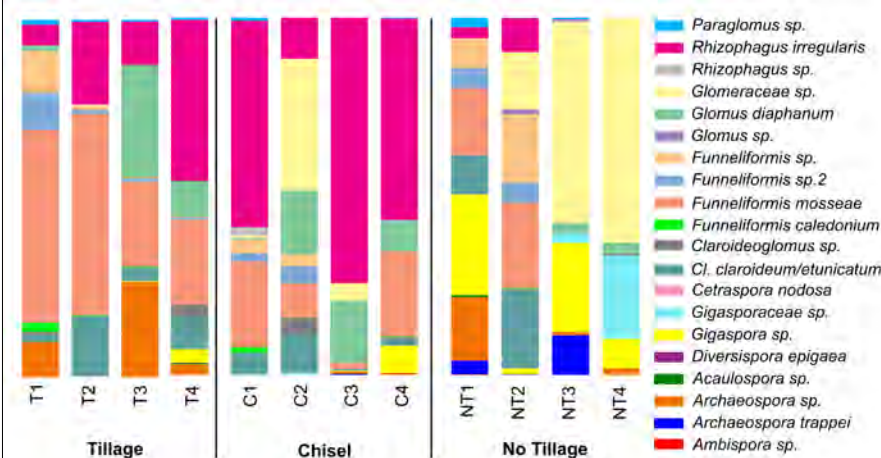
AM fungal diversity: disturbed (arable) < non-disturbed (grassland) ecosystems



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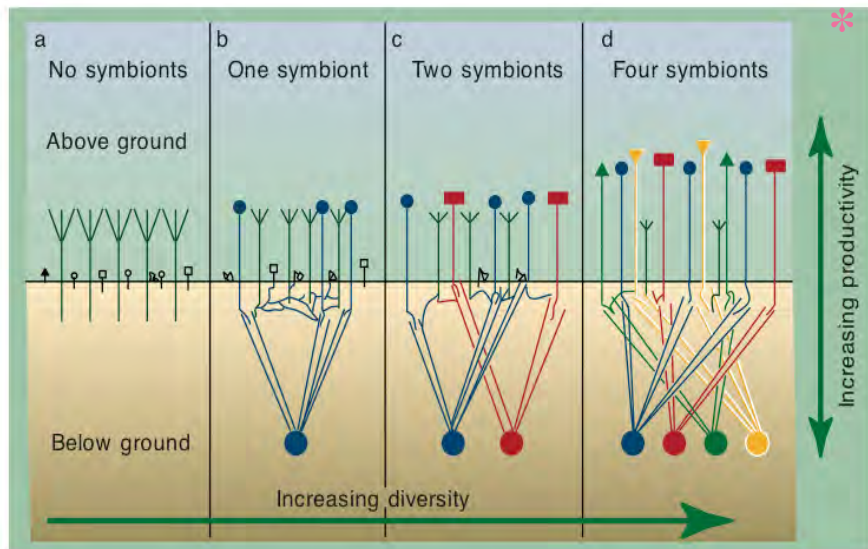
Community structure of AMF in a field experiment in Tänikon (CH) based on 454 Titanium FLX+ sequencing of an rpb1 region (Stockinger et al., 2014)



16 molecular taxa

15 molecular taxa

19 molecular taxa



Aboveground and belowground biodiversity are correlated
 (M. van der Heijden et al. / D. Read, Nature 396, 1998)

AMF effects on plant communities

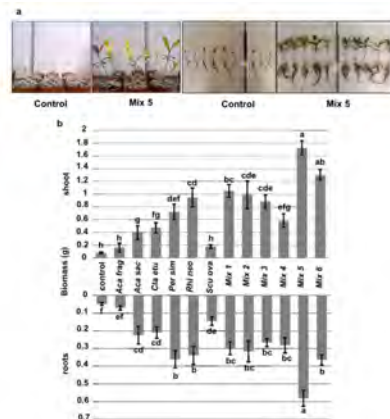


Encouraging diversity. Grasses prevail in poor soil conditions (right) but are overrun by a broader plant community when mycorrhizal fungi are added to the soil (left).

Photo: Marcel van der Heijden

Is a mixture of arbuscular mycorrhizal fungi better for plant growth than single-species inoculants? Crossay et al., 2019

Fig. 1 *Metrorhizon lanifolia* (a) control versus treatment Mix 5 (representing the extremes) after 8 months of growth on siliceous substrate. b Dry weight of shoot and root, respectively. Bars represent means, and error bars represent standard error of means ($n = 10$ except for control treatment $n = 9$). Means followed by the same letter above columns do not differ significantly by the Mann-Whitney post hoc test ($p \geq 0.05$).



A large part of AMF functional diversity may be found on the intraspecific level

- Need to take into account inter-isolate variation
- Strong inter-isolate variation of AMF-induced yield differences in crop plants with *F. mosseae* (Munkvold et al., 2004)
- Possible intra-species ecotypes grassland vs. elsewhere of *R. irregularis* (Börstler et al., 2010)

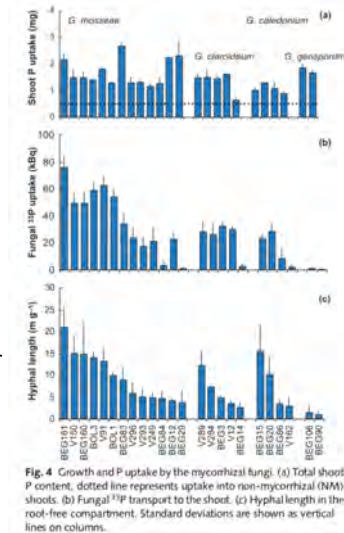
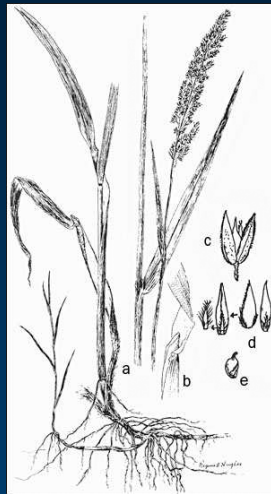


Fig. 4 Growth and P uptake by the mycorrhizal fungi. (a) Total shoot P content, dotted line represents uptake into non-mycorrhizal (NM) shoots. (b) Fungal ³²P transport to the shoot. (c) Hyphal length in the root-free compartment. Standard deviations are shown as vertical lines on columns.



Phalaris arundinacea

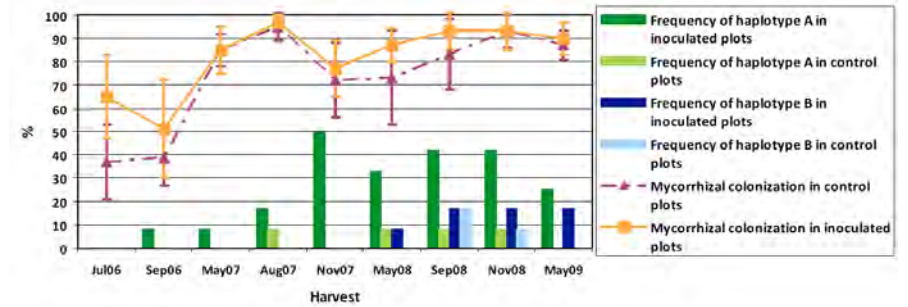
AMF are already there!
Using mtLSU markers to trace an inoculated strain in the field:

Sýkrova et al., 2011



Re-vegetated mine spoil at Chomutov, Czech Republic

Frequency of haplotypes A and B and mycorrhizal colonization in the roots of *P. arundinacea*



Using *Rh. irregularis* mtLSU haplotypes to demonstrate inoculation success:

Haplotype A was significantly more frequent in the inoculated than in uninoculated plots

(Analysis of variance Df error=11; F=24.43; p<0.001)

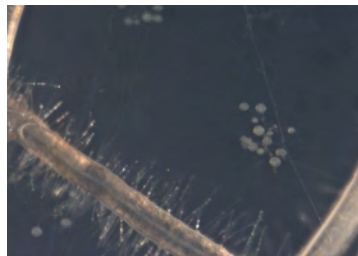


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The “*in vitro*” dilemma

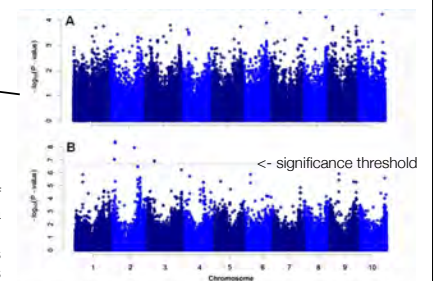
- “*in vivo*” inoculum production often has problems with purity or propagule density.
- Root organ culture offers the only way to produce axenic AMF biomass
- Higher technological effort necessary for large-scale production
- Only very few AMF species are available using this method (basically *Rhizophagus* isolates)
- Most *in vitro* inoculum produced worldwide is from a single isolate: *Rh. irregularis* DAOM 197198



Plant breeding, a possible door opener for mycorrhizal technology?

- Plant cultivars are known to differ in mycorrhizal growth response (MGR) and root colonization (Graham et al., 1991; Hetrick et al., 1993)
- Does domestication negatively affect MGR? Breeding for improved root soil exploration for P could result in selection for a lower MGR and such plants may limit colonisation (Graham et al., 1991)
- Breeding for high colonization may be difficult due to low heritability (Leiser et al., 2016)
- Is it more important to select for P use efficiency?

Genome-wide association study of
A: mycorrhiza root colonization,
B: single-tiller biomass of *Sorghum* cultivars



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Importance of mycorrhiza collections for sustainable agriculture

- ressource centers for research and inoculum production:
 - culturing methods, detection/identification, functional diversity, interactions with plant cultivars...
 - well-defined inoculum isolates
- represent intraspecific diversity of of “common colonizers” (*Rh. irregularis*, *C. claroideum*, *F. mosseae* etc.)
- represent maximum taxonomic (and functional?) diversity outside of “common colonizers”



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Thank you very much for your valuable
attention!



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