1st International Webinar

Parts In Plants And

Biodiversity in Plant Protection in Arid Region ID:BD0025

Onine October 18-19th, 2023



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IMPACT OF NATIVE ARBUSCULAR MYCORRHIZAL FUNGI ON *Tamarix articulata* GROWTH AND RHIZOSPHERIC MICROBIOTA BIODIVERSITY

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Introduction

• Soil degradation is an increasingly important problem in many parts of the world, particularly

in Mediterranean arid and semiarid areas.



Soil desertification in Algerian steppe



Revegetation programmes in degraded Algerian areas



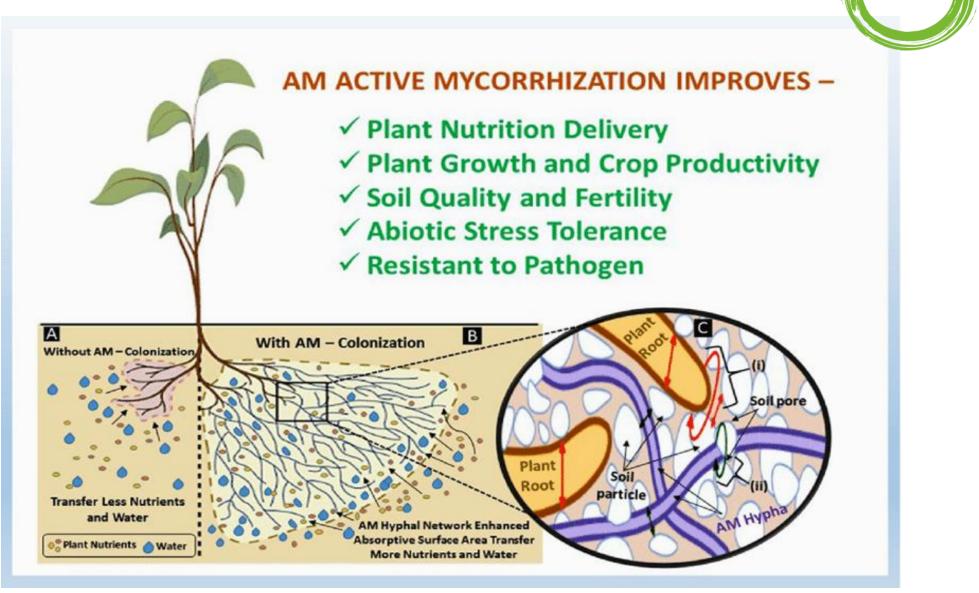
Failure of revegetation programmes in Algerian areas



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Arbuscular mycorrhizal fungi as biofertilizers



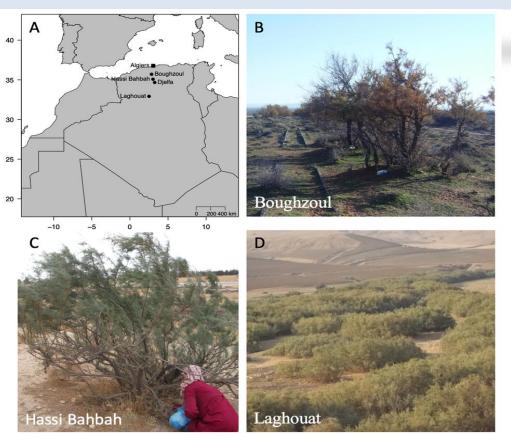




In the other hand, *Tamarix articulata* is one of the proposed halophytic shrubs in Algerian revegetation programs. This species has ecological and medicinal values, such as erosion control, wind break, and application in apiculture and cosmology.

Nevertheless, little is known about the importance of AMF inoculum sources on *Tamarix articulata* development in natural saline soils. Therefore, the current study aims at investigating the efficiency of two AMF-mixed inoculums on *T. articulata* growth, with consideration of its rhizosphere microbiota.

Material and Methods



A) Map showing the distribution of the sampling sites in Northern Algeria. Graduations on the Y and X axes are latitudes and longitudes. B to D) Pictures showing the type of environment for the sites sampled at Boughzoul, Hassi Bahbah and Laghouat.

Indigenous inoculum made of strains . originating from Algerian saline soils: Boughzoul region) (from and а commercial one were used to inoculate T. articulata in four saline soils with different salinity levels under microcosm conditions with evaluation of rhizosphere microbial biomasses. The studied soils were sampled from: Laghouat, Djelfa, Boughzoul and Hassi bahbah, they have different salinity level ranging between:

1.1 and 4.5 dS.m⁻¹.







Total mycorrhizal rate and plant biomasses were evaluated after six months of cultures.



The impact of soil salinity on AMF establishment and on rhizospheric biomass microbiota were evaluated using specific lipid biomarkers

Data were analyzed statistically by means of comparison by multivariate analysis with Excel stat 2020.

Results and discussion

AMF T. articulata Colonization

1.

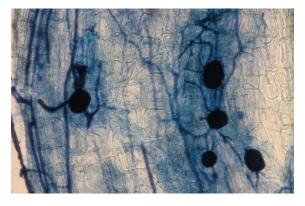
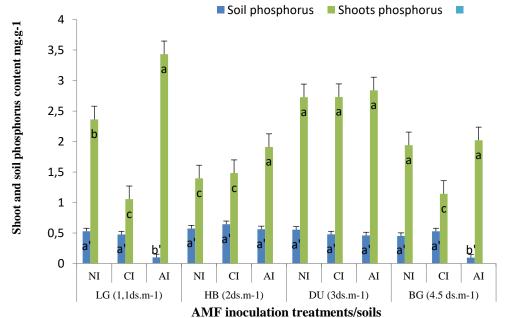


Table 1. Effect of AMF treatments on total AMF colonization of T. articulata cultivated in the four studied soils.

TreatmentNICINICININICININICINIMycorrhizal rate15.415.121.914.710.926.712.211.829.19.415.433.2Plant biomasses1.260.921.491.211.151.871.160.941.180.981.141.36		LG (1.1 dS.m-1)			HB (2.1 dS.m-1)			DU (3.1 dS.m-1)			BG (4.5 dS.m-1)		
rate Image: Constraint of the second sec	Treatment	NI	CI	NI	NI	CI	NI	NI	CI	NI	NI	CI	NI
	·	15.4	15.1	21.9	14.7	10.9 (26.7	12.2	11.8	29.1	9.4	15.4	33.2
		1.26	0.92	1.49	1.21	1.15 (1.87	1.16	0.94	1.18	0.98	1.14	1.36

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2. Phosphorus Contents in Soil and Shoots



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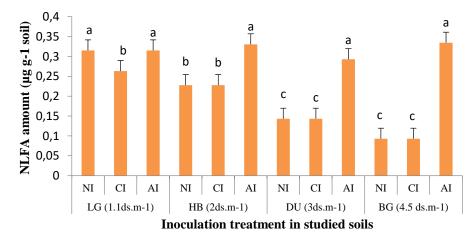
Plant biomasses are correlated with increasing shoot phosphorus content. In addition, Analysis of variance indicated that shoot phosphorus content was affected positively by indigenous AMF inoculation.

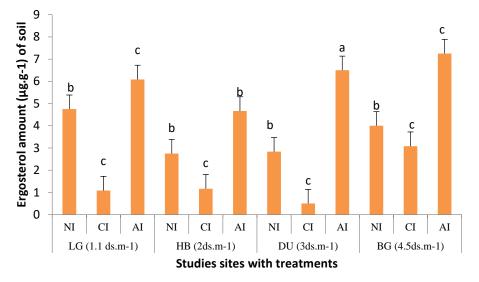
Impact of AMF inoculum sources on *T. articulata* shoots phosphorus content in the four studied soils. NI: non-inoculated. CI: commercial inoculum. AI: Indigenous inoculum.

2. Soil microbial biomasses:

Increase by 46% for AMF

Impact of AMF inoculum sources on C16:1w5 soil content of *T. articulata* in the four studied soils. NI: non-inoculated. CI: commercial inoculum. AI: Indigenous inoculum.



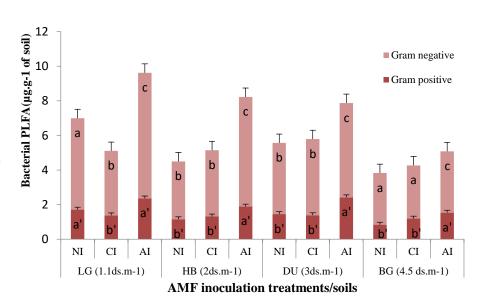


Impact of AMF inoculum sources on ergosterolsoil content of *T. articulata* in the four studied soils. NI: non-inoculated. CI: commercial inoculum. AI: Indigenous inoculum

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25% for saprotrophic fungi evaluated by ergosterol amount.

and 15% for bacterial biomasses.



Impact of AMF inoculum sources on PLFA C16:1w5 content of *T. articulata* in the four studied soils. NI: non-inoculated. CI: commercial inoculum. AI: Indigenous inoculum

Pearson's correlation coefficient between mycorrhizal colonization rate, shoot dry weight and the different measured parameters under AMF inoculation treatments:

	NI Trea	atment	CI Trea	atment	AI Treatment		
	MCR	SDW	MCR	SDW	MCR	SDW	
SDW	0.5*	-	0.7*	-	0.9**	-	
RDW	0.6*	-	-0.2	0.9***	0.8**	0.9***	
Shoot Phosphorus	0.6*	0.6*	0.7*	0.5*	0.9**	0.7*	
Soil phosphorus	-0.3	0.9***	0.3	0.7**	-0.8**	0.2	
Shannon index	0.3	0.1	0.3	0.2	0.9**	0.4**	
AMF species richness	0.5	0.3	-0.4	-0.5	0.9**	0.2**	
NLFA C16 :1ω5	0.6*	0.4 *	-0.1	-0.5	0.7*	0.5*	
PLFA C16 :1ω5	-0.00	0.6	-0.00	0.5	0.6*	0.2*	
NLFA/PLFA C16:1ω5	0.6*	0.3 **	-0.1	-0.3	0.6*	0.00*	
Ergosterol	0.4	0.6	0.0	0.3	0.6*	0.2*	
Gram positive bacteria	0.5	0.0	0.2	0.1	0.5*	0.2*	
Gram negative bacteria	0.4	0.3	0.00	0.1	0.6*	0.2**	

MCR: Mycorrhizal colonization rate. SDW: Shoot dry weight. RDW: root dry weight.*Correlation is significant at the 0.05 level. ** Correlation is significant at the 0.001 level. NI: non-inoculated. CI: commercial inoculum. AI: indigenous inoculum

Conclusion

• Present results open the way towards the preferential use of mycorrhizal inoculum, based on native AMF, to perform revegetation and to restore the saline soil microbiota biodiversity.





Thanks for your attention

