

Can Arbuscular Mycorrhizal Fungi Inoculum Improve Conditions of Various Golf Greens?

Florence Sessoms, Ryan Schwab, Eric Watkins

Department of Horticultural Science, University of Minnesota, St. Paul, MN



Objective: Determine if repeated inoculations of arbuscular mycorrhizal fungi would result in improved turfgrass quality, NDVI, and soil moisture of four different types of golf greens.

Materials & Methods

Experimental design and treatments

The experimental design was a split-plot design with 3 replications. The main plot was nitrogen treatment applied every 10 days, and the sub-plot was inoculum treatment applied at least once per month. The arbuscular mycorrhizal fungi (AMF) inoculum used was MycoApply® Injector Endo (Figure 1; Mycorrhizal Applications, Grants Pass, OR).

Main plot: Nitrogen treatment

- Low (0.15 g N m⁻²)
- Regular (0.49 g N m⁻²)

Sub-plot: Inoculum treatment

- AMF-inoculum (initial application rate of 2.6 mg inoculum m⁻², subsequent applications rate of 1.3 mg inoculum m⁻²)
- Water-treated

Figure 1. Bottles of AMF inoculum mixture. The inoculum contains propagules of *Glomus intraradices*, *Glomus mosseae*, *Glomus aggregatum*, and *Glomus etunicatum*.



Experiments

The study was implemented as four separate experiments, each on a different type of research green (Table 1). Each experiment was performed once throughout the 2020 growing season.

Table 1. Turfgrass species and rootzone type of each experiment

Experiment no.	Turfgrass species	Rootzone
1	Creeping bentgrass (<i>Agrostis stolonifera</i> L.)†	USGA sand-based
2	Creeping bentgrass	Push-up (silty loam)
3	Fine fescue mixture (<i>Festuca</i> spp.)‡	USGA sand-based
4	Fine fescue mixture	Push-up (silty loam)

† 'L-93' creeping bentgrass

‡ 'Radar' and 'Compass' Chewings fescue (*Festuca rubra* spp. *commutata* Gaudin), and 'Seabreeze GT' and 'Joppa' slender creeping red fescue [*Festuca rubra* spp. *litoralis* (G.F.W. Meyer) Auquier]

Data collection

The following data was collected 7 and 15 days after each inoculation:

- Turfgrass quality (1-9 scale; TQ)
- Normalized difference vegetation index (NDVI)
- Volumetric water content (VWC)

Statistical analysis

Each experiment was analyzed using a linear model representing a split-split-plot. The main plot was date, the sub-plot was nitrogen treatment, and the sub-sub-plot was inoculum treatment. Mean separation for any significant interactions was performed using estimated marginal means with Tukey's method at $\alpha=0.05$.

Results

Experiment 1

Table 2. ANOVA results of Experiment 1

Source of variation	df	TQ	NDVI	VWC
Rep	2	ns	ns	ns
Inoculum	1	ns	0.055	ns
Nitrogen (Ntrt)	1	ns	*	ns
Date	8	***	***	***
Ntrt:Inoculum	1	ns	ns	ns
Date:Ntrt	8	***	***	*
Date:Inoculum	8	ns	ns	ns
Date:Ntrt:Inoculum	8	ns	ns	ns

* Significant at 0.05 probability level; ** significant at 0.01 level; *** significant at 0.001 level; ns, nonsignificant at 0.05 level

In this experiment, there was a moderately significant (P -value = 0.055) effect of inoculum treatment on NDVI (Table 2). Averaged across date, the AMF-inoculated and water-treated plots had NDVI ratings of 0.782 and 0.778, respectively.

Experiment 2

Table 3. ANOVA results of Experiment 2

Source of variation	df	TQ	NDVI	VWC
Rep	2	ns	ns	ns
Inoculum	1	ns	ns	0.097
Nitrogen (Ntrt)	1	*	**	0.095
Date	8	***	***	***
Ntrt:Inoculum	1	ns	ns	ns
Date:Ntrt	8	*	ns	ns
Date:Inoculum	8	ns	ns	ns
Date:Ntrt:Inoculum	8	ns	ns	ns

* Significant at 0.05 probability level; ** significant at 0.01 level; *** significant at 0.001 level; ns, nonsignificant at 0.05 level

In this experiment, there was a moderately significant (P -value = 0.097) effect of inoculum treatment on VWC (Table 3). Averaged across date and nitrogen treatment, the AMF-inoculated plots and water-treated plots had VWC of 34.1% and 32.7%, respectively. This may be due to plot location.

Experiment 3

Table 4. ANOVA results of Experiment 3

Source of variation	df	TQ	NDVI	VWC
Rep	2	ns	ns	ns
Inoculum	1	ns	ns	ns
Nitrogen (Ntrt)	1	ns	***	0.059
Date	8	***	***	***
Ntrt:Inoculum	1	ns	0.059	ns
Date:Ntrt	8	**	***	ns
Date:Inoculum	8	ns	ns	ns
Date:Ntrt:Inoculum	8	ns	ns	ns

* Significant at 0.05 probability level; ** significant at 0.01 level; *** significant at 0.001 level; ns, nonsignificant at 0.05 level

In this experiment, there was a moderately significant (P -value = 0.059) effect of nitrogen treatment - inoculum treatment interaction on NDVI (Table 4). Averaged across date, the AMF-inoculated plots subjected to the low nitrogen treatment had the lowest NDVI and the AMF-inoculated plots subjected to the regular nitrogen treatment had the highest NDVI (Figure 2, 3).

Experiment 3 NDVI

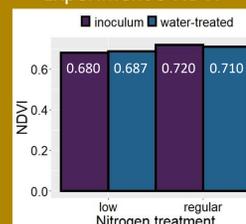


Figure 2. NDVI of inoculum treatment by nitrogen treatment, averaged across date.

Experiment 3 plots

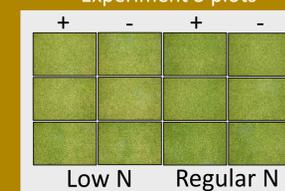


Figure 3. Plot images on 12 Aug 2020 of AMF-inoculated (+) and water-treated (-) plots by nitrogen treatment.

Experiment 4

Table 5. ANOVA results of Experiment 4

Source of variation	df	TQ	NDVI	VWC
Rep	2	ns	ns	ns
Inoculum	1	ns	ns	ns
Nitrogen (Ntrt)	1	*	**	0.086
Date	6	***	***	***
Ntrt:Inoculum	1	*	ns	ns
Date:Ntrt	6	*	***	ns
Date:Inoculum	6	ns	ns	ns
Date:Ntrt:Inoculum	6	ns	ns	ns

* Significant at 0.05 probability level; ** significant at 0.01 level; *** significant at 0.001 level; ns, nonsignificant at 0.05 level

In this experiment, there was a significant effect of nitrogen treatment - inoculum treatment interaction on NDVI (Table 5). Averaged across date, the AMF-inoculated plots subjected to low nitrogen had the lowest TQ (Figure 4). This was likely due to uniformity losses caused by the presence of weeds (Figure 5).

Experiment 4 TQ

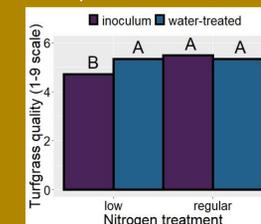


Figure 4. TQ of inoculum treatment by nitrogen treatment, averaged across date.

Experiment 4 plots

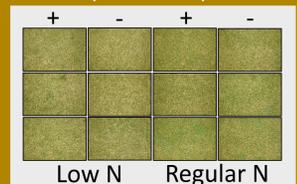


Figure 5. Plot images on 16 July 2020 of AMF-inoculated (+) and water-treated (-) plots by nitrogen treatment.

Conclusion: Additional research is needed on the use of arbuscular mycorrhizal fungi inoculations for established golf greens in northern climates.

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