





Emerging effects of selected rhizosphere properties on transpiration and leaf water potential of two Zea mays L. genotypes in semi-arid environments

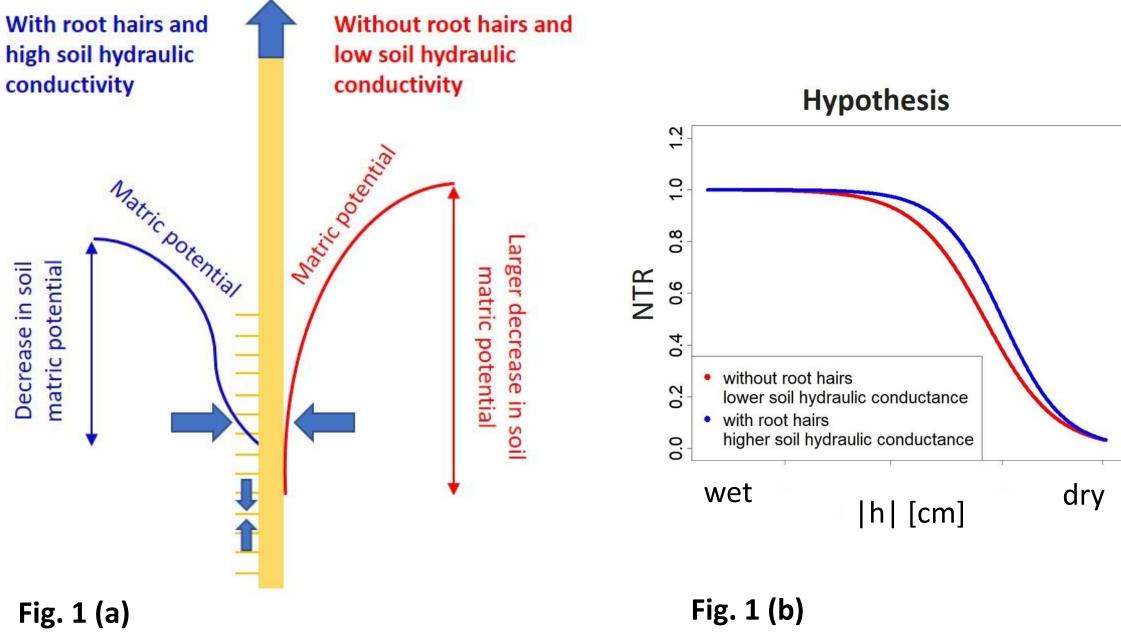
Tina Koehler¹, Daniel-Sebastian Moser¹, Ákos Botezatu¹, Tharanya Murugesan², Sivasakthi Kaliamoorthy², Jana Kholová², Mutez Ahmed¹, Andrea Carminati¹ ¹ Chair of Soil Physics, University of Bayreuth

² Dept. of Crop Physiology, ICRISAT Hyderabad

E-mail: tiekey94@web.de, Daniel-Sebastian.Moser@uni-bayreuth.de, Akos.Botezatu@uni-bayreuth.de, m.tharanya@gmail.com, K.Sivasakthi@cgiar.org, J.Kholova@cgiar.org, Mutez.Ahmed@uni-bayreuth.de, Andrea.Carminati@uni-bayreuth.de

Introduction

This study focused on the effect of belowground processes and rhizosphere traits on soil-plant water relations (Fig 1a and b). Therefore, experiments which investigate the effect of root hairs and water flow dynamics through different soil textures on transpiration, leaf water potential, soil-plant hydraulic conductance and stomata conductance have been tested in drying soils.



Material and Methods

Two maize (*Zea mays* L.) genotypes, a hairy wildtype (Fig. 2a) and a roothairless rth3-mutant (Fig. 2b) were grown in pots, filled with sandy loam, silty clay or sand. They were exposed to soil drying under glasshouse conditions. Soil water content (O) has been monitored by TDRmeasurements. Soil water potential (h) was calculated from the soil water content. Daily transpiration (E) was measured gravimetrically and normalized to facilitate comparison (NTR, [1]). Leaf water potential has been measured at four water stress levels (WW-WS3) during the day (Ψ_{leaf}) and under pre-dawn conditions (=indicator for soil water potential, [2]), using the Scholander pressure chamber. Stomata conductance (gs) has been calculated from the transpiration rate, normalized by leaf area and VPD, normalized by atmospheric pressure [3].

Results and Discussion

The effect of root hairs

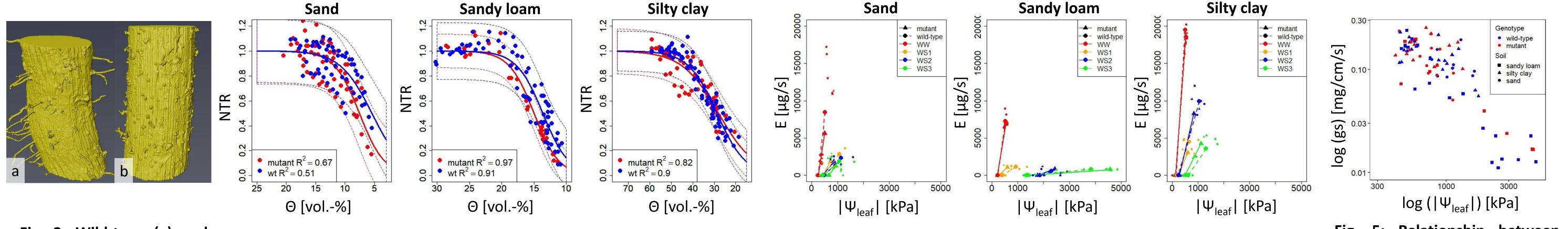


Fig. 2: Wild-type (a) and

Fig. 5: Relationship between

Fig. 3: Relationship between normalized transpiration ratio (NTR) and soil rth3-mutant (b) root. By water content (O) between genotypes. courtesy of P. Duddek

Fig. 4: Relationship between transpiration (E) and absolute leaf water potential ($|\Psi_{leaf}|$) between genotypes.

stomata conductance (gs) and $|\Psi_{leaf}|$ between genotypes.

The effect of root hairs manifested in a delayed drop in NTR with decreasing Θ and/or higher NTR post the transpiration breakpoint in the presence of root hairs (Fig. 3). The effect of root-hairs might be especially pronounced in silty clay because of their ability to bridge the gap between roots and soil as originated by cracks (Fig. 7, [4]) and thereby maintained the connectivity of the liquid phase and water flow eventually [5]. In sand, root hairs are believed to have attenuated the gradients in matric potential around the roots [6], which developed with a drastic drop in soil hydraulic conductivity in coarse-grained soils. No obvious differences between the genotypes were visible in the relationship between Ψ_{leaf} and E (Fig. 4, which is equivalent to the soil-plant hydraulic conductivity (K_{sp}) as well as between gs and E (Fig. 5). This is believed to be explained by the prompt stomata closure under natural conditions [7].

The effect of soil texture

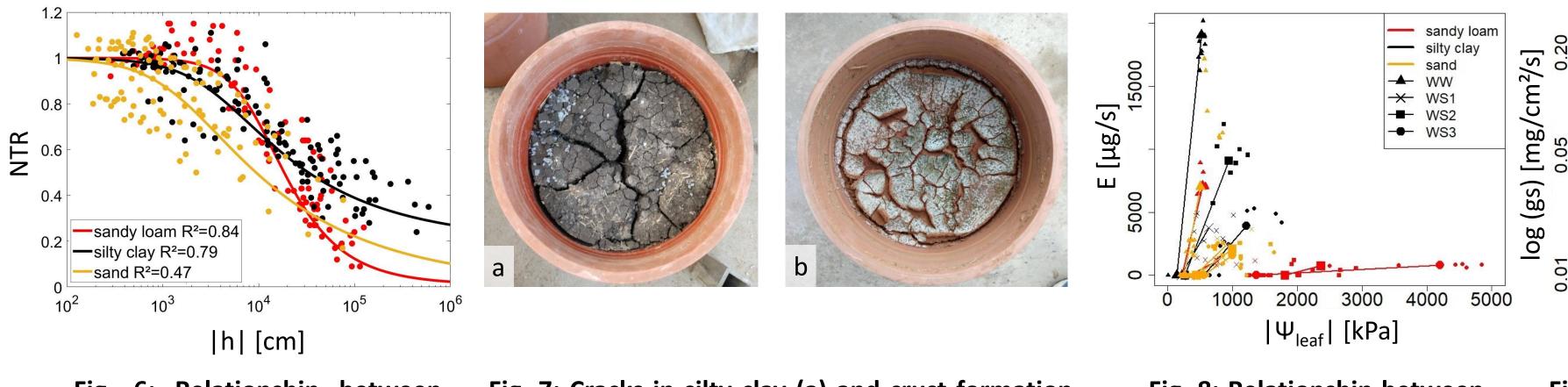


Fig. 6: Relationship between normalized transpiration ratio (NTR) and absolute soil water notontial (Ibl) botwaan cail

Fig. 7: Cracks in silty clay (a) and crust formation in sandy loam (b). By courtesy of D.-S. Moser

Fig. 8: Relationship between transpiation and leaf absolute water notantial (1111 1) hotwar

Fig. 9: Relationship between stomata conductance (gs) and absolute leaf water notontial (1111 1) hot

1000

 $\log(|\Psi_{leaf}|)$ [kPa]

2000

sandy loam $R^2 = 0.8$

silty clay $R^2 = 0.27$ sand $R^2 = 0.22$

500

Plants grown in different soil textures decreased NTR at different h, although not exactly as expected (Fig. 6). This was attributed to soil features like soil cracking or crust formation that might have changed the expected soil hydraulic conductivity (Fig. 7). Moreover, soil texture seemed to have mattered for the development ⁵⁰⁰⁰ of plant- as well as soil hydraulic conductivity, which can be deduced from the differences in K_{sp} at all WS-levels between soils (Fig. 8). Overlapping confidence intervals are suggesting that differences between soil textures in gs

| potential | ([n]) | between | SOII | |
|-----------|-------|---------|------|--|
| textures. | | | | |

COLOR Koehler et al. 2020

| potential ($ \Psi_{leaf} $) between | potential ($ \Psi_{leaf} $) between | | |
|---------------------------------------|---------------------------------------|---|--|
| soil textures. | soil textures. | response to Ψ_{leaf} were not significant (Fig. 9). | |

Conclusion and References

We conclude that: 1.) **root hairs** had a positive impact on root water uptake and therefore transpiration. However, a different experimental setup will be needed to investigate References genotypic differences in soil-plant hydraulic conductivity, as they are expected to become pronounced in dry soils under high transpiration rates, when the relationship between leaf water potential and transpiration becomes non-linear [8]. Furthermore, we have found that: 2.) soil texture had an effect on transpiration with soil drying. Soil texture likely mattered by determining soil hydraulic conductivity, which affected the resistance to water flow between soil and roots and consequently transpiration. There were no clear differences between soil textures in the relationship between gs and Ψ_{leaf} , but in K_{sp}. We conclude that soil texture indirectly affected stomata by changing the soil-plant conductance.

Acknowledgments

The study and the exchange with ICRISAT was funded by DAAD (PPP; Project Nr. 57390361) and DFG.

- [1] Devi, M.J.; Sinclair, T.R.; Vadez, V. (2010): Genotypic variability among peanut (Arachis hypogea L.) in sensitivity of nitrogen fixation to soil drying. Plant and Soil. 330(1-2). 139–148.
- [2] Munns, R.; Passioura, J.B. (1984): Hydraulic Resistance of Plants. III. Effects of NaCl in Barley and Lupin. Functional Plant Biology. 11(5). 351.

[3] Jarvis & McNaughton (1986): Stomatal Control of Transpiration: Scaling Up from Leaf to Region. 15. Elsevier. Sperry, J.S.; Love, D.M. (2015): What plant hydraulics can tell us about responses to climate-change droughts. The New phytologist. 207(1). 14-27.

[4] White, R. G., & Kirkegaard, J. A. (2010). The distribution and abundance of wheat roots in a dense, structured subsoil-implications for water uptake. Plant, cell & environment. 33(2). 133-148.

[5] Carminati, A.; Passioura, J.B.; Zarebanadkouki, M.; Ahmed, M.A.; Ryan, P.R.; Watt, M.; Delhaize, E. (2017): Root hairs enable high transpiration rates in drying soils. New Phytologist. 216(3). 771–781.

[6] Gardner WR (1960): Dynamic aspects of water availability to plants. Soil Science. 89. 63–73.

[7] Hacke, U.G.; Sperry, J.S.; Ewers, B.E.; Ellsworth, D.S.; Schäfer, K.V.R.; Oren, R. (2000): Influence of soil porosity on water use in Pinus taeda. Oecologia. 124(4). 495–505.

[8] Sperry, J.S.; Love, D.M., (2015): What plant hydraulics can tell us about responses to climate-change droughts. The New phytologist. 207(1). 14–27.