Does plant root architecture respond to potassium under water stress? A case from rice seedling root responses

Dipika S. Patel1, Bardhan Kirti2,* , P. Patle Dhiraji2, Parekh Vipulkumar2, Jena Suchismita2, V. Narwade Ajay1 and N. Chhatrola Harshadkumar2

1N.M. College of Agriculture, Navsari Agricultural University, Navsari 396 450, India
2ASPEE College of Horticulture and Forestry, Navsari Agricultural University, Navsari 396 450, India

In plants, roots sense the availability of potassium and water. This study examined the influence of potassium availability on root architecture and above-ground growth under water-deficit condition. We found that the growth of rice seedlings was further reduced by low potassium under water stress. We noted considerable reduction in the root projected area, maximum width and width-to-depth ratio. Furthermore, high potassium supply helped in sustaining the root top and bottom angles and prevented root steepness under mild water stress. However, high potassium availability did not result in better seedling growth. Root steepness was more influenced by water than potassium levels under severe water stress.

Keywords: Drought, Oryza sativa L., potassium, root plasticity.

Depending on the growing conditions, major factors that decide rice yields under drought are thickness, density, depth and distribution of the roots1. Furthermore, changes that occur in the early stage in the root angle, which contribute in the development of deep root systems, play a key role in drought tolerance of rice2. A cross-talk may occur between signalling underlying sensing of water and nutrients by the roots, and in response to the availability of water and nutrients, root signals can regulate the growth of shoots3. Potassium (K) fertilizers are commonly applied to alleviate the adverse effects of water stress on many crops, including rice4-5. High K availability can cause an increase in the root diameter and dry matter6. Hormonal mechanisms are responsible for the effects exerted by K on the roots7. K also plays key roles in maintaining cell-water status8, stomatal opening–closing9 and neutralizing reactive oxygen species10. In addition, during moisture stress, K might affect the spatial distribution of roots, termed as root architecture. However, because most previous studies have focused on root growth11 or have been conducted in the absence of water stress9, the effect of K on the spatial distribution of roots under moisture stress remains unclear. Moreover, although the effect of water deficit on the roots of rice was studied earlier12, the collaborative influence of the availability of water and nutrients remains unclear. Therefore, the present study aims to examine the interactive effects of K and water availability on root architecture and rice seedling growth.

Materials and methods

Rice (Oryza sativa L.) variety NAUR-1, adapted to upland and lowland cultivation was used. Between 2017 and 2018, we conducted randomized, factorial-design experiments with three replicates in a naturally ventilated polyhouse at Navsari Agricultural University (20.9248°N, 72.9079°E). We pre-soaked rice seeds in deionized water for 24 h, with five seeds being sown in each semi-transparent polythene bag (32 × 20.5 cm) containing 1.2 kg of a homogeneous mixture of sand and perlite (ratio of 2 : 1, 800 + 400 g). Post emergence of shoots, we retained only two seedlings in each bag.

We examined three levels of water stresses: no water stress (NWS; field capacity (FC) of 100%), mild water stress (MWS; FC of 60%), and severe water stress (SWS; FC of 40%). We used the gravimetric method to estimate FC. We used four water-saturated bags which had their tops covered to prevent evaporative loss, and kept them overnight to drain excess water. The following formula was used to calculate the amount of water retained at 100% FC: ((weight of saturated media – weight of oven-dried media)/weight of oven-dried media) × 100. The medium was oven-dried at 105 ± 1°C overnight to obtain a constant weight. Day-to-day evapotranspiration loss was recorded by weighing the four water-saturated bags daily and determining the difference in their weights at 100% FC. All the bags were initially replenished daily up to 100% FC using the unmodified Yoshida solution13 until the emergence of seedlings. Subsequently, we used the modified Yoshida solution to apply water stress (Figures 1a and b and 2).

---

*For correspondence. (e-mail: kirtivardhan@nau.in)
We modified the Yoshida solution to develop three treatments of K: K– (Yoshida solution containing 50% low K, 20 ppm K, 35.7 g/l K₂SO₄), K₀ (Yoshida solution containing optimal K, 40 ppm K, 71.4 g/l K₂SO₄), and K+ (Yoshida solution containing 50% high K, 60 ppm K, 107.1 g/l K₂SO₄). Besides maintaining all other nutrient concentrations unchanged, sulphur concentration was adjusted by the amount of H₂SO₄ (specific gravity: 1.84, purity: 98%) according to K₂SO₄ used in the treatments (50 ml/l H₂SO₄ in K₀, 61.14 ml/l in K–, and 38.85 ml/l in K+). pH of the solution was adjusted to 5.5 before application.

We examined 405 root samples that represented the root system of five seedlings in each treatment at three-time intervals (7, 14 and 21 days after emergence (DAE)). Before scanning, the roots were placed under water in a scanner acrylic tray (23.1 × 15.5 × 7 cm³) for 45 min and then stained with a natural red dye (0.25 g/l) to optimize the contrast and decrease the diffraction of scanner light from the roots. Root images were taken using a scanner (HP Scanjet G2410) at a resolution of 600 dpi. A 40 mm diameter circular scale marker placed under the scanner tray and all the images were captured with the scale marker (Figure 1 c). To calculate root dry weight, we used the root samples of 21 DAE. We kept these samples in a hot-air oven at 65°C ± 1°C overnight till a constant weight was achieved.

We computed the root architecture features through digital imaging of root traits (DIRT; figure 3)⁴. All scanned images were adjusted at a masking threshold of 3 before computation (Figure 1 d and e). We converted DIRT results in pixels to metric units, except for the...
average root density (which is the ratio of foreground to background pixels within the root shape) using the following formula

\[
\text{Calibration factor} = \frac{\text{Scale marker diameter}}{\text{Circle ratio (pixels)}}.
\]

The circle ratio (pixels) was computed by the DIRT platform.

Root depth (mm) = Length (pixels) \times \text{calibration factor}.

Maximum width (mm) = \text{Maximum width (pixels)} \times \text{calibration factor}.

Projected root area (mm\(^2\)) = \text{Area (pixels)} \times \text{calibration factor}^2.

At 21 DAE, we measured the length (cm) and dry weight (g) of shoots of the same seedlings from each treatment that were sampled for root imaging. Analysis of variance was used to estimate treatment effects and we considered differences between mean values to be statistically significant if \( p \geq 0.05 \) (Table 1).\(^5\)

### Results

#### Vertical features of root architecture

We found that the projected root area (PRA), reduced with an increase in water stress, and a low K level caused a larger decrease in PRA (Figure 4a). A reduction of 51% and 66% was observed in PRA under MWS and SWS respectively, with a low K supply (K\(^-\)) when compared with NWS seedlings grown with optimal K (K\(^0\)). In addition, a 25% decrease in PRA was noted due to low K supply under the water stress treatments (K\(^-\)-MWS and K\(^-\)-SWS) compared with that under optimal K level. We found no significant effect of high K level (K\(^+\)) under MWS and SWS treatments on root area compared to MWS and SWS treatments with optimum K supply.

Although seedling root depth (RD), increased under water stress, it decreased by up to 28.7% and 31.1% under MWS and SWS respectively, at low K level when compared with MWS and SWS at optimal K level (Figure 4b). However, in contrast to low K level, a high K level increased RD by 9.3% and 10% in MWS and SWS over optimum K level respectively. Moreover, the influence of high K on RD under the absence of water stress was not strong.
We observed that the average root density decreased with increase in water stress (Figure 4c). Compared with an optimal K level under MWS, a low K level under MWS reduced root density by 33%. However, the root density values from seedlings grown at a high K level under MWS were statistically equal to those obtained with an optimal K level under NWS. We did not find any significant difference in root density between SWS with a low or high K levels and SWS with optimum K level.

**Horizontal features of root architecture**

We observed that the maximum width (MW) of the roots reduced with an increase in water deficit (Figure 5a). The root width substantially decreased under low K level. Compared with MWS and SWS with optimal K supply, a reduction of 41% and 43% in root width was noted under MWS and SWS respectively, with low K availability.

**Table 1.** Summary of analysis of variance for water stress (WS), potassium (K) and their interactions (WS × K) on root architecture and seedling growth

| Parameters                       | WS | K | WS × K |
|----------------------------------|----|---|--------|
| Projected root area (mm²) at     |    |   |        |
| 7 DAE                            | *  |   | ns     |
| 14 DAE                           | *  |   |        |
| 21 DAE                           | *  | * |        |
| Root depth (mm) at               |    |   |        |
| 7 DAE                            | *  | * | ns     |
| 14 DAE                           | *  | * |        |
| 21 DAE                           | *  | * |        |
| Maximum width (mm) at            |    |   |        |
| 7 DAE                            | *  | * | ns     |
| 14 DAE                           | *  | * |        |
| 21 DAE                           | *  | * |        |
| Average root density at          |    |   |        |
| 7 DAE                            | *  | * | ns     |
| 14 DAE                           | *  | * |        |
| 21 DAE                           | *  | * |        |
| Maximum width-to-depth ratio at  |    |   |        |
| 7 DAE                            | *  | * |        |
| 14 DAE                           | *  | * |        |
| 21 DAE                           | *  | * |        |
| Root top angle (°) at            |    |   |        |
| 7 DAE                            | *  | * | ns     |
| 14 DAE                           | *  | * | ns     |
| 21 DAE                           | *  | * |        |
| Root bottom angle (°) at         |    |   |        |
| 7 DAE                            | *  | * | ns     |
| 14 DAE                           | *  | * | ns     |
| 21 DAE                           | *  | * |        |

*Root parameters of 7 and 14 DAE (days after emergence) are presented in the Supplementary Figures 1 and 2 and total dry weight (g) in Supplementary Figure 3. *Significant at p ≥ 0.05, ns: not significant.

However, we found that seedlings supplied with high K under MWS and SWS showed 20% and 48% more wide roots respectively, against seedlings of MWS and SWS with optimal K supply.

Water stress caused a reduction in the MW/RD ratio (Figure 5b). We observed that the MW/RD ratio increased by 6.5% at high K level, but decreased by 12.3% with low K supply compared with that under optimal K level for seedling under no water stress. Although K availability did not affect the MW/RD ratio at 21 DAE under different water stress treatments, there was a significant interaction at 7 and 14 DAE (Supplementary Figure 2a and b).

We observed the narrowest root top angle (RTA) under NWS and the widest SWS (Figure 5c). Compared with NWS, RTA increased by 6° and 17.2° under MWS and SWS respectively, with optimal K level. However, similar effects of different K levels on RTA were found under MSW. Rice seedlings that were exposed to SWS with a high K level exhibited narrower RTA (49.9°) compared with those exposed to SWS with a low K supply (61.9°). However, RTA determined under SWS with high K level was similar to that under MWS with optimal K level (46.0°).

Water stress also increased the root bottom angle (RBA, Figure 5d). Compared with NWS (55.0°), RBA values were 63.7° and 74.1° under MWS and SWS respectively, with optimal K supply. Furthermore, rice seedlings demonstrated RBA values of 71.7° and 79.4° when treated with MWS and SWS respectively, with low K level. The values increased by 8° and 5.3° under MWS and SWS with optimal K supply respectively. These findings indicate that roots are more narrowly spread at the
bottom under water stress with low K availability. However, no difference in RBA was observed between seedlings grown with high K supply under MWS and those supplied with optimum K under MWS. Under SWS, high K supply reduced RBA by 10° over optimal K level.

**Root dry weight**

We observed that root dry weight (RDW), decreased with increase in water stress (Figure 6b). Compared with RDW of seedlings grown under MWS and SWS with optimal K supply, a reduction of 31% and 49% was noted in RDW of rice seedlings grown under MWS and SWS at a low K level respectively. Also, compared with rice seedlings grown under SWS with optimum K level, an increase of 15.6% was observed in the RDW of rice seedlings grown under SWS at high K level. However, we did not find such an increase in seedlings treated with MWS.

**Seedling growth**

The shoot length of rice seedlings subjected to MWS and SWS with optimum K supply was 16% and 30%, lower than that of seedlings under NWS respectively (Figure 6a). Furthermore, no significant difference in shoot length was observed between seedlings supplied with low and high K solution under NWS. However, we found that the shoot length of seedlings exposed to MWS and SWS at low K level was 21% and 24% lower respectively, than that of seedlings grown under MWS and SWS with optimum K. Furthermore, the supply of high K to rice seedlings exposed to MWS and SWS showed increased shoot length by 12.8% and 24.8% respectively, against to those grown under MWS and SWS with optimum K supply. The shoot dry weight of seedlings reduced with an increase in water stress (Figure 6b). This reduction was higher by 40% in rice seedlings exposed to MWS at low K level than those exposed to MWS at optimum K level. The shoot dry weight values were similar between seedlings subjected to MWS at high K level and those subjected to MWS at optimum K level. However, no significant difference in shoot dry weight values was observed between seedlings supplied with low and high K under SWS. The total dry weight values were similar between seedlings under MWS and SWS with high K level and those under MWS and SWS at optimum K level (Supplementary Figure 3). However, seedlings from MWS and SWS at low K level exhibited a significant decrease in total dry weight.

**Discussion**

Although the genetics of the rice variety regulates its root architecture, a broad variation of phenotypic plasticity is observed due to the availability of nutrients and water, and this is crucial for the drought tolerance of rice. In the present study, we found water and K availability both interact in regulating the root system of rice seedlings (Figures 7 and 8). We observed a decrease in PRA and average root density with an increase in water stress. This detrimental effect was more at a low K level; however, such reduction in PRA and average root density was not seen in seedlings exposed to MWS and SWS at optimum and high K levels. Similarly, a study reported that K increased the root surface area, volume and root tip number in tomato under atmospheric drought. The strong effects of water deficit on these traits indicate that both K and water are limiting factors that may independently regulate these root traits. The translocation of sugar depends on K availability which triggers cell division, whereas cell elongation depends on cell turgor and water content. The results of this study indicate that an optimum K level is crucial for the development of the root system because rice seedlings exposed to NWS with a low K level exhibited decreased root area compared with seedlings exposed to NWS with an optimum K level. This finding may be attributed to the loss of turgidity resulting from low water absorption and the sub-optimal levels of auxin in the root tip under low K availability.

The horizontal and vertical elongation of the roots is responsible for the exploration of soil area and uptake of water.
soil resources. In the present study, we calculated the root depth, maximum width and ratio of maximum width to root depth to determine whether the roots expand preferentially in the horizontal or vertical direction. We found that the root depth increased and maximum width decreased with increase in water stress, resulting in a low MW/RD ratio. This finding indicates that with increasing water deficit, the roots expand vertically rather than horizontally to reach the water source deep down in the soil, thus resulting in steeper root growth. However, with K sufficiency under water stress, the roots expanded horizontally, resulting in an increased root width and MW/RD ratio. We also observed that roots exhibiting steep growth had wide root top and bottom angles under MWS and SWS, suggesting that the roots had lesser surface contact area with the surrounding medium. On farmlands, these characteristics can restrict the acquisition of water and nutrients efficiently from the soil. In this study, high K supply helped maintain the top and bottom angles in seedlings exposed to MWS. However, high K availability did not exert this positive effect in seedlings exposed to SWS. These results suggest that in addition to increasing the translocation of mineral nutrients in the xylem under water stress, K fertilization maintains RTA and RBA, thus increasing the root–soil contact area and may play a key role to nutrient acquisition by reaching more soil areas. The synthesis, transport and signalling of auxin are the main factors involved in the development of rice roots. An auxin-inducible DRO gene controls root depth. This gene regulates the root top angle, thus deciding the direction of root elongation. However, the mechanisms through which K availability results in variations in root top and bottom angles remain unclear. However, K was found to affect auxin signalling and...
translocation in rice; the K transporter altered membrane-bound auxin efflux proteins and may change the root angle.

Water stress caused a reduction in seedling growth, and reduction in root dry weight, shoot dry weight, and seedling height was more with low K availability. However, high K level could ameliorate these effects and increase shoot length, shoot dry weight and root dry weight. The beneficial effects of a high K level were more pronounced in seedlings exposed to MWS. The application of high K under SWS did not increase total dry weight compared with optimum K level under SWS. These results are in agreement with that of a previous study.

Conclusion

This study shows that the interaction of K and water stress influences the three-dimensional distribution of roots. This can be beneficial for exploring underground resources in order to promote the growth of rice seedlings above the ground under MWS depending upon K availability. The results emphasize the need to maintain K availability in soils under water stress. These findings provide insights into the role of K in drought tolerance through modification of root architecture. Additional studies should examine the degree to which root architecture is shaped by water and K availability in a broad genetic background.

1. Gowda, V. R. P., Henry, A., Yamcuachi, A., Shashidhar, H. E. and Serraj, R., Root biology and genetic improvement for drought avoidance in rice. *Field Crops Res.*, 2011, **122**, 1–13.

2. Uga, Y., Sugimoto, K., Ogawa, S., Rane, J., Ishitani, M. and Hara, N., Control of root system architecture by DEEPER ROOTING 1 increases rice yield under drought conditions. *Nature Genet.*, 2013, **45**, 1097–1102.

3. Kudoyarova, G. R., Dodd, I. C., Veselov, D. S., Rothwell, S. A. and Yu Veselov, S., Common and specific responses to availability of mineral nutrients and water. *J. Exp. Bot.*, 2015, **66**, 2133–2144.

4. Gezebsiz, W., Gransee, A., Szczepaniak, W. and Diatta, J., The effects of potassium fertilization on water-use efficiency in crop plants. *J. Plant Nutr. Soil Sci.*, 2013, **176**, 355–374.

5. Zain, N. A. M. and Ismail, M. R., Effects of potassium rates and types on growth, leaf gas exchange and biochemical changes in rice (*Oryza sativa*) planted under cyclic water stress. *Agric. Water Manage.*, 2016, **164**, 83–90.

6. Filho, A. C. A. C., Crusciol, C. A. C., Nascente, A. S., Mauad, M. and Garcia, R. A., Influence of potassium levels on root growth and nutrient uptake of upland rice cultivars. *Rev. Coatinga*, 2017, **30**(1), 32–44.

7. Yasuni, J., Endo, N. and Kono, Y., Root and partitioning of 14C-labelled photosynthate in the seminal root of corn seedlings as affected by light intensity. *Jpn. J. Crop Sci.*, 1992, **61**, 271–278.

8. Mengel, K. and Arneke, W. W., Effect of potassium on the water potential, the pressure potential, the osmotic potential and cell elongation in leaves of *Phaseolus vulgaris*. *Physiol. Plant.*, 1982, **54**, 402–408.

9. Benlloch-González, M., Arquero, O. J., Fournier, M., Barranco, D. and Benlloch, M., K starvation inhibits water-stress-induced stomatal closure. *J. Plant Physiol.*, 2008, **165**, 623–630.

10. Cakmak, I., The role of potassium in alleviating detrimental effects of abiotic stresses in plants. *J. Plant Nutr. Soil Sci.*, 2005, **168**, 521–530.

11. Jordan-Meille, L., Martineau, E., Bornot, Y., Lavres, J., Abreu Jr, H. C. and Domec, J. C., How does water-stressed corn respond to potassium nutrition? A shoot-root scale approach study under controlled conditions. *Agricultura*, 2018, **8**, 180.

12. Kano-Nakata, M. et al., Functional roles of the plasticity of root system development in biomass production and water uptake under rainfed lowland conditions. *Field Crops Res.*, 2013, **144**, 288–296.

13. Yoshida, S., Forno, D. A., Cock, J. H. and Gomez, K. A., *Laboratory Manual for Physiological Studies of Rice*, The International Rice Research Institute, Philippines, 1976, 2nd edn.

14. Das, A. et al., Digital imaging of root traits (DIRT): a high throughput computing and collaboration platform for field-based root phenomics. *Plant Methods*, 2015, **11**, 51.

15. Gomez, K. A. and Gomez, A. A., *Statistical Procedures for Agricultural Research*, John Wiley, New York, USA, 1984, 2nd edn.

16. Shahzad, Z. and Antmann, A., Food for thought: how nutrients regulate root system architecture. *Curr. Opin. Plant Biol.*, 2017, **39**, 80–87.

17. Giehl, R. F. H. and Von Wieren, N., Hydropatterning – how roots test the waters. *Science*, 2018, **362**, 1358–1359.

18. Zhang, J., Jiao, X., Du, Q., Song, X., Ding, J. and Li, J., Effects of vapor pressure deficit and potassium supply on root morphology, potassium uptake, and biomass allocation of tomato seedlings. *J. Plant Growth Regul.*, 2020; https://doi.org/10.1007/s00344-020-10115-2.

19. Martineau, E., Domec, J. C., Bosc, A., Dannoura, M., Gibon, Y., Bernard, C. and Jordan-Meille, L., The role of potassium on maize leaf carbon exportation under drought condition. *Acta Physiol. Plant.*, 2017, **39**, 219.

20. Wang, L. and Ruan, Y. L., Regulation of cell division and expansion by sugar and auxin signaling. *Frontiers Plant Sci.*, 2013, **4**, 163.

21. Gerarddeaux, E., Jordan-Meille, L., Constantin, J., Pellerin, S. and Dingkuhn, M., Changes in plant morphology and dry matter partitioning caused by potassium deficiency in *Gossypium hirsutum* (L.). *Environ. Exp. Bot.*, 2010, **67**, 451–459.

22. Song, W. et al., Potassium deficiency inhibits lateral root development in tobacco seedlings by changing auxin distribution. *Plant Soil*, 2015, **396**(12), 163–173.

23. Hasanuzzaman, M. et al., Potassium: a vital regulator of plant responses and tolerance to abiotic stresses. *Agronomy*, 2018, **8**, 31.

24. Wang, Y., Zhang, T., Wang, R. and Zhao, Y., Recent advances in auxin research in rice and their implications for crop improvement. *J. Exp. Bot.*, 2018, **69**, 255–263.

25. Yang, T. et al., The potassium transporter OsHAK5 alters rice architecture via ATP-dependent transmembrane auxin fluxes. *Plant Commun.*, 2020, **1**, 100052.

26. Suatr, M., Soukup, A. and Tylova, E., Potassium in root growth and development. *Plants*, 2019, **8**, 435.

27. Zain, N. A. and Ismail, M. R., Effects of potassium rates and types on growth, leaf gas exchange and biochemical changes in rice (*Oryza sativa*) planted under cyclic water stress. *Agric. Water Manage.*, 2016, **164**, 83–90.

Received 22 January 2020; revised accepted 4 December 2020