

# Salinity Tolerance Mechanisms in Plants: A Physiological, Biochemical and Molecular Characterization Study: A Review

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**Abstract – Salinity is a significant abiotic stress that limits plant growth and profitability in numerous areas of the world because of the increasing use of helpless water quality for water systems and soil salinization. Complex physiological characteristics, metabolic pathways, and atomic or quality networks are involved in plant variation or resistance to salinity pressure. Diverse versatile responses to salinity stress at sub-atomic, cell, metabolic, and physiological levels have been recognized in late research, in spite of the fact that the mechanisms underlying salinity resistance are a long way from completely understood. This paper offers a comprehensive audit of significant advances in research on biochemical, physiological, and atomic mechanisms that control plant transformation and salinity stress resilience.**

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## INTRODUCTION

Salinity is one of the main threats to farming, having been one reason for the demise of the old Sumerian human progress in Mesopotamia (1) and right now causing enormous yearly financial losses of more than USD 10 billion (this figure exceeds the gross domestic result of in excess of 50 less created countries in the advanced world (2). The urgent starting point for determining salinity resistance in plants is particle transport: this includes cation and anion transport through the root cell plasma membranes, transport through the vacuolar membranes, xylem and phloem significant distance particle transport, and salt discharge and collection by specialized cells.

Numerous aspects of particle transport under salinity stress are not yet surely known right now of the aggregation of this research theme. The particle channel and transporter structure-work relationships are slowly being interpreted, however remain backcountry in essence. Different links between particle fluxes, electrophysiology and other physiological processes are often not described in any detail, leading to salinity resistance. Under salinity stress, totally unforeseen features of particle transport might be waiting to be discovered. Researchers in particle transport and salinity resilience of plants (and parasites) were pulled in to the research subject and we combined our efforts to accomplish a more extensive, more definite understanding of salt resistance in particle transport-intervened plants.

The mechanisms of salt stress response and resistance have eluded classification, despite fair progress in identifying their physiological manifestations. The integrated salt metabolism of plants is considered in this study, essentially as an issue in meganutrient physiology, to discuss advances in research on the complex physiological and atomic mechanisms involved in plant salinity resistance.

## PHYSIOLOGICAL AND BIOCHEMICAL MECHANISMS OF SALT TOLERANCE:

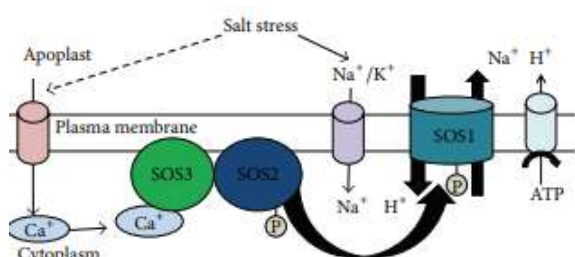
### Ion Homeostasis and Salt Tolerance:

NaCl is the main type of salt present in the soil, so the essential focus of research is the study of the Na<sup>+</sup> particle transport mechanism and its compartmentalization. The Na<sup>+</sup> particle entering the cytoplasm is then transported through the Na<sup>+</sup>/H<sup>+</sup> antiporter to the vacuole. In the vacuolar film, there are two types of H<sup>+</sup> pumps: vacuolar type H<sup>+</sup>-ATPase (V-ATPase) and vacuolar pyrophosphatase (V-PPase). Of these, the most dominant H<sup>+</sup> siphon present within the plant cell is V-ATPase. In non-stress environments, it plays an important role in sustaining solute homeostasis, energising secondary transport, and promoting vesicle fusion. The survivability of the plant depends on V-ATPase development under stressed circumstances (3, 4). Increasing proof demonstrates the significance of a Salt Overly Sensitive (SOS) stress signaling pathway in

molecule homeostasis and salt resilience (Figure 1).

A powerful method has been created by numerous plants to sustain the union of ions in the cytoplasm at a low level. In preserving molecule obsession inside the cytosol by controlling molecule take-up and transport during the hour of stress, membranes assume an integral occupation alongside their associated components. The transport wonder is controlled by various transporter proteins, channel proteins, antiporters and symporters.

Mama et al. showed that Arabidopsis NADPH oxidases AtrbohD and AtrbohF work under salt stress in the ROS-subordinate  $\text{Na}^+/\text{K}^+$  homeostasis guidelines in Arabidopsis. The plants sustain a raised  $\text{K}^+$  level of around 100 mM inside the cytosol, suitable for cytoplasmic protein development. Within the vacuole, the  $\text{K}^+$  obsession varies between 10 mM and 200 mM. The vacuole acts as the largest pool of  $\text{K}^+$  inside the plant cell. In keeping the turgor within the cell,  $\text{K}^+$  plays a significant job. It is passed on through  $\text{K}^+$  transporter and layer channels into the plant cell against the focus angle. When the extracellular  $\text{K}^+$  fixation is low, high affinity  $\text{K}^+$  take-up mechanisms are intervened by  $\text{K}^+$  transporters, while low affinity take-up is done by  $\text{K}^+$  channels when the extracellular  $\text{K}^+$  focus is high. The mechanism of take-up is thus mainly determined by the convergence of  $\text{K}^+$  accessible in the soil. Then again, the cytosol maintains an extremely low  $\text{Na}^+$  particle fixation (around 1 mM or less).  $\text{Na}^+$  particles contend with  $\text{K}^+$  for the transporter because of increased  $\text{Na}^+$  obsession in the soil during salinity stress, as both share the same transport mechanism, thus reducing  $\text{K}^+$  take-up (5).



**Figure 1: Model of SOS pathway for salinity stress responses.**

#### Compatible solute & osmotic defence accumulation:

At the hour of exposure to salinity stress, amino acids such as cysteine, arginine and methionine, which constitute around 55 percent of absolute free amino acids, decrease while proline obsession increases in response to salinity stress. Proline gathering is a notable measure for salinity stress help that has been embraced. Intracellular proline, which is aggregated during salinity stress, provides stress resilience, yet additionally serves during stress recuperation as a natural nitrogen reserve. Proline is

synthesized either from ornithine or from glutamate. Glutamate in osmotically stressed cells works as the essential precursor. Two significant enzymes, pyrroline carboxylic corrosive synthetase and pyrroline carboxylic corrosive reductase, are essential for the biosynthetic pathway. In the overproduction of proline in plants, both these administrative steps are used. It operates as an  $\text{O}_2$  quencher and thus reveals its cancer prevention agent limit. This was noted in a study directed by Matysik et al. Ben Ahmed et al. observed that proline supplements improved olive salt resistance (*Olea europaea*) by improving the movement of certain cell reinforcement enzymes, photosynthetic action and plant growth, and maintaining a fitting plant water status under salinity conditions (6-9).

Polyols are compounds accessible for natural reactions with various hydroxyl useful groups. Sugar alcohols are a class of polyols that work as viable solutes, as chaperones of low atomic weight and as compounds for ROS scavenging. They can be arranged into two main types, cyclic (for example pinitol) and non-cyclic (for example pinitol) (e.g., mannitol). During stressed periods, Mannitol synthesis is induced in plants by the activity of NADPH-subordinate mannose-6-phosphate reductase. These viable solutes go about as a compound or layer structure defender or stabilizer that is susceptible to parchedness or ionically induced harm. Pinitol is collected when the plant is subjected to salinity stress within the plant cell. The biosynthetic pathway consists of two main steps: myo-inositol methylation, leading to the arrangement of an intermediate compound, ononitol, which is epimerized to frame pinitol. The imt-encoded inositol methyl transferase compound plays a significant part in the synthesis of pinitol. In plants, the transformation of the imt quality shows a result similar to that observed for the mltid quality (10).

#### Antioxidant Regulation of Salinity Tolerance:

Anthocyanin is a flavonoid ordinarily revealed for collection in plants exposed to salt stress. Van Oosten et al. isolated the anthocyanin-impaired response-1 (air1) crack that can not gather anthocyanins under salt stress. The air1 crack showed a deformation in anthocyanin advancement in response to salt stress, yet not to other stresses such as raised light, low phosphorous, high temperature, or dry season stress. This specificity indicated that the air1 transformation in response to salt stress didn't influence anthocyanin biosynthesis, yet rather its guideline. The discovery and description of AIR1 opens avenues for dissecting the connections between abiotic stress and cancer prevention agent gathering in the type of flavonoids and anthocyanins (11).

### **Polyamines' Function in Salinity Tolerance:**

The PA biosynthetic pathway has been completely investigated and investigated inside and out in various species, including plants. PUT is the smallest polyamine and is synthesized from either ornithine or arginine through the activity of ornithine decarboxylase (ODC) and arginine decarboxylase (ADC) enzymes, respectively. N-carbamoyl-putrescine is gone over to PUT through the substance N-carbamoyl-putrescine aminohydrolase. The PUT thus acts as a significant substrate for higher polyamines such as SPD and SPM to be biosynthesized. By successively extending the aminopropyl assortment to PUT and SPD, triamine SPD and tetramine SPM are synthesized by the enzymes spermidine synthase (SPDS) and spermine synthase (SPMS). The ODC pathway is the most widely recognized polyamine synthesis pathway found in plants. They separated and cloned most genes involved in the ODC pathway. In either case, there are numerous plants where the ODC pathway is absent; polyamines are synthesized through the ADC pathway in Arabidopsis, for instance. All the genes involved in the pathways of polyamine biosynthesis, similar to Arabidopsis, have been recognized from various plant species. Again, SPM synthase, thermospermine synthase, agmatine iminohydrolase and amidohydrolase N-carbamoylputrescine are simply expressed by single genes (12).

### **Roles of Nitric Oxide in Salinity Tolerance:**

Cancer prevention agent activities and the adjustment of the ROS detoxification system have been credited to the positive effects of NO on salinity resistance or stress relief. Increases in disease avoidance specialist enzymes such as SOD, CAT, GPX, APX, and GR, and the turn of events or lipid peroxidation of malondialdehyde (MDA) suppression were linked to improved plant growth under salinity stress through exogenous use of NO. The effects of NO on salinity resistance are also associated with its plasma film H<sup>+</sup>-ATPase and Na<sup>+</sup>/K<sup>+</sup> proportion guideline. NO stimulates H<sup>+</sup>-ATPase (H<sup>+</sup>-PPase), thus generating a H<sup>+</sup> slope and providing the Na<sup>+</sup>/H<sup>+</sup> trade power. Such an increase in the trading of Na<sup>+</sup>/H<sup>+</sup> may prompt homeostasis of K<sup>+</sup> and Na<sup>+</sup>. Albeit NO acts under salt stress as a signal particle and induces salt resistance by increasing the movement of PM H<sup>+</sup>-ATPase, research results from Zhang et al. with *Populus euphratica* calluses also indicated that NO, in any event in vitro, cannot actuate purged PM H<sup>+</sup>-ATPase action. Initially, they hypothesized that ABA or H<sub>2</sub>O<sub>2</sub> may be downstream signal molecules to manage PM H<sup>+</sup>-ATPase movement. Extra results showed that the substance of H<sub>2</sub>O<sub>2</sub> increased significantly under salt stress. Since H<sub>2</sub>O<sub>2</sub> could be the candidate downstream signal atom, through the expansion of H<sub>2</sub>O<sub>2</sub>, Zhang

et al. tested PM H<sup>+</sup>-ATPase action and K to Na proportion in calluses. The results indicated that H<sub>2</sub>O<sub>2</sub> inducing increased action of PM H<sup>+</sup>-ATPase resulted in an increased K to Na proportion leading to transformation of NaCl stress (13-15).

### **Hormone Regulation of Salinity Tolerance:**

Significant phytohormone ABA whose application to plants enhances the impact of stress conditions (s). As a chemical that is upregulated because of soil water deficiency around the root, it has for some time been perceived. Osmotic stress and water insufficiency are caused by salinity stress, increasing the creation of ABA in shoots and roots. The inhibitory impact of salinity on the photosynthesis, growth, and translocation of assimilates can be moderated by ABA aggregation. The positive relationship between the aggregation of ABA and the resilience of salinity has been in any event somewhat credited to the collection of K<sup>+</sup>, Ca<sup>2+</sup> and viable solutes in root vacuoles, such as proline and sugars, which balance the absorption of Na<sup>+</sup> and Cl<sup>-</sup>. ABA is a fundamental cell signal that modulates the expression of various genes responsive to salt and water deficits. The effects of ABA on the expression of two genes, HVP1 and HVP10, for vacuolar H<sup>+</sup>-inorganic pyrophosphatase, and of HvVHA-A, for the vacuolar H<sup>+</sup>-ATPase synergist (subunit A) in *Hordeum vulgare* under salinity stress were demonstrated by Fukuda and Tanaka. ABA treatment in wheat induced the expression under salinity stress of MAPK4-like, TIP 1, and GLP 1 genes (16).

### **Transcriptional Regulation and Gene Expression of Salinity Tolerance:**

The guideline of quality expression in salinity stress includes a wide scope of mechanisms used by plants to manage or decrease (increase or decrease) the creation of specific quality products (protein or RNA). During the focal creed, various mechanisms of quality guideline, from transcriptional initiation to RNA processing and posttranslational adjustment of a protein, have been recognized.

The most significant regulators that control quality expressions are considered transcription factors. Among them, countless stress-responsive members include bZIP, WRKY, AP2, NAC, C2H2 zinc finger quality, and DREB families. Through binding to the specific cis-acting component in the promoters of these genes, these transcription factor genes are fit for controlling the expression of an expansive scope of target genes. Johnson et al. observed that when exposed to long haul salinity, the expression of bZIP genes was upregulated in salt-sensitive wheat cultivars yet decreased in salt-

open minded varieties. In both rice and wheat, overexpression of a NAC transcription factor confers salt resilience, thus predicting their job in stress alleviation. DREB1/CBF, DREB2, and AREB/ABF are involved in transcriptional rice regulators that have been shown to assume a significant part in reactions to abiotic stress. Under salinity stress, transcription factors such as OsNAC5 and ZFP179 demonstrate upregulation that can compel protein synthesis and accumulation such as proline, sugar and LEA, which in turn plays an integral job in resistance to stress. Salt stress results in the upregulation of AtWRKY8 in Arabidopsis, which straightforwardly binds to the RD29A advertiser, suggesting that it is one of AtWRKY88's objective genes (17, 18).

There are several transcription variables that are restricted by various kinases and have been discovered to be significant players in the transition to salinity stress of plants. Serra et al. showed that OsRMC encodes a receptor-like kinase perceived in rice as a controller of negative salt stress response. In the OsRMC advertiser, two transcription factors belonging to the AP2/ERF family, OsEREBP1 and OsEREBP2, were shown to bind to the same GCC-like DNA theme and direct its quality expression the opposite way. Also, Serra et al. uncovered that the transcript level of OsEREBP1 is not significantly influenced by salt, ABA, or severe cold (5° C) and is just slightly constrained by dry spell and moderate virus. Analysis of value expression with contrasting salt resilience in rice varieties further suggests that OsEREBP2 is involved in the salt stress response of rice. Rice has also been perceived as an ABRE binding transcription factor class bZIP, known as OSBZ8, and has been shown to be unusually expressed in salt tolerant varieties as opposed to salt sensitive varieties. Likewise, during salinity stress, OSBZ8 was shown to be actuated/phosphorylated by the SNF-1 serine/threonine kinase bunch in the presence of Spd (19).

### Bioengineering for Improving Salinity Tolerance

The innovation of hereditary transformation allows scientists to precisely and typically accomplish quality transmission. Hereditary engineering approaches would therefore be useful to control the biosynthetic pathways of osmoprotectants to collect such molecules that demonstration by scavenging ROS, reducing lipid peroxidation, preserving the structure and functions of protein. Several works on the transformation of plants focus on genes controlling particle transport to improve salinity resilience, as Na<sup>+</sup> take-up and compartmentalization guideline is a basically significant mechanism for plant survival under salinity stress, and numerous candidate genes have been distinguished to control this mechanism. Engineering plants are recognized as a viable strategy for generating salt-open minded plants for the over-expression of genes encoding antiporters (Table 1). Contrasted and the use of

inducible promoters or cell type-specific promoters, quality expression studies using constitutive promoters give restricted natural information. The selection of promoters can significantly affect the result of transgenic control. Through successful fine-tuning of the stress response by engineering novel administrative targets, appropriate understanding of post-translational modifications that control plant growth execution under stress, over-expression of miRNAs or their targets, maintenance of chemical homeostasis to forestall pleiotropic effects under stress, and use of plant synthetic science, salt open minded crops could therefore be engineered (20).

**Table 1: Improving plant salt tolerance through engineering genes for various membrane antiporters**

Transgenic host	Gene engineered	Source	Improved functions under salinity stress
Arabidopsis	Vacuolar Na <sup>+</sup> /H <sup>+</sup> antiporter AtNHX1	Albida (Medicago sativa)	Increased osmotic tolerance; ME1A content rises.
Rice	Vacuolar Na <sup>+</sup> /H <sup>+</sup> Antiporter PgNHX1	Pennisetum glaucum	Elaborate root system.
Wheat	Vacuolar Na <sup>+</sup> /H <sup>+</sup> Antiporter AtNHX1	Arabidopsis thaliana L.	Increase in grain yield and biomass production; Accumulation of K <sup>+</sup> in leaf; Reduced aggregation of Na <sup>+</sup> .
Tobacco	Vacuolar Na <sup>+</sup> /H <sup>+</sup> antiporter GhNHX1	Gossypium hirsutum	Na <sup>+</sup> compartmentalization.
Tomato	Vacuolar Na <sup>+</sup> /H <sup>+</sup> antiporter AtNHX1	Arabidopsis thaliana L.	Over production of vacuolar Na <sup>+</sup> /H <sup>+</sup> antiporter.
Tobacco	Vacuolar Na <sup>+</sup> /H <sup>+</sup> antiporter AtNHX1	Arabidopsis thaliana L.	Compartmentalization of Na in roots; Maintenance of K <sup>+</sup> /Na <sup>+</sup> ratio in the leaf.
Brassica	Vacuolar Na <sup>+</sup> /H <sup>+</sup> antiporter AtNHX1	Arabidopsis thaliana L.	Increased proline content; Improved growth rate; Mitigate the toxic effect of Na <sup>+</sup> .
Arabidopsis	Plasma membrane Na <sup>+</sup> /H <sup>+</sup> antiporter SOS1	Arabidopsis thaliana L. (wild type)	Improved germination rate, root growth, and chlorophyll content; Reduced accumulation of Na <sup>+</sup> .
Maize	Vacuolar Na <sup>+</sup> /H <sup>+</sup> antiporter AtNHX1	Arabidopsis thaliana L.	Increased rate of germination.

### CONCLUSION:

At atomic, cell, metabolic, physiological, and entire plant levels, salinity resilience involves a complex of responses. Extensive research through cell, metabolic, and physiological analysis has explained that particle take-up, transport and equilibrium, osmotic guideline, chemical metabolism, cancer prevention agent metabolism, and stress signaling assume basic roles in plant transformation to salinity stress among various salinity responses, mechanisms or strategies. Taking preferred position of the latest advances in genomic, transcriptomic, proteomic, and metabolomic techniques, plant biologists are focusing on the improvement of a total quality, protein, and metabolite profile that is responsible for various salinity resilience mechanisms in various species of plants. There is, notwithstanding, an absence of information integration from genomic, transcriptomic, proteomic, and metabolomic studies, and a

combined methodology is essential for identifying key pathways or processes controlling salinity resistance.

## REFERENCES:

- Jacobsen, T., and Adams, R. M. (1958). Salt and silt in ancient Mesopotamian agriculture. *Science* 128, pp. 1251–1258. DOI: 10.1126/science.128.3334.1251
- Qadir, M., Quillerou, E., Nangia, V., Murtaza, G., Singh, M., Thomas, R. J., et. al. (2014). Economics of salt-induced land degradation and restoration. *Nat. Resour. Forum* 38, pp. 282–295. DOI: 10.1111/1477-8947. 12054
- M. De Lourdes Oliveira Otoch, A. C. Menezes Sobreira, M. E. Farias De Aragao, E. G. Orellano, M. Da Guia Silva Lima, and ~ D. Fernandes De Melo (2001). "Salt modulation of vacuolar H<sup>+</sup>-ATPase and H<sup>+</sup>-Pyrophosphatase activities in *Vigna unguiculata*," *Journal of Plant Physiology*, vol. 158, no. 5, pp. 545–551.
- B. Wang, U. Luttge, and R. Ratajczak (2001). "Effects of salt treatment " and osmotic stress on V-ATPase and V-PPase in leaves of the halophyte *Suaeda salsa*," *Journal of Experimental Botany*, vol. 52, no. 365, pp. 2355–2365.
- L. Ma, H. Zhang, L. Sun et. al. (2012). "NADPH oxidase Atrboh D. and Atrboh F. function in ROS-dependent regulation of Na<sup>+</sup>/K<sup>+</sup> homeostasis in *Arabidopsis* under salt stress," *Journal of Experimental Botany*, vol. 63, no. 1, pp. 305–317.
- J. Matysik, A. Alia, B. Bhalu, and P. Mohanty (2010). "Molecular mechanisms of quenching of reactive oxygen species by proline under stress in plants," *Current Science*, vol. 82, no. 5, pp. 525– 532.
- C. Ben Ahmed, B. Ben Rouina, S. Sensoy, M. Boukhriss, and F. Ben Abdullah (2010). "Exogenous proline effects on photosynthetic performance and antioxidant defense system of young olive tree," *Journal of Agricultural and Food Chemistry*, vol. 58, no. 7, pp. 4216–4222.
- M. A. Hoque, M. N. A. Banu, Y. Nakamura, Y. Shimoishi, and Y. Murata (2008). "Proline and glycinebetaine enhance antioxidant defense and methylglyoxal detoxification systems and reduce NaCl-induced damage in cultured tobacco cells," *Journal of Plant Physiology*, vol. 165, no. 8, pp. 813–824.
- S. Deivanai, R. Xavier, V. Vinod, K. Timalata, and O. F. Lim (2011). "Role of exogenous proline in ameliorating salt stress at early stage in two rice cultivars," *Journal of Stress Physiology & Biochemistry*, vol. 7, pp. 157–174.
- H. J. Bohnert, D. E. Nelson, and R. G. Jensen (1995). "Adaptations to environmental stresses," *Plant Cell*, vol. 7, no. 7, pp. 1099–1111.
- M. J. Van Oosten, A. Sharkhuu, G. Batelli, R. A. Bressan, and A. Maggio (2013). "The *Arabidopsis thaliana* mutant *air1* implicates SOS3 in the regulation of anthocyanins under salt stress," *Plant Molecular Biology*, vol. 83, pp. 405–415.
- T. Kusano, K. Yamaguchi, T. Berberich, and Y. Takahashi (2007). "The polyamine spermine rescues *Arabidopsis* from salinity and drought stresses (Plant Signaling and Behavior)," *Plant Signaling and Behavior*, vol. 2, no. 4, pp. 251–252.
- L. Zhao, F. Zhang, J. Guo, Y. Yang, B. Li, and L. Zhang (2004). "Nitric oxide functions as a signal in salt resistance in the calluses from two ecotypes of reed," *Plant Physiology*, vol. 134, no. 2, pp. 849– 857.
- A. M. Nalouisi, S. Ahmadiyan, A. Hatamzadeh, and M. Ghasemnezhad (2012). "Protective role of exogenous nitric oxide against oxidative stress induced by salt stress in bell-pepper (*Capsicum annum* L.)," *American-Eurasian Journal of Agricultural & Environmental Science*, vol. 12, no. 8, pp. 1085–1090.
- F. Zhang, Y. Wang, Y. Yang, H. Wu, D. Wang, and J. Liu (2007). "Involvement of hydrogen peroxide and nitric oxide in salt resistance in the calluses from *Populus euphratica*," *Plant, Cell and Environment*, vol. 30, no. 7, pp. 775–785.
- B. C. Keskin, A. T. Sarikaya, B. Yuksel, and A. R. Memon (2011). "Absciscic acid regulated gene expression in bread wheat (*Triticum aestivum* L.)," *Australian Journal of Crop Science*, vol. 4, no. 8, pp. 617–625.
- S.-Y. Song, Y. Chen, J. Chen, X.-Y. Dai, and W.-H. Zhang (2011). "Physiological mechanisms underlying OsNAC5-dependent tolerance of rice plants to

abiotic stress,” *Planta*, vol. 234, no. 2, pp. 331–345.

18. Y. Hu, L. Chen, H. Wang, L. Zhang, F. Wang, and D. Yu (2013). “Arabidopsis transcription factor WRKY8 functions antagonistically with its interacting partner VQ9 to modulate salinity stress tolerance,” *The Plant Journal*, vol. 74, pp. 730–745.
19. T. Serra, D. D. Figueiredo, A. M. Cordeiro et. al. (2013). “OsRMC, a negative regulator of salt stress response in rice, is regulated by two AP2/ERF transcription factors,” *Plant Molecular Biology*, vol. 82, pp. 439–455.
20. J. V. Cabello, A. F. Lodeyro, and M. D. Zurbriggen (2014). “Novel perspectives for the engineering of abiotic stress tolerance in plants,” *Current Opinion in Biotechnology*, vol. 26, pp. pp. 62–70.

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