The Na\textsuperscript{+}/H\textsuperscript{+} exchanger SOS1 controls extrusion and distribution of Na\textsuperscript{+} in tomato plants under salinity conditions

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Maintaining a high K\textsuperscript{+}/Na\textsuperscript{+} ratio in the cell cytosol, along with the transport processes implicated in the xylem and phloem loading/unloading of Na\textsuperscript{+} in plants (long-distance transport) are key aspects in plant salt tolerance. The Ca\textsuperscript{2+}-dependent SOS pathway regulating Na\textsuperscript{+} and K\textsuperscript{+} homeostasis and long-distance Na\textsuperscript{+} transport has been reported in Arabidopsis. However, Arabidopsis might not be the best model to analyze the involvement of the SOS pathway in long-distance Na\textsuperscript{+} transport due to the very short stem of these plants which do not allow a precise dissection of the relative content of Na\textsuperscript{+} in stem versus leaf. This separation would be critical to assess the role of SOS1 in xylem loading/unloading, Na\textsuperscript{+} export by roots, retention in stems and the differential distribution/accumulation in old leaves. Towards this goal, tomato might represent a superior model due to its anatomical structure and agricultural significance. We recently demonstrated the key role played by the plasma membrane Na\textsuperscript{+}/H\textsuperscript{+} antiporter SOS1 in salt tolerance in tomato by maintaining ion homeostasis under salinity stress and in the partitioning of Na\textsuperscript{+} in plant organs.

Ion homeostasis under salt stress conditions is essential in salt tolerance and involves a network of transport processes regulating uptake, extrusion through the plasma membrane, compartmentalization of salts into cell vacuoles and recirculation of ions through the plant organs, thus allowing the osmotic adjustment and maintenance of high K\textsuperscript{+}/Na\textsuperscript{+} ratios in the cytosol of plants.\textsuperscript{1,2} The SOS signaling pathway is essential in the salt tolerance of Arabidopsis.\textsuperscript{3} This pathway regulates Na\textsuperscript{+} and K\textsuperscript{+} homeostasis as well as long-distance Na\textsuperscript{+} transport from roots to shoots.\textsuperscript{3} In the SOS pathway, a calcium-binding protein, SOS3, senses cytosolic calcium changes elicited by salt stress.\textsuperscript{4} SOS3 physically interacts with and activates the serine/threonine protein kinase, SOS2.\textsuperscript{5} The SOS3/SOS2 kinase complex phosphorylates and activates the plasma membrane Na\textsuperscript{+}/H\textsuperscript{+} exchanger encoded by the SOS1 gene.\textsuperscript{6,7} AtSOS1 expression was observed at the epidermal cells of the root tip implying a role of this transporter in extruding Na\textsuperscript{+} to the soil.\textsuperscript{8} Additionally, the preferential expression of AtSOS1 in the cells surrounding the vasculature also suggested a role of this transporter in the control of long-distance Na\textsuperscript{+} transport in plants, since this ion is transported from the root to the shoot via the xylem.\textsuperscript{8} The great significance of this Na\textsuperscript{+}/H\textsuperscript{+} antiporter in salt tolerance was further recognized when overexpression of AtSOS1 restricted Na\textsuperscript{+} accumulation in plant cells and improved the salt tolerance of Arabidopsis.\textsuperscript{9}

Although orthologous SOS genes have been identified in crop species like rice and wheat,\textsuperscript{10,11} the relevance of this pathway for salt tolerance in crop plants has not yet been demonstrated. Furthermore, the study of the involvement of the SOS pathway in long-distance Na\textsuperscript{+} transport by retention of salt in the stem, and the differential distribution/accumulation in old leaves, is a difficult task in Arabidopsis due to the small size of the stem. Tomato is a good alternative to study Na\textsuperscript{+} transport, plasma membrane Na\textsuperscript{+}/H\textsuperscript{+} antiporter, salt tolerance, posttranscriptional gene silencing, SLOX1, Solanum lycopersicum (tomato)

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long-distance transport due to its anatomical structure, and the high genotypic diversity related to salt tolerance. The relatively more tolerant tomato species accumulate higher amounts of salt in stems and leaves while the more sensitive species accumulate salt mainly in roots. Physiological evidences indicate that tomato roots could determine the Na⁺ concentration reaching aerial parts depending on the intensity of the stress, probably involving the SOS pathway.

We isolated the SlSOS1 gene, encoding a Na⁺/H⁺ antiporter from tomato, and confirmed through gene silencing that it plays a crucial role in the survival of tomato plants under saline conditions. SlSOS1 silencing in tomato renders these plants more sensitive to salt stress (Fig. 1A), thus extending the observations from the model system Arabidopsis to a crop plant. Net uptake rates of Na⁺ in whole plants were three-fold higher in SlSOS1-silenced plants than in wild-type plants subjected to mild (25 mM NaCl) and severe salt stress (100 mM NaCl), indicating that the main action of SOS1 is the extrusion Na⁺ out of the root. As previously discussed, the rates of unidirectional entry of Na⁺ (influx) in roots are very high but they do not translate into a commensurate Na⁺ accumulation. The much lower rate of net Na⁺ uptake relative to unidirectional influx (about ten-fold) implies a substantial efflux rate of ca. 90% of the incoming Na⁺. Thus, even small changes in the rate of Na⁺ efflux, to which SOS1 is known to be critical, would have dramatic effects in the root Na⁺ content. However, what we would like to highlight here is the critical importance of SOS1 to control Na⁺ distribution among plant organs. The analyses of the fluxes of Na⁺ and K⁺ to individual organs supported an important function of this Na⁺/H⁺ antiporter in the distribution of Na⁺ throughout the tomato plant. Arabidopsis sos1 mutants accumulated more Na⁺ in shoots under severe salt stress than the wild-type, presumably because of uncontrolled Na⁺ uptake by roots. However, under mild stress conditions, the mutant accumulated less Na⁺ in the shoot than wild-type, suggesting that AtSOS1 mediated xylem loading.
In tomato, \textit{SISOS1}-silenced plants under saline conditions accumulated more Na\(^+\) in roots and leaves than control plants under both mild and severe regimes of salt stress (Fig. 1B). By contrast, the stems of suppressed plants accumulated significantly less Na\(^+\) under severe stress (100 mM NaCl). The ability to retain Na\(^+\) in stems was clearly lost in suppressed plants, in which an acropetal gradient of Na\(^+\) was established. As most glycophytes, tomato plants exclude salts from the photosynthetic tissues in leaves by removing Na\(^+\) from the root while retaining it mainly in the stem,\(^{15,16}\) and SOS1 seems critical to this process. The greatest Na\(^+\) accumulation in control plants was found in stems at any time point or salt regime, whereas the suppressed plants showed preferential deposition in roots, particularly at longer times of treatment. Average Na\(^+\) contents in stems of suppressed plants remained mostly unchanged. Net Na\(^+\) fluxes carried over by xylem sap movement was much lower in suppressed plants than in controls. The greater accumulation of Na\(^+\) in the roots of suppressed plants is coherent with the known role of SOS1 in Na\(^+\) efflux in roots, but the contribution of SOS1 to the retention of Na\(^+\) in the stem of the control plants is perplexing and not immediately reconciled with the proposal that SOS1 loads Na\(^+\) into the xylem. We hypothesize that the activity of SOS1 should be coordinated with other transporters in order to control the Na\(^+\) that reaches the photosynthetic tissues (Fig. 2).

A functional link and possible interplay between \textit{AtSOS1} and \textit{AtHKT1};1 has been previously suggested.\(^{17}\) It appears that the transport function of the SOS1 and HKT1 systems is coordinated to achieve Na\(^+\) (and K\(^+\)) homeostasis\(^{18}\) (Fig. 2). Dysfunction of either system alters long-distance transport and adequate partition of Na\(^+\), thereby resulting in salt-sensitive phenotypes in Arabidopsis.\(^{17}\) Our results are coherent with a model in which SOS1 mediates the transfer of Na\(^+\) from the xylem parenchyma to xylem vessels, preferentially in roots, whereas HKT1 would mediate the reverse flux, i.e., Na\(^+\) unloading from the xylem in the stem. The coordinated actions of these transporters would ultimately determine the amount of Na\(^+\) that leaves the roots, is retained in the stem, or transferred to leaves. It remains to be shown whether the depletion of SOS1 in suppressed tomato does affect the function of the HKT1 orthologous protein in the xylem parenchyma of tomato. At the same time, SOS1 might contribute in the exclusion of Na\(^+\) from the cytosol of young leaves toward the leaf apoplast, avoiding the accumulation of intracellular Na\(^+\) in cells lacking a well-developed vacuolar system, while HKT1 might act predominantly in the fully expanded or mature leaves with a well-developed vacuolar system that can accumulate Na\(^+\). Part of apoplastic Na\(^+\) could also be retrieved by the action of HKT1 at phloem companion cells. This might contribute, to some extent, to Na\(^+\) recirculation through phloem (Fig. 2). It will be of interest to experimentally determine the expression pattern of \textit{SOS1} (and perhaps \textit{HKTI}-like genes) in relation to the developmental stage of the plant organs, and also the possibility that these proteins are recruited to specific membrane domains near the xylem parenchyma-xylem vessel interface to mediate vectorial movements of Na\(^+\) in and out of the xylem sap.

Our results provide evidence of the importance of the plasma membrane Na\(^+\)/H\(^+\) antiporter, SOS1, in maintaining ion homeostasis in tomato, a crop plant that is a Na\(^+\) excluder with the capacity to tolerate moderate salt concentrations.\(^{12,15,16}\) We also show that, besides its main action in extruding Na\(^+\) out the root, \textit{SOS1} is critical for the partitioning of Na\(^+\) in plant organs and the ability of tomato plants to retain Na\(^+\) in the stems, thus preventing Na\(^+\) from reaching the photosynthetic tissues.

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**Figure 2.** Schematic diagram showing Na\(^+\) fluxes mediated by SISOS1 within the plant. The main action of SISOS1 is extruding Na\(^+\) out the root and the partitioning of Na\(^+\) in plant organs. Suppression of SISOS1 causes a root-to-leaf gradient of Na\(^+\) distribution together with a reduced ability to accumulate Na\(^+\) in the stem. The transport function of the SIS1 and HKT systems may be coordinated to achieve Na\(^+\) and K\(^+\) homeostasis.
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