

Review

Horticultural Salinity-Stress Phenotyping and Tolerance Inference: A Critical Evidence Map and Validation Framework for AI-Related Claims

Xiongwei Liang ^{1,2} , Shaopeng Yu ¹, Yongfu Ju ¹, Yingning Wang ^{1,2} and Dawei Yin ^{3,*} 

¹ Cold Region Wetland Ecology and Environment Research Key Laboratory of Heilongjiang Province, Harbin University, Harbin 150086, China; liangxiongwei007@163.com (X.L.); juyongfu@163.com (Y.J.)

² State Key Laboratory of Urban Water Resource and Environment, Harbin Institute of Technology, Harbin 150086, China

³ College of Agricultural Science, Heilongjiang Bayi Agricultural University, Daqing 163319, China

* Correspondence: yindazhiyindawei@126.com

Highlights

What are the main findings?

- Horticultural salinity-stress tolerance depends on crop-specific ion homeostasis, osmotic adjustment, ROS control, photosynthetic protection and production-system context.
- Only a small AI/sensing/phenotyping subset directly addresses salinity; current AI evidence is strongest as a phenotyping and validation layer, not as autonomous management proof.

What are the implications of the main findings?

- Validation ladders, leakage controls, biological metadata and reporting checklists should align AI claims with physiological, biochemical and omics evidence.
- Field-ready and commercially defensible claims require external validation, stress-confusion testing, economic feasibility assessment and prospective decision trials.

Abstract

Salinity stress constrains horticultural production in protected cultivation, hydroponics, coastal agriculture and reclaimed-water irrigation. This critical narrative review and evidence map synthesizes a DOI-verified core corpus of 160 peer-reviewed journal articles to ask how artificial intelligence (AI) can support, rather than overstate, salinity-stress inference in horticultural crops. The evidence base is uneven: 22 retained records were AI-, sensing- or phenotyping-relevant, six treated ML, deep learning, edge intelligence or agentic AI as a central method, and four directly tested salinity- or water-stress AI/sensor phenotyping in a crop-relevant system. Among the six explicit-AI records, none externally validated a salinity-specific AI model; one distinguished salinity from drought, and none reported a prospective AI-guided intervention trial. Accordingly, this article is framed as a validation and reporting framework, not as a quantitative meta-analysis of model or intervention efficacy. Across the broader corpus, evidence for salinity tolerance centres on osmotic limitation, Na⁺ and Cl[−] toxicity, K⁺ retention, ROS regulation, photosynthetic protection, hormonal signalling, root hydraulics, rhizosphere processes and metabolic re-programming. The review links these mechanisms to measurable traits and to claim-specific AI validation requirements. We propose a mechanism-to-AI map, a validation ladder, an intervention maturity framework and a reporting checklist. The conclusion is deliberately conservative: AI can improve salinity research when it is constrained by rigorous metadata,



Academic Editor: Dimitrios Fanourakis

Received: 30 June 2026

Revised: 22 July 2026

Accepted: 23 July 2026

Published: 25 July 2026

Copyright: © 2026 by the authors.

Licensee MDPI, Basel, Switzerland.

This article is an open access article distributed under the terms and

conditions of the [Creative Commons Attribution \(CC BY\)](https://creativecommons.org/licenses/by/4.0/) license.

physiological grounding, external validation, stress-confusion testing and explicit uncertainty, but current evidence is insufficient to support autonomous, economically validated or field-ready AI-based salinity management.

Keywords: artificial intelligence; salinity stress; horticultural crops; high-throughput phenotyping; machine learning; multi-omics; external validation; evidence synthesis; rootstock; biostimulants

1. Introduction

Soil salinization and irrigation-water salinization remain persistent constraints on high-value horticultural production, particularly in protected cultivation, hydroponic systems, grafted orchards, reclaimed-water irrigation and coastal or arid-zone fields. Existing reviews have summarized salinity physiology, high-throughput phenotyping, machine learning for plant-stress detection, multi-omics integration and agronomic innovation in greenhouse systems [1–12]. The unresolved question is narrower: which AI-related claims are justified by the current horticultural salinity evidence, and which remain measurement, validation or future-standard claims rather than management-ready conclusions?

Salt stress is sequential, organ-specific and production-system-dependent. Early osmotic limitation restricts water status, stomatal conductance, leaf expansion and root hydraulic adjustment; later ionic stress reflects Na^+ and Cl^- accumulation, K^+ competition, Ca^{2+} signalling disruption, membrane injury and oxidative imbalance. These phases interact with energy costs, developmental stage, cultivar, rootstock, substrate and greenhouse technology level, so a single composite salt-stress score is usually too coarse to support mechanistic interpretation [1,2,4,6,11,12].

The horticultural literature is broad but fragmented. Studies on tomato, cucumber, lettuce, citrus, grapevine, pepper, melon, watermelon, eggplant, strawberry and potato report growth suppression, ion imbalance, oxidative damage, reduced fluorescence, altered osmolyte profiles and partial rescue through grafting, microbial inoculants, biostimulants, nanoparticles or signal molecules. However, doses, substrates, cultivars, rootstock combinations, stress duration, developmental stage, sampled tissue and production targets differ widely enough that success in a single crop or greenhouse system rarely supports general salinity-management claims.

AI is relevant because this evidence base is heterogeneous, but heterogeneity alone does not make AI evidence mature. Machine learning, computer vision, hyperspectral sensing, chlorophyll fluorescence, UAV imagery, multi-omics integration and AI-assisted workflows can improve detection sensitivity, phenotyping scale, data linkage and auditability. They cannot, by architecture alone, transform treatment classification into mechanistic understanding, internal predictive accuracy into field usefulness, or single-system evidence into cross-crop transferability [7–10].

This review therefore addresses four explicit questions: what salinity mechanisms and crop contexts must anchor AI phenotypes; what portion of the current corpus provides direct AI/sensing evidence; what validation designs are required for detection, transfer, mechanism and decision claims; and what reporting standards would make future horticultural salinity AI studies reusable. The novelty is not another broad salinity-physiology review or a pooled model ranking, but a claim-calibration framework linking biological mechanism, evidence maturity and AI validation boundaries. Figure 1 presents a mechanism-to-AI framework linking salinity exposure and biological responses to measurable traits, sensing and AI inference, and claim-specific validation boundaries.

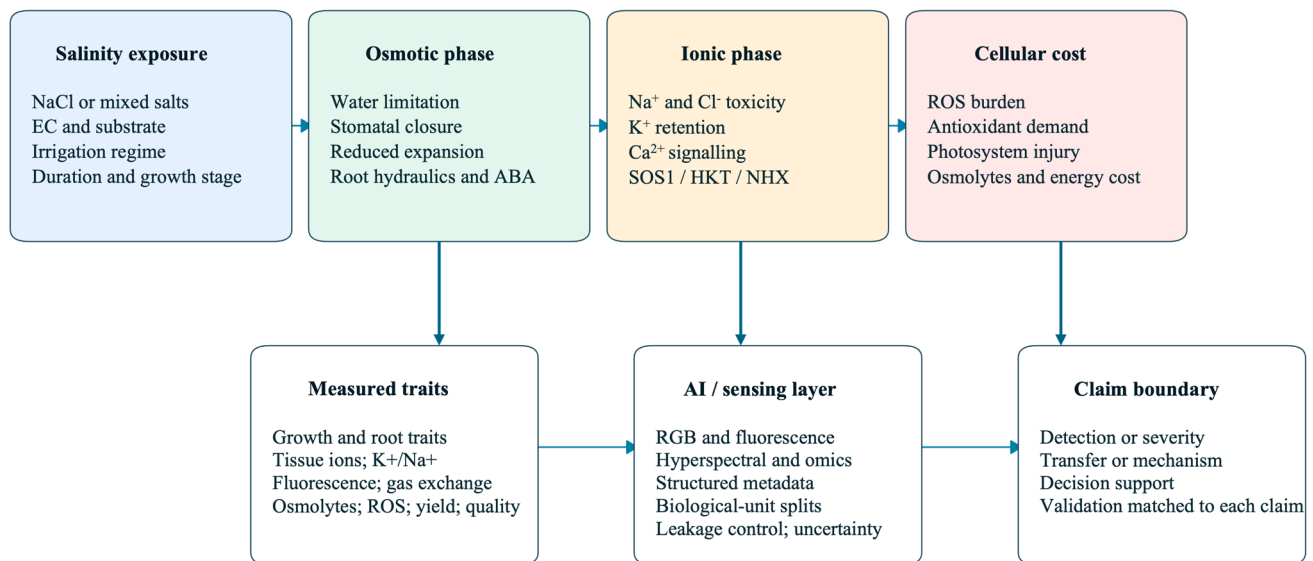


Figure 1. Mechanism-to-AI map for salinity-stress tolerance inference. This figure links osmotic limitation, Na^+/Cl^- toxicity, K^+ retention, ROS control, root hydraulics, stomatal regulation and energy cost to measurable phenotypes, sensor streams and AI validation boundaries.

2. Review Methodology and Evidence-Map Construction

This article uses a structured critical narrative review supported by an auditable evidence map. The design was chosen to calibrate claims across biologically heterogeneous studies, rather than to estimate pooled effects, study prevalence or comparative model performance. Quantitative pooling was avoided because the corpus varies sharply in crop, cultivar, rootstock, substrate, salt composition, dose, duration, endpoint, intervention timing, sampled organ, validation design and claim level. Pooling such studies would risk producing a false average across biologically non-equivalent experiments.

The structured database search was conducted in Web of Science, Scopus and PubMed up to 27 May 2026. Search blocks combined crop terms, salinity terms, physiological and biochemical response terms, multi-omics terms, sensing terms, machine-learning terms and AI-workflow terms. Google Scholar was used only after database retrieval for supplementary discovery and backward or forward citation chasing. Its ranked result set was neither treated as a bounded screening frame nor counted as a standalone source denominator. Any record first encountered through Google Scholar entered the same DOI/source verification and scope-anchoring procedure before consideration for the evidence map. Supplementary Information S1 reports the search date, 87 query expressions preserved in the source log, candidate-log categories, core-citation selection rationale, AI validation audit, study-level AI-subset evidence and operational definitions in Tables S1–S7. Database-specific syntax, searched fields, per-database hit counts, language restrictions and publication-type restrictions were not recorded contemporaneously and cannot be reconstructed without false precision. Supplementary Table S1 now reports each unavailable item explicitly, and Table S7 labels the 87 entries as preserved discovery phrases rather than database-executable search strings. Accordingly, this review does not claim PRISMA-level search reproducibility or exhaustive or bounded literature coverage.

To keep the 172-reference main bibliography manageable without oversized citation strings, the DOI-verified 160-paper core corpus is introduced once, in full, as an ordered citation-spine table. Table 1 is an audit device: it establishes first-appearance order for the core corpus, while later sections cite only the sources needed for a specific claim. Two additional contextual references were added during revision to address greenhouse species specificity and technology-context transferability [11,12].

Table 1. Ordered citation spine for the DOI-verified retained corpus. Contiguous reference ranges identify audit segments in the core corpus; later narrative sections use claim-specific citations. The supplementary study-level audit is provided in Supplementary Information S1, Tables S3–S6.

Corpus Segment	Evidence Use in Main Review	Detailed Coding
AI, phenotyping and validation-method support [13–34]	Defines sensor, imaging, ML, workflow and validation boundaries.	S1: AI subset and validation audit
Molecular, omics and mechanism evidence [35–64]	Links visible phenotypes to candidate regulatory, metabolic, genomic and rhizosphere mechanisms.	S1: omics and mechanism tags
Intervention evidence [65–96]	Maps rootstocks, grafting, biostimulants, microbial treatments, nanomaterials and signal molecules.	S1: intervention maturity coding
Crop-specific physiology and agronomy I [97–112]	Supports crop-specific response interpretation across lettuce, tomato, cucumber, grapevine and citrus.	S1: crop and endpoint tags
Crop-specific physiology and agronomy II [113–127]	Adds potato, grapevine, cucumber, melon, pepper and eggplant physiology or regulatory evidence.	S1: crop and endpoint tags
Crop-specific physiology and agronomy III [128–142]	Covers drought–salinity contrasts, rootstock studies, nanoparticles, hormones and cultivar variation.	S1: crop and intervention tags
Crop-specific physiology and agronomy IV [143–157]	Extends citrus, tomato, strawberry, eggplant, pepper, grapevine and microbial/field-context evidence.	S1: crop and mechanism tags
Crop-specific physiology and agronomy V [158–172]	Completes the retained corpus with compound stress, rootstock, genotype, salinity–drought and cultivar-response evidence.	S1: final corpus audit

The evidence map distinguishes direct from supporting evidence and avoids using supporting-method papers as proof of crop-specific salt tolerance. Direct evidence required a salinity, salt–alkali or closely related stress treatment applied to a horticultural crop, rootstock or crop-relevant system, with physiological, biochemical, ionomic, omics, microbial or agronomic endpoints. Supporting evidence comprised broader phenotyping, sensing, machine-learning and workflow papers that inform measurement or validation standards but do not, on their own, demonstrate crop-specific salinity tolerance.

The candidate log contained 542 unique DOI-bearing records. A curated core of 160 peer-reviewed journal articles was assembled to support the manuscript’s biological, intervention, sensing and validation claims. The remaining 382 records remained in the candidate log but were not selected for core citation because they were outside the target crop/stress scope, provided indirect methodological support beyond the review boundary, overlapped with a more directly relevant source for a peripheral illustrative role, or had source/type ambiguity. The 69 records in the overlap category met the broad topic scope but were not selected to duplicate a peripheral illustration already supported by a retained source. This was a core-citation selection decision, not a systematic-review eligibility exclusion or a scientific-quality judgement. The category cannot establish that retained and nonselected records are scientifically equivalent or that contradictory evidence is absent. The 160-paper corpus is therefore a curated citation set, not an eligible-study denominator. These disposition counts document only the article’s internal evidence-map

workflow and are not used to estimate field prevalence, study quality, intervention efficacy or model superiority. Of the 160 core records, 22 were assigned to the AI, phenotyping and validation-method segment. Figure 2 reports corpus construction and the associated audit trail.

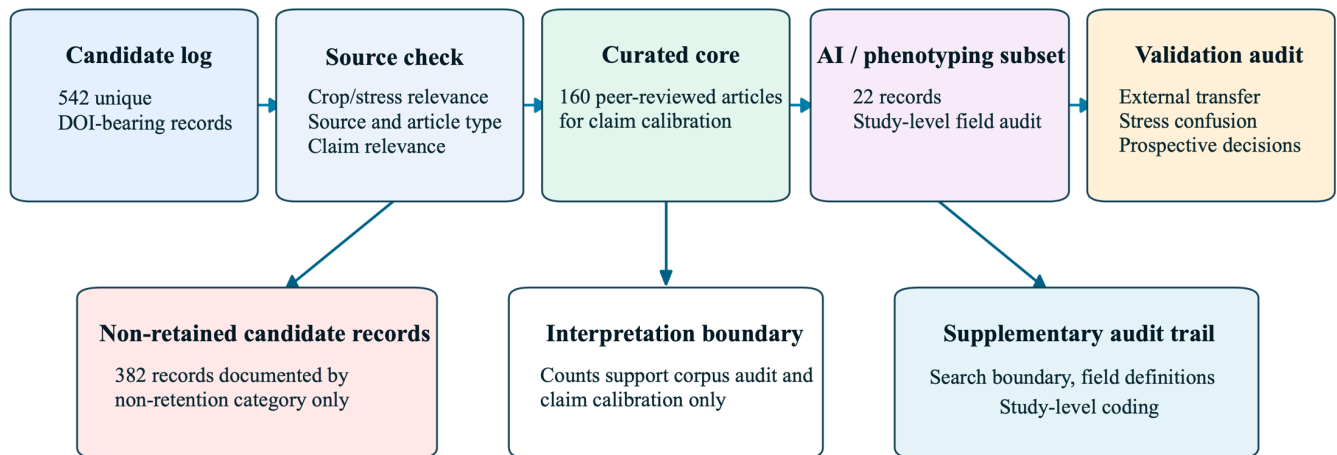


Figure 2. Evidence-map corpus construction and audit trail. The diagram documents how the DOI-bearing candidate log informed the curated core corpus and the AI/phenotyping validation audit. Counts describe the internal evidence-map workflow and are used only for claim calibration. The corresponding supplementary audit trail is provided in Supplementary Information S1 (Tables S1–S7).

Each retained study was tagged by crop group, stress composition where available, developmental context, experimental scale, evidence type, intervention class, data modality, validation design and claim level. Claim levels distinguished descriptive association, internal prediction, external validation, mechanistic triangulation and prospective decision testing. For the 22 AI-, sensing- or phenotyping-relevant records, Supplementary Information S1 provides study-level information in Table S4A–C, covering crop/system, data stream, model or sensor modality, prediction or phenotyping target, validation unit, external transfer, stress-confusion testing, prospective decision testing, performance metrics, data/code availability and claim level. For the remaining non-AI corpus, coding was limited to the biological and intervention evidence needed for narrative claim calibration, rather than full extraction of model-performance metrics.

Figure 3 summarizes the publication-year distribution, overlapping evidence tags, and main crop or methodological focus of the 160-paper core corpus.

The AI/phenotyping subset was audited separately because it carries the highest risk of evidence inflation. For denominator reconciliation, each of the 22 records was assigned one primary category according to the methodological role for which it is cited; secondary modalities remain listed in Supplementary Table S4A. The mutually exclusive primary categories are 14 image/fluorescence phenotyping records, five explicit ML/AI-method records, one remote/hyperspectral record, one agentic-workflow record and one non-AI physiological-support record. The 6/22 explicit ML/deep/agentic result is a nested analytical flag rather than a sixth category: it comprises the five explicit ML/AI-method records (refs. [21,24,27,29,34]) plus the agentic-workflow record (ref. [33]). Four records directly tested salinity- or water-stress AI/sensor phenotyping in a crop-relevant system; none of the six explicit-AI records externally validated a salinity-specific AI model; one record directly tested salt-versus-drought stress confusion; and none of the 22 records reported a prospective AI-guided intervention trial [13–34]. Table 2 therefore functions as a validation-risk audit, not as a ranking of model quality. These counts support a claim about evidence maturity, not model superiority.

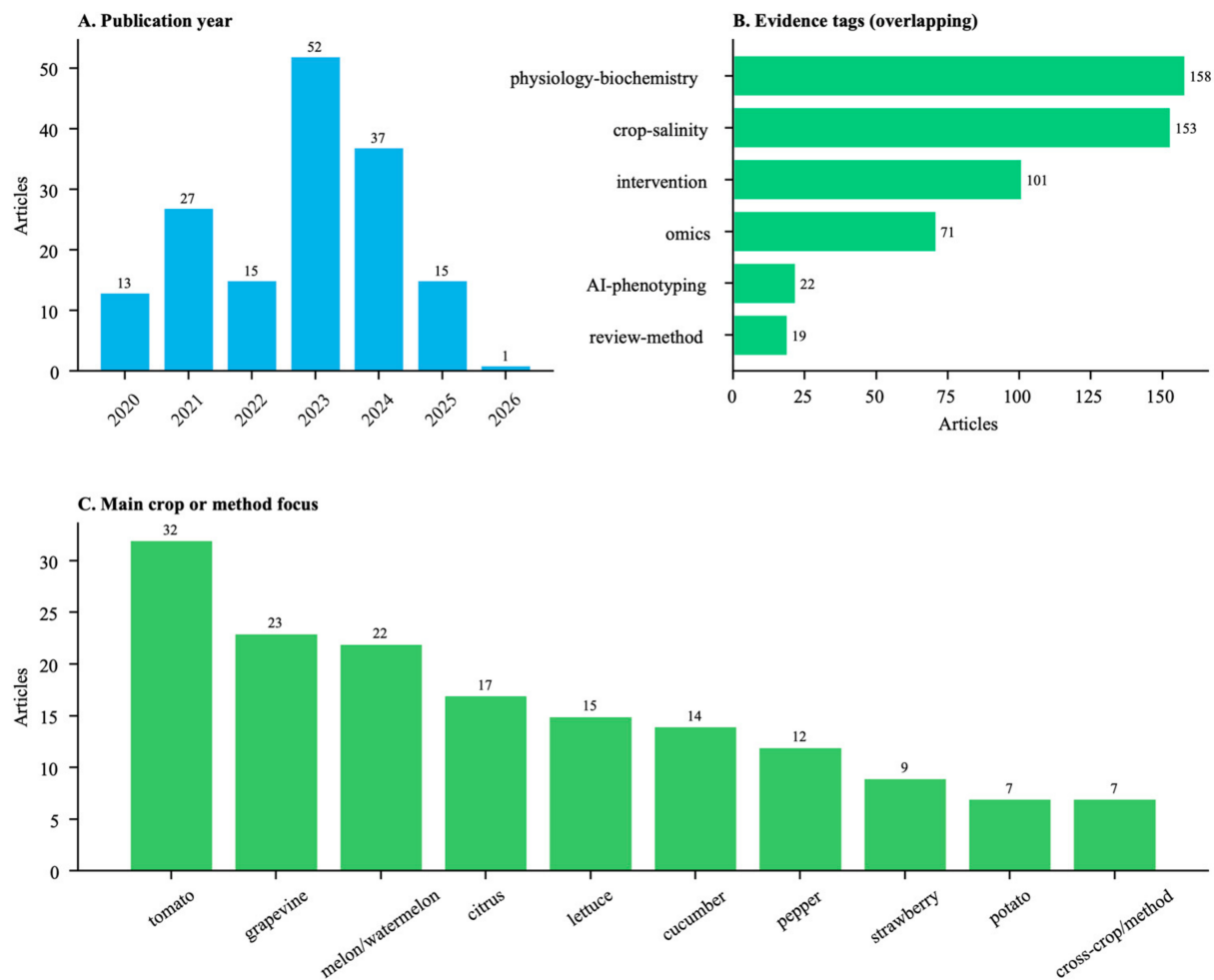


Figure 3. Evidence profile of the 160-paper core corpus. Evidence tags overlap because one article can contribute simultaneously to physiology, omics, intervention and AI-method evidence. Counts indicate evidence availability, not effect size or certainty.

Table 2. AI/phenotyping validation audit supporting the claim that decision-grade AI evidence remains scarce.

Audit Dimension	Operational Coding	Result	Interpretation	Qualifying/Reference Basis
AI/phenotyping-relevant subset	Core-corpus records in the AI, phenotyping and validation-method segment, including conventional non-AI physiological support used to define claim boundaries.	22/160	AI is important to the review question but is not the dominant direct evidence base.	Full subset in Supplementary Information S1, Tables S3 and S4A–S4C; representative sources: [19,21,24,29,33].
Explicit ML/deep/agentic AI	Machine learning, deep learning, edge intelligence, review-level AI analytics or multi-agent AI stated as central method or workflow.	6/22	Most AI-relevant records support measurement or workflow standards rather than full decision systems.	Qualifying records: [21,24,27,29,33,34].
Direct salinity AI/sensor phenotyping	Salinity- or water-stress treatment plus image, fluorescence, hyperspectral, sensor or ML phenotype endpoint in a crop or crop-relevant system; broad reviews excluded from the count.	4/22	Direct evidence exists but remains too sparse for broad management claims.	Qualifying examples: [19,21,24,31].

Table 2. Cont.

Audit Dimension	Operational Coding	Result	Interpretation	Qualifying/Reference Basis
External salinity-model validation	Test of a salinity-specific AI model on an independent dataset beyond the original experimental split and acquisition context.	0/6 explicit-AI records	No study met the strict external salinity-model definition; transfer and deployment claims remain unsupported.	No qualifying study identified.
Stress-confusion testing	Explicit salt-vs-drought or salt-vs-other-stress discrimination.	1/22	Most models do not yet show that salt stress is separated from visually or physiologically similar stresses.	Qualifying example: [21].
Prospective decision trial	AI-guided intervention tested prospectively against a management or experimental baseline.	0/22	No reviewed AI subset record supports autonomous intervention recommendation.	No eligible study identified.
Reusable benchmark metadata	Crop, genotype, stress composition, growth stage, imaging/sensor protocol and validation split sufficiently specified for transfer testing.	Heterogeneous	Benchmark-quality datasets and consistent metadata are needed before strong cross-study AI claims are credible.	Useful partial sources: [19,21,24,31,33].

3. Salinity Biology and Crop-Specific Evidence Signals

Salt stress produces interacting constraints rather than a single uniform response. The osmotic phase reduces water uptake, leaf expansion, stomatal conductance and carbon gain before large tissue Na^+ or Cl^- accumulation is necessarily detectable. The ionic phase reflects Na^+ and Cl^- accumulation, impaired K^+ retention, disrupted Ca^{2+} signalling, membrane leakage and progressive photosynthetic inhibition. Because these phases overlap in time and vary by organ, growth stage and crop architecture, mechanistic claims must remain crop-, organ- and stage-specific.

Most reviewed studies measured biomass, leaf area, root morphology, gas exchange, chlorophyll fluorescence, relative water content, membrane leakage, malondialdehyde, proline, soluble sugars, antioxidant enzymes, tissue ions and yield-quality traits [16,18,20,22,26,67,113,137,172]. These endpoints are informative but not interchangeable. Elevated proline may indicate osmotic adjustment, cellular injury or delayed development; antioxidant activity may indicate protection or a stronger oxidative challenge; and lower shoot Na^+ may reflect exclusion, vacuolar sequestration, altered transpiration or reduced uptake by smaller plants.

Distinguishing genuine tolerance from injury or escape therefore requires complementary measurements. A defensible interpretation should connect growth or image signals with at least some combination of tissue Na^+ and Cl^- , K^+/Na^+ ratio, osmotic potential, stomatal conductance, chlorophyll fluorescence, ROS or antioxidant markers, root hydraulic traits, yield or quality, and recovery after stress removal. Without these convergent measurements, AI models risk learning treatment labels rather than salinity physiology.

Crop-specific biology determines which proxy traits are worth modelling. In leafy vegetables, canopy expansion and marketable biomass sit close to the production target itself. In grapevine and citrus, rootstock-mediated Na^+ or Cl^- exclusion, perennial ion accumulation and scion–rootstock compatibility are central [14,54,55,75,129,143,171]. In fruiting vegetables, early fluorescence, root-trait or omics signals become useful only when linked to later yield, quality, recovery or greenhouse-management context [11,12].

4. AI as a Measurement and Evidence-Integration Layer

AI enters salinity research through several distinct routes that should not be pooled indiscriminately: image-based phenotyping, optical sensing, fluorescence and hyperspectral measurement, omics analytics, remote sensing and integrated decision-support workflows. RGB and root images, thermal data, fluorescence, hyperspectral reflectance, transcriptomes, metabolomes, ionomes and microbiome profiles all generate useful signals, but each supports a different level of inference [19,24,27–29,33,34].

The main methodological danger is endpoint substitution. Images and spectral indices can be acquired cheaply and repeatedly, whereas yield, fruit quality, postharvest performance, tissue ion content, and recovery are slower and more costly to measure. A model that detects early stress is valuable only when its claim is explicitly limited to detection, or when the proxy traits it relies on are explicitly linked to agronomic outcomes [19,21,24,31].

Each data stream carries its own claim boundary. RGB or root images can reveal differences in growth or morphology, but not ion homeostasis. Fluorescence and hyperspectral data can support early detection under a clearly stated protocol, but not robust diagnosis across all compound stresses. Omics models can prioritize candidate pathways, but cannot establish causal regulation without perturbation experiments, independent validation, or convergent physiological evidence [36,37,46,57,72].

Standard computer-vision assumptions require particular caution in plant-stress biology. Repeated images or temporal observations from the same biological unit can create direct train–test leakage, as the study-level designs in refs. [19,21,24] illustrate. In ref. [21], five-fold cross-validation was applied to 240 temporal observations derived from 48 pots, but pot-grouped partitioning was not reported. In ref. [24], a random 80/20 partition and repeated 10-fold cross-validation were reported, but grouping by recombinant inbred line and repeated time point was not specified. Their performance estimates are therefore treated here as internally exploratory and potentially optimistic, not as deployment evidence. Technical replicates, tray or chamber effects, acquisition date, lighting background, preprocessing before partitioning and full-dataset feature selection create additional acquisition-context leakage risks emphasized in the sensing and phenotyping literature [27–29]. A rigorous study should therefore split by plant or genotype for internal biological claims and by experiment, date, sensor, site or production system for the corresponding transfer claim; workflow papers such as ref. [33] are used here only to support auditable task separation, not salinity-model validation.

Label quality is often the weaker link, not model architecture. In many datasets, “salt stress” denotes treatment assignment rather than a validated physiological state. Stronger labels would combine treatment metadata with leaf Na^+ or Cl^- content, K^+/Na^+ ratio, fluorescence decline, gas exchange, osmotic potential, yield penalty, or recovery time [19,21,24,31,119,171].

Modelling should begin with the intended claim. Early detection requires earlier time points and independent plants, with repeated observations kept within one split [19]. Stress diagnosis requires explicit comparison with biologically similar non-salt stresses [21], whereas genotype ranking requires validation on unseen genotypes or trials [24]. Mechanism claims require alignment with ions, ROS, hormones or omics, and intervention recommendation requires prospective testing against a decision baseline. Model complexity is secondary to validation design; workflow evidence [33] is treated as methodological support rather than crop-specific validation. Figure 4 translates this claim hierarchy into a validation ladder.

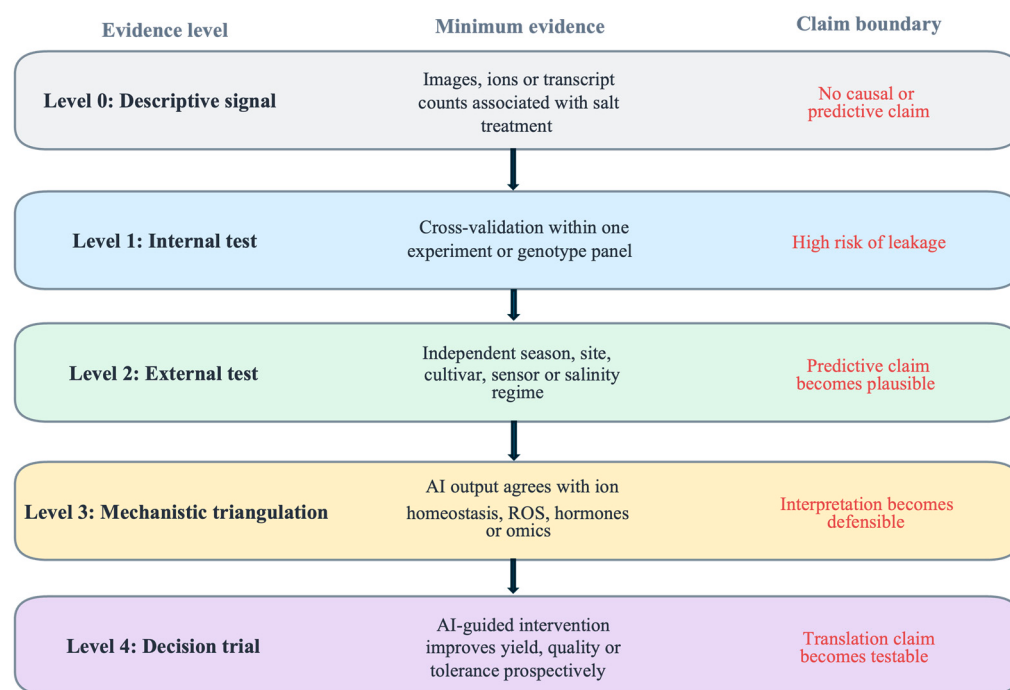


Figure 4. Validation ladder for AI claims in horticultural salinity research. Stronger biological and translational claims require external tests, mechanistic triangulation and prospective decision trials.

The validation ladder is a claim-control device. Level 0 and Level 1 studies remain useful when framed as descriptive or internal phenotyping evidence, as illustrated by the internal designs in refs. [21,24]. External-transfer, mechanistic and prospective decision levels are proposed requirements for stronger claims rather than observed properties of the current corpus. The goal is not to punish early studies but to prevent internal accuracy from being sold as deployment readiness.

5. Multi-Omics and Knowledge-Structured Inference

Multi-omics studies bridge visible phenotypes and cellular mechanisms by linking ion transport, osmoprotection, antioxidant regulation, photosynthesis, hormone pathways, cell-wall remodelling, aquaporins, transcription factors and rhizosphere processes [36–38,42,46–49,57,59,72,103,114,163]. They define the feature space that future AI systems should use, but they also expose the danger of naive feature selection.

The useful role for AI is not to discover tolerance *de novo* from sparse, high-dimensional matrices, but rather to organize candidate mechanisms, test whether features remain stable across genotypes or time points and render underlying assumptions visible. Omics-informed AI should rank pathways for confirmation, not replace physiological interpretation [34,36,37,46,57,62,72].

Small sample size relative to feature dimensionality remains a major risk. A transcriptome classifier may appear to separate treatment from control while having, in fact, learned batch structure, tissue age, or sampling time. Feature importance should therefore be checked across resampling schemes, normalization choices, genotypes, stress durations, and the biological direction of the effect [37,46,57,59,62,103,111,114].

Knowledge graphs and mechanism-constrained models are particularly promising because they encode prior relations among Na^+ exclusion, K^+ retention, ROS, ABA, ethylene, auxin, aquaporins, photosystem impairment, root architecture, and compatible solutes [34,36,46,57,72,91,147]. Even a simple graph linking crop, cultivar, dose, duration, tissue, intervention, response direction, and outcome quality would help surface contradictions rather than smoothing them away.

6. Intervention Evidence and Translational Maturity

The intervention literature is rich but uneven in maturity. Rootstocks and grafting offer the clearest translational logic because they are durable and already embedded in horticultural production, but they remain genotype-, scion-, site- and management-dependent [13,31,54,73,75,162]. Microbial inoculants are deployable but strongly conditioned by rhizosphere and substrate context [43,70,71,82]. Plant-derived biostimulants can improve selected endpoints, but responses vary with product, crop, cultivar and dose [41,84,88]. Nanomaterial studies report strong short-term responses, but the reviewed experiments provide limited evidence on dose transfer, environmental fate or regulatory readiness [44,53,74,85]. Signal molecules can also improve stress markers, but recovery dynamics and long-term performance remain incompletely tested [16,39,45,67,79].

AI should first triage existing evidence rather than recommend interventions autonomously. It can help identify where evidence for a given crop, stress type, endpoint, and intervention is dense enough to support synthesis, and where it remains confined to isolated case studies [27–29,33,34,50]. Figure 5 visualizes this uneven crop-by-intervention evidence density. Tomato, lettuce, citrus, grapevine, strawberry and melon do not have equal evidence maturity, so they should not receive equal translational confidence.

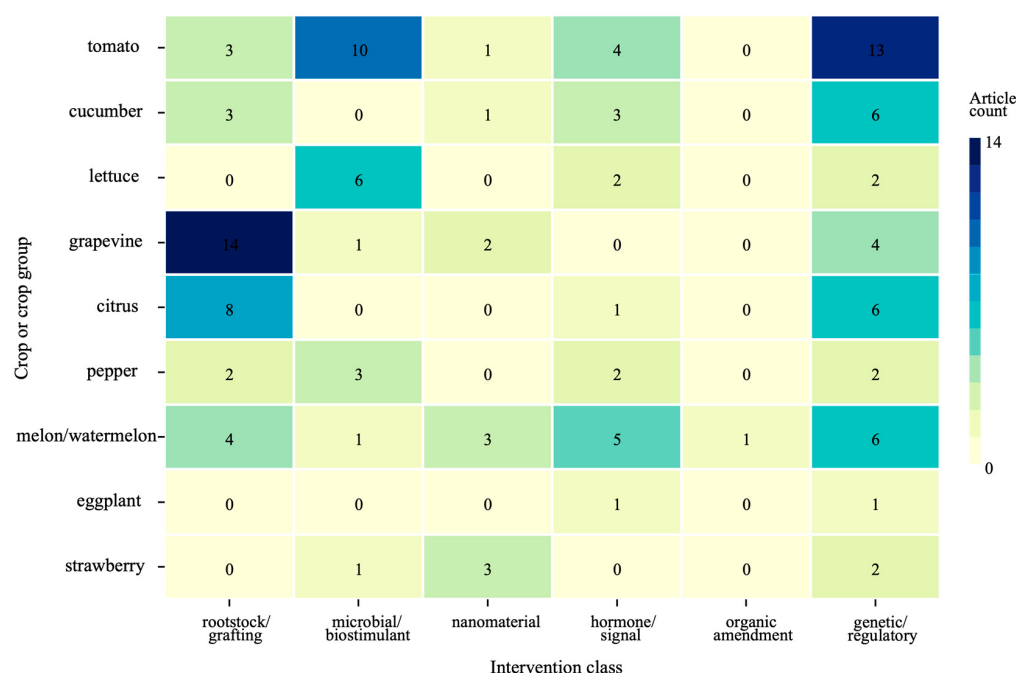


Figure 5. Crop-by-intervention evidence density in the reviewed literature. Counts indicate article co-occurrences in the evidence map; they do not indicate effect size, certainty or management readiness.

Rootstocks and grafting are not simply another input class: they can alter Na^+ and Cl^- transport, K^+/Na^+ balance, hydraulic function, root architecture, photosynthetic stability, and root-zone interactions. Their usefulness depends on scion–rootstock compatibility, stress duration, ion species, substrate, growth stage, and production target [55,83,86,95,96,131].

Biostimulants, microbial inoculants and nanomaterials introduce different sources of uncertainty. Protein hydrolysates and seaweed-derived biostimulants can improve selected physiological endpoints, but the evidence is often product-, cultivar-, dose- or duration-specific [41,84,88]. Microbial treatments, including *Bacillus* and mycorrhizal inoculants, show context-dependent benefits shaped by substrate and rhizosphere conditions [43,70,71,82]. Silicon- and selenium-based formulations modify antioxidant, ion-balance and photosynthetic responses, but transferability remains limited by short experi-

mental windows and incomplete dose assessment [74,81,138,140]. Metal-oxide nanoparticles, graphene oxide nanomaterials and hybrid lipid nanoparticles require separate toxicity and environmental-fate evaluation [44,53,85,92]. Biochar and related carbon amendments may alter yield and rhizosphere responses, but the evidence remains intervention- and production-system-specific [35,69].

Mechanistic maturity also differs among intervention classes. Rootstock effects should be interpreted through ion exclusion, vacuolar sequestration, K^+/Na^+ maintenance, aquaporin-mediated hydraulic conductance, hormonal signalling, photosynthetic stability and recovery after stress removal. Microbial and biostimulant effects require evidence for rhizosphere modification, nutrient uptake, oxidative regulation and persistence across substrates rather than short-term endpoint improvement alone.

For these interventions, AI is most useful when it models heterogeneity, failure cases, and dose–response relationships. A toxicity-aware response surface with explicit uncertainty boundaries is far more informative than a binary classifier that merely separates treated from untreated plants after the fact; negative results and cases of non-transfer are especially valuable for model calibration [27–29,33,34,50,53,138,140].

The next experimental step is response-surface design. AI can help select salinity doses, genotype–intervention combinations, scion–rootstock pairings and sampling times that avoid ceiling effects and reveal nonlinear windows. Table 3 translates the intervention evidence into maturity classes using five criteria: commercial embedding, mechanistic plausibility, durability across stress duration, transfer across genotype or production system, and unresolved safety or regulatory uncertainty. This would move the field beyond binary treatment–control studies toward designs that are more useful for growers and breeders [12,50,53,73,138,140,162].

Table 3. Translation maturity of major salinity-intervention classes and AI endpoints worth modelling.

Intervention Class	Translation Maturity	Strongest Evidence Signal	AI-Relevant Endpoint	Main Failure Mode
Rootstocks and grafting	Commercially embedded but genotype- and site-dependent.	Ion exclusion, hydraulic stability and durable scion–rootstock interactions [13,54,73,75,162].	Genotype-by-salinity response surfaces and yield–quality trade-offs.	Overgeneralizing from one scion–rootstock combination.
Microbial inoculants and biostimulants	Greenhouse promising; field establishment variable.	Root-zone modulation, antioxidant activation, nutrient uptake and partial yield rescue [36,43,70,77,84].	Treatment-response heterogeneity and failure-case prediction.	Substrate context and microbiome context determine transfer.
Nanomaterials	Mechanistically plausible but regulatory and dose–response maturity low.	Antioxidant defence, ion balance, photosynthetic protection and priming [44,53,74,85,138].	Toxicity-aware dose–response modelling.	Positive short-term physiology may not equal safe agronomic deployment.
Signal molecules and osmoprotectants	Experimental; timing- and concentration-sensitive.	ABA/ethylene/NO/H ₂ S/melatonin/salicylic signalling and osmolyte balance [16,26,67,79,115].	Temporal response trajectories and recovery prediction.	Biochemical marker increases may encode injury rather than tolerance.
Genetic and regulatory approaches	Discovery-stage to breeding-candidate level.	Transporters, transcription factors, aquaporins, polyploidy and gene-editing candidates [15,61,64,89,160].	Candidate prioritization across omics and phenotypes.	Gene-level evidence often stops before agronomic validation.

Transferability should be treated as a set of separate tests rather than a single claim. Evidence may transfer from one cultivar to another, from one scion–rootstock combination to another, across related species, across seasons or sites, across sensors, or from low-tech

to high-tech greenhouse systems. A model or intervention can pass one transfer boundary while failing another.

7. Reporting Standards, Failure Modes and Research Priorities

The central reporting rule is straightforward: every AI claim should be paired with the weakest condition under which it has been tested. Internal cross-validation supports internal discrimination, not field-level generalization [21,24]. In refs. [21,24], grouping of repeated temporal observations by pot or recombinant inbred line was not reported, so leakage through non-independent observations cannot be excluded. Repeated images or measurements from the same plant, pot or genotype must remain within one partition [19]. Training a model to distinguish salt-treated from control plants does not, by itself, support discrimination from drought, heat, nutrient deficiency, hypoxia, alkalinity or disease; among the audited subset, only ref. [21] explicitly compared salt with drought.

Reporting standards should specify the salinity protocol, plant age, developmental stage, cultivar, rootstock, sampled organ, tissue position, sampling time, number of independent plants, biological replication, technical replication, experimental unit, imaging or sensor protocol, preprocessing steps, data-split strategy, leakage controls, class balance, external-test definition, uncertainty estimates and code or model availability where possible. Table 4 turns these requirements into a minimum reporting checklist. Omics studies should add accession, tissue and time point, normalization, batch correction, multiple-testing procedure and validation criteria [36,37,46,57,72].

Table 4. Minimum reporting checklist for AI-enabled salinity studies in horticultural crops.

Checklist Item	Minimum Reporting Expectation	Why it Matters	Reviewer Decision if Absent
Stress protocol	NaCl or mixed-salt composition, EC, duration, timing, substrate and irrigation context.	Defines the biological state being modelled.	Downgrade all generalization claims.
Biological metadata	Cultivar, rootstock, growth stage, tissue, replication and environmental conditions.	Prevents hidden biological and environmental confounding.	Require clarification or reanalysis.
Data split and leakage controls	Plant-level or experiment-level separation; no image/time-point leakage; external test if claiming transfer.	Separates memorization from prediction.	Reject broad AI performance claims.
Mechanistic alignment	At least one convergent physiological, biochemical, ionic or omics layer for mechanism claims.	Prevents accuracy from being misread as understanding.	Restrict conclusion to detection.
Uncertainty and negative results	Confidence intervals, failure cases, stress confusions and boundary conditions.	Makes the model useful outside ideal conditions.	Treat as exploratory only.
Reproducibility	Code, model, raw/processed data or clear reason for restrictions.	Allows verification and reuse.	Flag as incomplete for decision-grade claims.

Data availability deserves particular attention. Study-level data or Supplementary Material was available in some audited records [19,21,24], but images, growth measurements, ion data, omics tables and environmental metadata were not consistently released as one integrated reusable dataset. Workflow and data-organization papers [33,34] support the need for structured provenance, but they are not counted as direct salinity-model evidence. Without standardized metadata, future AI systems risk reproducing narrow, study-specific validation patterns.

Auditable workflows can usefully separate planning, trait extraction, visualization, model checking and human review [33,34]. Figure 6 shows how this separation can be adapted to salinity phenotyping. Its value is not autonomous interpretation but structured accountability: requesting missing metadata, flagging leakage, comparing outputs against physiological priors and documenting uncertainty.

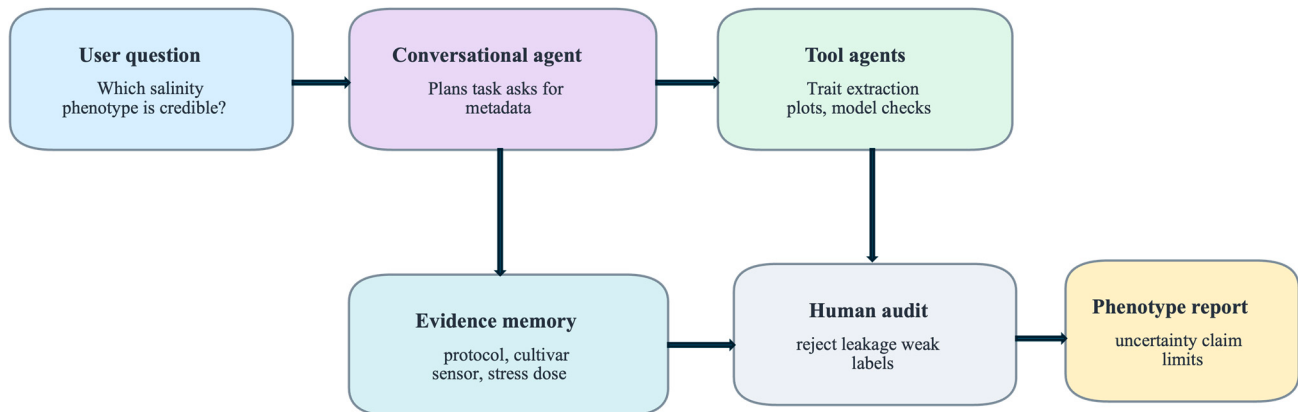


Figure 6. Proposed auditable AI workflow for salinity phenotyping. The workflow is redrawn as an original conceptual figure adapted from task-separated plant phenotyping architectures.

Commercial deployment requires evidence beyond biological detection and internal model accuracy. A deployable system must address sensor cost, camera or sensor mounting, calibration frequency, lighting variability, computational requirements, maintenance, data connectivity, compatibility with irrigation and fertigation infrastructure, grower usability, technical support and failure consequences.

False positives can trigger unnecessary leaching, fertigation changes or intervention costs, whereas false negatives can allow irreversible yield or quality losses. Deployment claims should therefore distinguish laboratory prototypes, advisory decision-support tools and autonomous commercial systems. None of the reviewed AI subset provides enough prospective, economic or regulatory evidence to support autonomous salinity management.

Six failure modes recur across the evidence map: data leakage, weak labels, stress confusion, intervention overgeneralization, field-transfer failure and premature commercialization. The AI validation risks are illustrated by the study-level evidence in refs. [19,21,24] and by the methodological workflow boundary in ref. [33]. Intervention overgeneralization and short experimental windows are separately supported by refs. [31,50,108,138,163,170]. These are not minor technical issues; they explain why internal accuracy or a short-term rescue effect cannot on its own be treated as decision-grade salinity evidence.

This review has six material limitations. First, the source log lacks database-specific syntax, searched fields, per-database hit counts, language and publication-type restrictions, and a bounded Google Scholar screening depth; therefore, this review cannot support a database-executable rerun, PRISMA-style search reproducibility or exhaustive literature-coverage claim. Second, evidence coding was conducted by one author, and no independent duplicate coding was performed. Third, treatment effects and model-performance estimates were not pooled, so model or intervention classes could not be ranked by average efficacy. Fourth, the 22-record AI, phenotyping and validation-method segment is small and methodologically heterogeneous. Fifth, none of the six explicit-AI records externally validated a salinity-specific AI model, and none of the 22 records conducted a prospective AI-guided decision trial. Sixth, the evidence cannot establish economic feasibility, field robustness, cultivar-to-cultivar transfer, cross-species applicability or autonomous salinity management.

Direct AI-specific salinity evidence remains a minority of the corpus. Many AI, sensing and workflow papers function as methodological support rather than direct evidence, so the frameworks proposed here should be read as standards for future evidence generation rather than proof that current systems have already reached decision-grade performance.

Initial evidence coding was performed by one author and then converted into the auditable operational categories in Supplementary Information S1. The coding supports claim calibration within this article and is not used for prevalence inference or pooled effect estimates. Nevertheless, source-verification notes, non-retention categories, preserved query expressions and the AI-subset matrix reduce, but do not eliminate, subjective judgement. Independent screening and evidence coding by a second author would be required before stronger prevalence or efficacy claims could be made.

The next phase of research should design studies around explicit claims: detection, severity estimation, genotype ranking, mechanism inference, intervention recommendation or commercial decision support. External tests should match the prediction claim and experimental unit, as the limitations of the internal designs in refs. [21,24] make clear; workflow evidence such as ref. [33] supplies audit structure rather than external validation. Mechanistic interpretation should be constrained by convergent physiological and omics evidence [36,37,57,72], while transfer across greenhouse technology levels requires explicit production-system context [12].

Factorial stress testing is urgently needed. Real horticultural systems combine salinity with heat, drought, high vapour-pressure deficit, nutrient imbalance, alkalinity, or pathogen pressure, and a single-factor NaCl model may fail precisely when saline irrigation coincides with heat or nutrient stress [21,31,32,98,108,138,159,163,170].

Experiments should also move toward dose–response and response-surface designs. AI can help select informative doses, prioritize sensors that predict later yield penalties, and identify genotype-by-intervention interactions—but only if failure cases and boundary conditions are consistently reported [50,53,73,138,140,162].

Cross-study learning requires shared metadata, including crop, cultivar, rootstock, growth stage, salt composition, electrical conductivity (EC), substrate, irrigation context, greenhouse technology level, sensor calibration, sampling time, endpoint definition and economic or management target. The audited sensing and modelling studies illustrate the need for stable experimental-unit and sensor metadata [19,21,24], and workflow evidence supports explicit provenance [33]. Crop, rootstock and production-system studies separately show why biological transfer metadata are necessary [11,12,54,119,171].

Review writing should be held to the same standard it asks of primary studies. Search terms, inclusion logic, evidence weighting, citation provenance, and figure licencing must all be transparent. The safest high-level claim is not that AI will solve salinity stress, but that it can improve how salinity evidence is measured, compared, and challenged.

8. Conclusions

The reviewed literature supports a cautious but constructive conclusion. Established biological evidence shows that horticultural salinity responses involve recurring processes, including osmotic limitation, ion homeostasis, K⁺ retention, ROS control, photosynthetic protection, hormone signalling, root architecture, root hydraulics, microbiome shifts and metabolic reprogramming. Their relative importance differs by crop, cultivar, developmental stage, organ, rootstock and production system.

The limited AI subset supports a narrower conclusion. Current evidence does not justify claims that AI can independently manage salinity stress. Most AI/phenotyping studies remain at the level of descriptive phenotyping, internal prediction or workflow support. Among the six explicit-AI records, none externally validated a salinity-specific AI

model; only one explicitly distinguished salinity from drought. No prospective AI-guided decision trial was identified, and the evidence is insufficient for economic claims.

The defensible path forward is hybrid and claim-limited. AI should function as an inference-support layer embedded within salinity biology, not as a replacement for physiological reasoning. Claims regarding early detection, phenotype stratification, evidence triage and experiment prioritization are defensible when metadata, validation split and biological endpoints are transparent. Causal, management, cross-species or autonomous-control claims based on internal accuracy alone should remain outside the conclusions that the current evidence can support.

Supplementary Materials: The following supporting information can be downloaded at: <https://www.mdpi.com/article/10.3390/horticulturae12080921/s1>. Supplementary Information S1: Table S1. Review design and search-source record. Unavailable fields are reported as not preserved and were not reconstructed retrospectively; Table S2. Candidate-log disposition and core-citation selection rationale. Categories describe evidence-map construction, not formal eligibility exclusion or scientific quality; Table S3. AI/phenotyping validation audit. Counts are based on the 22-record AI-, sensing- or phenotyping-relevant subset; Table S4A. Study identity, modality and evidence role. Review/method papers are not treated as original model validations; Table S4B. Biological design and validation strategy. Grouping omissions in refs. [21,24] are retained as validation-risk findings; Table S4C. Target, metrics, availability and claim boundary. External validation refers only to an independent salinity-specific model test beyond the original split and acquisition context; Table S5. Minimum reporting checklist for reusable salinity AI/phenotyping studies. These are proposed reporting requirements, not evidence that current systems are deployment-ready; Table S6. Operational field definitions and inference boundaries. Definitions were applied to prevent internal validation, repeated measurements and review papers from being upgraded to external model evidence; Table S7. Discovery phrases preserved in the original source log (search date: 27 May 2026). These are the 87 phrases exactly as recorded. They are not database-specific executable search strings; database syntax, searched fields, language restrictions, publication-type restrictions and per-database hit counts were not preserved.

Author Contributions: Conceptualization, X.L., D.Y. and S.Y.; Methodology, X.L.; Software, X.L.; Validation, X.L., S.Y., Y.W., D.Y. and Y.J.; Formal Analysis, X.L.; Investigation, X.L.; Resources, X.L.; Data Curation, X.L.; Writing—Original Draft Preparation, X.L.; Writing—Review and Editing, X.L., S.Y., Y.J., Y.W. and D.Y.; Visualization, X.L.; Supervision, S.Y. and D.Y.; Project Administration, S.Y.; Funding Acquisition, Y.J. All authors have read and agreed to the published version of the manuscript.

Funding: This article was funded by the Heilongjiang Provincial Natural Science Foundation of China, grant number PL2025D007.

Data Availability Statement: The original contributions presented in this study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

Acknowledgments: The authors used DeepSeek-v3 solely for English-language polishing. All AI-assisted edits were reviewed and revised by the authors, who take full responsibility for the content of this publication.

Conflicts of Interest: The authors declare no conflicts of interest.

Abbreviations

The following abbreviations are used in this manuscript:

ABA	Absciscic acid
AI	Artificial intelligence
CNN	Convolutional neural network
DEG	Differentially expressed gene

EC	Electrical conductivity
ML	Machine learning
NO	Nitric oxide
PGPB	Plant-growth-promoting bacteria
ROS	Reactive oxygen species
UAV	Unmanned aerial vehicle
XAI	Explainable artificial intelligence

References

- Munns, R.; Tester, M. Mechanisms of Salinity Tolerance. *Annu. Rev. Plant Biol.* **2008**, *59*, 651–681. [\[CrossRef\]](#)
- van Zelm, E.; Zhang, Y.; Testerink, C. Salt Tolerance Mechanisms of Plants. *Annu. Rev. Plant Biol.* **2020**, *71*, 403–433. [\[CrossRef\]](#) [\[PubMed\]](#)
- Zhu, J.K. Abiotic Stress Signaling and Responses in Plants. *Cell* **2016**, *167*, 313–324. [\[CrossRef\]](#) [\[PubMed\]](#)
- Yang, Y.; Guo, Y. Elucidating the molecular mechanisms mediating plant salt-stress responses. *New Phytol.* **2018**, *217*, 523–539. [\[CrossRef\]](#) [\[PubMed\]](#)
- Shabala, S. Learning from halophytes: Physiological basis and strategies to improve abiotic stress tolerance in crops. *Ann. Bot.* **2013**, *112*, 1209–1221. [\[CrossRef\]](#) [\[PubMed\]](#)
- Munns, R.; Day, D.A.; Fricke, W.; Watt, M.; Arsova, B.; Barkla, B.J.; Bose, J.; Byrt, C.S.; Chen, Z.-H.; Foster, K.J.; et al. Energy costs of salt tolerance in crop plants. *New Phytol.* **2020**, *225*, 1072–1090. [\[CrossRef\]](#) [\[PubMed\]](#)
- Araus, J.L.; Cairns, J.E. Field high-throughput phenotyping: The new crop breeding frontier. *Trends Plant Sci.* **2014**, *19*, 52–61. [\[CrossRef\]](#) [\[PubMed\]](#)
- Furbank, R.T.; Tester, M. Phenomics—Technologies to relieve the phenotyping bottleneck. *Trends Plant Sci.* **2011**, *16*, 635–644. [\[CrossRef\]](#) [\[PubMed\]](#)
- Singh, A.; Ganapathysubramanian, B.; Singh, A.K.; Sarkar, S. Machine Learning for High-Throughput Stress Phenotyping in Plants. *Trends Plant Sci.* **2016**, *21*, 110–124. [\[CrossRef\]](#) [\[PubMed\]](#)
- Ubbens, J.R.; Stavness, I. Deep Plant Phenomics: A Deep Learning Platform for Complex Plant Phenotyping Tasks. *Front. Plant Sci.* **2017**, *8*, 1190. [\[CrossRef\]](#) [\[PubMed\]](#)
- Fanourakis, D.; Makraki, T.; Vlachogiannakis, E.; Tsaniklidis, G.; Körner, O.; Ntatsi, G. Innovations in Agronomy and Their Impact on Greenhouse Vegetable Yields: Species-Specific Perspectives. *Horticulturae* **2026**, *12*, 684. [\[CrossRef\]](#)
- Fanourakis, D.; Tsaniklidis, G.; Makraki, T.; Ntanasi, T.; Giannothanas, E.; Papasotiropoulos, V.; Prohens, J.; Ntatsi, G. Adapting Breeding Goals to Greenhouse Technological Contexts for Sustainable Vegetable Production: A Review. *Hortic. Plant J.* **2026**, *in press*. [\[CrossRef\]](#)
- Mozafarian, M.; Hawrylak-Nowak, B.; Kappel, N. Effect of Different Rootstocks on the Salt Stress Tolerance and Fruit Quality of Grafted Eggplants (*Solanum melongena* L.). *Plants* **2023**, *12*, 3631. [\[CrossRef\]](#) [\[PubMed\]](#)
- Othman, Y.A.; Hani, M.B.; Ayad, J.Y.; St Hilaire, R. Salinity level influenced morpho-physiology and nutrient uptake of young citrus rootstocks. *Heliyon* **2023**, *9*, e13336. [\[CrossRef\]](#) [\[PubMed\]](#)
- Song, X.; Zhang, M.; Wang, T.T.; Duan, Y.Y.; Ren, J.; Gao, H.; Fan, Y.J.; Xia, Q.M.; Cao, H.X.; Xie, K.D.; et al. Polyploidization leads to salt stress resilience via ethylene signaling in citrus plants. *New Phytol.* **2025**, *246*, 176–191. [\[CrossRef\]](#) [\[PubMed\]](#)
- Yang, Y.; Yao, Y.; Li, J.; Zhang, J.; Zhang, X.; Hu, L.; Ding, D.; Bakpa, E.P.; Xie, J. Trehalose Alleviated Salt Stress in Tomato by Regulating ROS Metabolism, Photosynthesis, Osmolyte Synthesis, and Trehalose Metabolic Pathways. *Front. Plant Sci.* **2022**, *13*, 772948. [\[CrossRef\]](#) [\[PubMed\]](#)
- Saeedi, R.; Seyedi, A.; Esmaeilzadeh, M.; Seyedi, N.; Morteza Zahedi, S.; Malekzadeh, M.R. Improving the performance of the photosynthetic apparatus of *Citrus sinensis* with the use of chitosan-selenium nanocomposite (CS + Se NPs) under salinity stress. *BMC Plant Biol.* **2024**, *24*, 745. [\[CrossRef\]](#) [\[PubMed\]](#)
- Shin, Y.K.; Bhandari, S.R.; Jo, J.S.; Song, J.W.; Cho, M.C.; Yang, E.Y.; Lee, J.G. Response to Salt Stress in Lettuce: Changes in Chlorophyll Fluorescence Parameters, Phytochemical Contents, and Antioxidant Activities. *Agronomy* **2020**, *10*, 1627. [\[CrossRef\]](#)
- Montanaro, G.; Briglia, N.; Petrozza, A.; Carlomagno, A.; Rustioni, L.; Cellini, F.; Nuzzo, V. Image-based sensing of salt stress in grapevine. *OENO One* **2024**, *58*, 7757. [\[CrossRef\]](#)
- Avestan, S.; Ghasemnezhad, M.; Esfahani, M.; Barker, A.V. Effects of nanosilicon dioxide on leaf anatomy, chlorophyll fluorescence, and mineral element composition of strawberry under salinity stress. *J. Plant Nutr.* **2021**, *44*, 3005–3019. [\[CrossRef\]](#)
- Mohammadi, P.; Asefpour Vakilian, K. Machine learning provides specific detection of salt and drought stresses in cucumber based on miRNA characteristics. *Plant Methods* **2023**, *19*, 123. [\[CrossRef\]](#) [\[PubMed\]](#)

22. Fátima, R.T.; Dantas, M.V.; Lima, G.S.; Oliveira, V.K.N.; Soares, L.A.A.; Silva, A.A.R.; Gheyi, H.R.; Guedes, M.A.; Nóbrega, J.S.; Fernandes, P.D. Salicylic acid does not relieve salt stress on gas exchange, chlorophyll fluorescence, and hydroponic melon growth. *Braz. J. Biol.* **2023**, *83*, e274595. [[CrossRef](#)] [[PubMed](#)]
23. AlNeyadi, S.; Kappachery, S.; Khan, T.A.; Karumannil, S.; AlHosani, M.; Anand Gururani, M. Ectopic expression of potato *ARP1* encoding auxin-repressed protein confers salinity stress tolerance in *Arabidopsis thaliana*. *PLoS ONE* **2024**, *19*, e0309452. [[CrossRef](#)] [[PubMed](#)]
24. Kumar, P.; Eriksen, R.L.; Simko, I.; Mou, B. Molecular Mapping of Water-Stress Responsive Genomic Loci in Lettuce (*Lactuca* spp.) Using Kinetics Chlorophyll Fluorescence, Hyperspectral Imaging and Machine Learning. *Front. Genet.* **2021**, *12*, 634554. [[CrossRef](#)] [[PubMed](#)]
25. Gohari, G.; Panahirad, S.; Sadeghi, M.; Akbari, A.; Zareei, E.; Zahedi, S.M.; Bahrami, M.K.; Fotopoulos, V. Putrescine-functionalized carbon quantum dot (put-CQD) nanoparticles effectively prime grapevine (*Vitis vinifera* cv. 'Sultana') against salt stress. *BMC Plant Biol.* **2021**, *21*, 120. [[CrossRef](#)] [[PubMed](#)]
26. Wei, L.; Zhang, J.; Wei, S.; Hu, D.; Liu, Y.; Feng, L.; Li, C.; Qi, N.; Wang, C.; Liao, W. Nitric Oxide Enhanced Salt Stress Tolerance in Tomato Seedlings, Involving Phytohormone Equilibrium and Photosynthesis. *Int. J. Mol. Sci.* **2022**, *23*, 4539. [[CrossRef](#)] [[PubMed](#)]
27. Liu, J.; Xiang, J.; Jin, Y.; Liu, R.; Yan, J.; Wang, L. Boost Precision Agriculture with Unmanned Aerial Vehicle Remote Sensing and Edge Intelligence: A Survey. *Remote Sens.* **2021**, *13*, 4387. [[CrossRef](#)]
28. Omia, E.; Bae, H.; Park, E.; Kim, M.S.; Baek, I.; Kabenge, I.; Cho, B.K. Remote Sensing in Field Crop Monitoring: A Comprehensive Review of Sensor Systems, Data Analyses and Recent Advances. *Remote Sens.* **2023**, *15*, 354. [[CrossRef](#)]
29. Al-Tamimi, N.; Langan, P.; Bernád, V.; Walsh, J.; Mangina, E.; Negrão, S. Capturing crop adaptation to abiotic stress using image-based technologies. *Open Biol.* **2022**, *12*, 210353. [[CrossRef](#)] [[PubMed](#)]
30. Ahmed, W.; Imran, M.; Yaseen, M.; Haq, T.U.; Jamshaid, M.U.; Rukh, S.; Ikram, R.M.; Ali, M.; Ali, A.; Maqbool, M.; et al. Role of salicylic acid in regulating ethylene and physiological characteristics for alleviating salinity stress on germination, growth and yield of sweet pepper. *PeerJ* **2020**, *8*, e8475. [[CrossRef](#)] [[PubMed](#)]
31. Dunlevy, J.D.; Blackmore, D.H.; Betts, A.; Jewell, N.; Brien, C.; Berger, B.; Walker, R.R.; Edwards, E.J.; Walker, A.R. Investigating the effects of elevated temperature on salinity tolerance traits in grapevine rootstocks using high-throughput phenotyping. *Aust. J. Grape Wine Res.* **2022**, *28*, 276–291. [[CrossRef](#)]
32. Gao, Y.; Li, X.; Han, C.; Huang, Q.; Wu, R.; Zhao, C.; She, K. Photosystem vulnerabilities under compound abiotic stresses: Mechanisms, diagnostics, and engineering for resilient crops. *Plant Stress* **2026**, *19*, 101193. [[CrossRef](#)]
33. Chen, F.; Stogiannidis, I.; Wood, A.; Bueno, D.; Williams, D.; Macfarlane, F.; Grieve, B.D.; Wells, D.; Atkinson, J.A.; Hawkesford, M.J.; et al. A conversational multi-agent AI system for automated plant phenotyping. *Nat. Commun.* **2026**, *17*, 6391. [[CrossRef](#)] [[PubMed](#)]
34. Zhao, L.; Walkowiak, S.; Fernando, W.G.D. Artificial Intelligence: A Promising Tool in Exploring the Phytomicrobiome in Managing Disease and Promoting Plant Health. *Plants* **2023**, *12*, 1852. [[CrossRef](#)] [[PubMed](#)]
35. Sahin, O.; Gunes, A.; Yagcioglu, K.D.; Kadioglu, Y.K. Mitigating Combined Boron and Salt Stress in Lettuce (*Lactuca sativa* L. Semental) through Salicylic Acid-Modified Rice Husk Biochar. *J. Soil Sci. Plant Nutr.* **2024**, *24*, 5220–5234. [[CrossRef](#)]
36. Monterisi, S.; Zhang, L.; Garcia-Perez, P.; Alzate Zuluaga, M.Y.; Ciriello, M.; El-Nakhel, C.; Buffagni, V.; Cardarelli, M.; Colla, G.; Roupahel, Y.; et al. Integrated multi-omic approach reveals the effect of a Gramineae-derived biostimulant and its lighter fraction on salt-stressed lettuce plants. *Sci. Rep.* **2024**, *14*, 10710. [[CrossRef](#)] [[PubMed](#)]
37. Gan, J.; Qiu, Y.; Tao, Y.; Zhang, L.; Okita, T.W.; Yan, Y.; Tian, L. RNA-seq analysis reveals transcriptome reprogramming and alternative splicing during early response to salt stress in tomato root. *Front. Plant Sci.* **2024**, *15*, 1394223. [[CrossRef](#)] [[PubMed](#)]
38. Ikuyinminu, E.; Goñi, O.; Łangowski, Ł.; O'Connell, S. Transcriptome, Biochemical and Phenotypic Analysis of the Effects of a Precision Engineered Biostimulant for Inducing Salinity Stress Tolerance in Tomato. *Int. J. Mol. Sci.* **2023**, *24*, 6988. [[CrossRef](#)] [[PubMed](#)]
39. Yan, M.; Mao, J.; Wu, T.; Xiong, T.; Huang, Q.; Wu, H.; Hu, G. Transcriptomic Analysis of Salicylic Acid Promoting Seed Germination of Melon under Salt Stress. *Horticulturae* **2023**, *9*, 375. [[CrossRef](#)]
40. Yang, W.; Ling, Y.; Li, M.; Zhang, X.; Liu, B. Screening and Identification of Saline-Tolerant Germplasm in Melon. *Agriculture* **2023**, *13*, 2051. [[CrossRef](#)]
41. Zhang, L.; Freschi, G.; Roupahel, Y.; De Pascale, S.; Lucini, L. The differential modulation of secondary metabolism induced by a protein hydrolysate and a seaweed extract in tomato plants under salinity. *Front. Plant Sci.* **2023**, *13*, 1072782. [[CrossRef](#)] [[PubMed](#)]
42. Miao, C.; Zhang, Y.; Cui, J.; Zhang, H.; Wang, H.; Jin, H.; Lu, P.; He, L.; Zhou, Q.; Yu, J.; et al. An Enhanced Interaction of Graft and Exogenous SA on Photosynthesis, Phytohormone, and Transcriptome Analysis in Tomato under Salinity Stress. *Int. J. Mol. Sci.* **2024**, *25*, 10799. [[CrossRef](#)] [[PubMed](#)]

43. Benito, P.; Celdrán, M.; Bellón, J.; Arbona, V.; González-Guzmán, M.; Porcel, R.; Yenush, L.; Mulet, J.M. The combination of a microbial and a non-microbial biostimulant increases yield in lettuce (*Lactuca sativa*) under salt stress conditions by up-regulating cytokinin biosynthesis. *J. Integr. Plant Biol.* **2024**, *66*, 2140–2157. [\[CrossRef\]](#) [\[PubMed\]](#)
44. Guerriero, G.; Sutura, F.M.; Hoffmann, J.; Leclercq, C.C.; Planchon, S.; Berni, R.; Hausman, J.-F.; Renaut, J.; Torabi-Pour, N.; Pennington, H.C.; et al. Nanoporous Quercetin-Loaded Silicon-Stabilized Hybrid Lipid Nanoparticles Alleviate Salt Stress in Tomato Plants. *ACS Appl. Nano Mater.* **2023**, *6*, 3647–3660. [\[CrossRef\]](#)
45. Liu, J.; Li, J.; Li, X.; Song, Y.; Zhang, Z.; Sun, J.; Sun, X. Melatonin-induced transcriptome variation of melon seedlings under salt stress. *Mater. Express* **2023**, *13*, 495–507. [\[CrossRef\]](#)
46. Kashyap, S.P.; Prasanna, H.C.; Kumari, N.; Mishra, P.; Singh, B. Understanding salt tolerance mechanism using transcriptome profiling and de novo assembly of wild tomato *Solanum chilense*. *Sci. Rep.* **2020**, *10*, 15835. [\[CrossRef\]](#) [\[PubMed\]](#)
47. Mahmoud, L.M.; Killiny, N.; Holden, P.; Gmitter, F.G.; Grosser, J.W.; Dutt, M. Physiological and Biochemical Evaluation of Salt Stress Tolerance in a Citrus Tetraploid Somatic Hybrid. *Horticulturae* **2023**, *9*, 1215. [\[CrossRef\]](#)
48. Wang, B.; Wang, X.; Wang, Z.; Zhu, K.; Wu, W. Comparative metagenomic analysis reveals rhizosphere microbial community composition and functions help protect grapevines against salt stress. *Front. Microbiol.* **2023**, *14*, 1102547. [\[CrossRef\]](#) [\[PubMed\]](#)
49. Wang, Y.; Zhou, J.; Wen, W.; Sun, J.; Shu, S.; Guo, S. Transcriptome and Proteome Analysis Identifies Salt Stress Response Genes in Bottle Gourd Rootstock-Grafted Watermelon Seedlings. *Agronomy* **2023**, *13*, 618. [\[CrossRef\]](#)
50. Melino, V.; Tester, M. Salt-Tolerant Crops: Time to Deliver. *Annu. Rev. Plant Biol.* **2023**, *74*, 671–696. [\[CrossRef\]](#) [\[PubMed\]](#)
51. Fu, L.; Tian, X.; Wang, W.; Wu, C. Metabolome and Transcriptome Analysis Reveals Molecular Mechanisms of Soil Amendment (Volcanic Ash) Alleviating Salt-Alkali Stress in Melons (*Cucumis melo* L.). *Agronomy* **2024**, *14*, 2478. [\[CrossRef\]](#)
52. Attia, M.S.; Osman, M.S.; Mohamed, A.S.; Mahgoub, H.A.; Garada, M.O.; Abdelmouty, E.S.; Abdel Latef, A.A.H. Impact of Foliar Application of Chitosan Dissolved in Different Organic Acids on Isozymes, Protein Patterns and Physio-Biochemical Characteristics of Tomato Grown under Salinity Stress. *Plants* **2021**, *10*, 388. [\[CrossRef\]](#) [\[PubMed\]](#)
53. Albaladejo-Marico, L.; Thameur, A.; Garcia-Martinez, A.; Carvajal, M.; Yepes-Molina, L. Nanoparticles as modulators of stress tolerance: Physiological and molecular insights into TiO₂ and ZnO effects in *Cucumis melo* L. subjected to salt shock. *Environ. Technol. Innov.* **2025**, *38*, 104101. [\[CrossRef\]](#)
54. Modica, G.; Di Guardo, M.; Puglisi, I.; Baglieri, A.; Fortuna, S.; Arcidiacono, F.; Costantino, D.; La Malfa, S.; Gentile, A.; Arbona, V.; et al. Novel and widely spread citrus rootstocks behavior in response to salt stress. *Environ. Exp. Bot.* **2024**, *225*, 105835. [\[CrossRef\]](#)
55. Zhou-Tsang, A.; Wu, Y.; Henderson, S.W.; Walker, A.R.; Borneman, A.R.; Walker, R.R.; Gilliam, M. Grapevine salt tolerance. *Aust. J. Grape Wine Res.* **2021**, *27*, 149–168. [\[CrossRef\]](#)
56. Sun, H.; Wang, Y.; Cao, L.; Wang, Y.; Wei, Z.; Song, L.; Jiang, J.; Liu, J.; Tian, S. Transcriptome profiling reveals key genes in eggplant (*Solanum melongena*) roots under salt stress. *BMC Genom.* **2025**, *26*, 635. [\[CrossRef\]](#) [\[PubMed\]](#)
57. Wang, B.; Wang, J.; Yang, T.; Wang, J.; Dai, Q.; Zhang, F.; Xi, R.; Yu, Q.; Li, N. The transcriptional regulatory network of hormones and genes under salt stress in tomato plants (*Solanum lycopersicum* L.). *Front. Plant Sci.* **2023**, *14*, 1115593. [\[CrossRef\]](#) [\[PubMed\]](#)
58. Zhang, L.; Wang, M.; Tang, X.; Yang, X.; Zhang, Z.; Wu, J. Genome-Wide Identification of β -Ketoacyl CoA Synthase Gene Family in Melon (*Cucumis melo* L.) and Its Expression Analysis in Autotoxicity, Saline-Alkali, and Microplastic Exposure Environments. *Curr. Issues Mol. Biol.* **2025**, *47*, 195. [\[CrossRef\]](#) [\[PubMed\]](#)
59. Liu, P.; Gao, C.; Li, S.; Wang, X.; Dong, Y.; Wang, C.; Jiao, Z.; Sun, J. Comparative Transcriptome Analysis of Gene Responses of Salt-Tolerant and Salt-Sensitive Watermelon Cultivars' Roots to Salt Stress. *Plants* **2025**, *14*, 1013. [\[CrossRef\]](#) [\[PubMed\]](#)
60. Vives-Peris, V.; López-Climent, M.F.; Moliner-Sabater, M.; Gómez-Cadenas, A.; Pérez-Clemente, R.M. Morphological, physiological, and molecular scion traits are determinant for salt-stress tolerance of grafted citrus plants. *Front. Plant Sci.* **2023**, *14*, 1145625. [\[CrossRef\]](#) [\[PubMed\]](#)
61. Zhang, A.; Shang, J.; Xiao, K.; Zhang, M.; Wang, S.; Zhu, W.; Wu, X.; Zha, D. WRKY transcription factor 40 from eggplant (*Solanum melongena* L.) regulates ABA and salt stress responses. *Sci. Rep.* **2024**, *14*, 19289. [\[CrossRef\]](#) [\[PubMed\]](#)
62. Wu, X.; Yuan, D.; Chen, L.; Sun, Y.; Wang, Y.; Jiang, S.; Sun, X.; Zhang, X.; Li, J.; Gao, H. Comparative transcriptome analysis revealed that tomato response to salt stress induced by exogenous GABA and functional verification of *SIGAD1*. *BMC Plant Biol.* **2025**, *25*, 1788. [\[CrossRef\]](#) [\[PubMed\]](#)
63. Liu, K.; Li, X.; Wang, C.; Han, Y.; Zhu, Z.; Li, B. Genome-wide identification and characterization of the LRX gene family in grapevine (*Vitis vinifera* L.) and functional characterization of *VvLRX7* in plant salt response. *BMC Genom.* **2024**, *25*, 1155. [\[CrossRef\]](#) [\[PubMed\]](#)
64. Li, S.; Jiang, Y.; Xie, H.; Wang, K.; Yang, K.; Cao, Q.; Xue, H. *FvWRKY75* Positively Regulates *FvCRK5* to Enhance Salt Stress Tolerance. *Plants* **2025**, *14*, 1804. [\[CrossRef\]](#) [\[PubMed\]](#)
65. İköz, B.; Dasgan, H.Y.; Balik, S.; Kusvuran, S.; Gruda, N.S. The use of biostimulants as a key to sustainable hydroponic lettuce farming under saline water stress. *BMC Plant Biol.* **2024**, *24*, 808. [\[CrossRef\]](#) [\[PubMed\]](#)

66. Amerian, M.; Palangi, A.; Gohari, G.; Ntatsi, G. Humic acid and grafting as sustainable agronomic practices for increased growth and secondary metabolism in cucumber subjected to salt stress. *Sci. Rep.* **2024**, *14*, 15883. [\[CrossRef\]](#) [\[PubMed\]](#)
67. Sardar, H.; Khalid, Z.; Ahsan, M.; Naz, S.; Nawaz, A.; Ahmad, R.; Razzaq, K.; Wabaidur, S.M.; Jacquard, C.; Širić, I.; et al. Enhancement of Salinity Stress Tolerance in Lettuce (*Lactuca sativa* L.) via Foliar Application of Nitric Oxide. *Plants* **2023**, *12*, 1115. [\[CrossRef\]](#) [\[PubMed\]](#)
68. Moreno-Lora, A.; Calero-Velázquez, R.; Arenas-Arenas, F.J. Physiological Responses of New Citrus Rootstocks To Salinity Stress Induced by Increasing Sodium and Chloride Concentrations in the Irrigation Solution. *J. Soil Sci. Plant Nutr.* **2025**, *25*, 7868–7877. [\[CrossRef\]](#)
69. Wang, Y.; Lu, Q.; Zhang, F.; Wang, W.; Wu, C. Effects of Biochar on the Yield of Melon and the Diversity of Rhizosphere Soil Microbial Communities Under Saline-Alkali Stress. *Plants* **2025**, *14*, 1423. [\[CrossRef\]](#) [\[PubMed\]](#)
70. Sánchez, P.; Castro-Cegrí, A.; Sierra, S.; Garrido, D.; Llamas, I.; Sampedro, I.; Palma, F. The synergy of halotolerant PGPB and mauran mitigates salt stress in tomato (*Solanum lycopersicum*) via osmoprotectants accumulation. *Physiol. Plant.* **2023**, *175*, e14111. [\[CrossRef\]](#) [\[PubMed\]](#)
71. Pal, S.C.; Hossain, M.B.; Mallick, D.; Bushra, F.; Abdullah, S.R.; Dash, P.K.; Das, D. Combined use of seaweed extract and arbuscular mycorrhizal fungi for alleviating salt stress in bell pepper (*Capsicum annuum* L.). *Sci. Hortic.* **2024**, *325*, 112597. [\[CrossRef\]](#)
72. Daldoul, S.; Gargouri, M.; Weinert, C.; Jarrar, A.; Egert, B.; Mliki, A.; Nick, P. A Tunisian wild grape leads to metabolic fingerprints of salt tolerance. *Plant Physiol.* **2023**, *193*, 371–388. [\[CrossRef\]](#) [\[PubMed\]](#)
73. Amerian, M.; Palangi, A.; Gohari, G.; Ntatsi, G. Enhancing salinity tolerance in cucumber through Selenium biofortification and grafting. *BMC Plant Biol.* **2024**, *24*, 24. [\[CrossRef\]](#) [\[PubMed\]](#)
74. Sheikhalipour, M.; Esmailpour, B.; Behnamian, M.; Gohari, G.; Giglou, M.T.; Vachova, P.; Rastogi, A.; Brestic, M.; Skalicky, M. Chitosan-Selenium Nanoparticle (Cs-Se NP) Foliar Spray Alleviates Salt Stress in Bitter Melon. *Nanomaterials* **2021**, *11*, 684. [\[CrossRef\]](#) [\[PubMed\]](#)
75. Silva, E.F.D.; Santos, H.R.B.; Ometto, J.P.H.B.; Jardim, A.M.D.R.F.; Silva, T.G.F.D.; Hermínio, P.J.; Simões, A.N.; Souza, E.; Ferreira-Silva, S.L. Salt-excluder rootstock improves physio-biochemical responses of grafted grapevine plants subjected to salinity stress. *Curr. Plant Biol.* **2024**, *37*, 100316. [\[CrossRef\]](#)
76. Abdelkader, M.; Voronina, L.; Shelepova, O.; Puchkov, M.; Loktionova, E.; Zhanbyrshina, N.; Yelnazarkyzy, R.; Tleppayeva, A.; Ksenofontov, A. Monitoring Role of Exogenous Amino Acids on the Proteinogenic and Ionic Responses of Lettuce Plants under Salinity Stress Conditions. *Horticulturae* **2023**, *9*, 626. [\[CrossRef\]](#)
77. Zuzunaga-Rosas, J.; Calone, R.; Mircea, D.M.; Shakya, R.; Ibáñez-Asensio, S.; Boscaiu, M.; Fita, A.; Moreno-Ramón, H.; Vicente, O. Mitigation of salt stress in lettuce by a biostimulant that protects the root absorption zone and improves biochemical responses. *Front. Plant Sci.* **2024**, *15*, 1341714. [\[CrossRef\]](#) [\[PubMed\]](#)
78. Zhang, M.; Li, X.; Wang, X.; Feng, J.; Zhu, S. Potassium fulvic acid alleviates salt stress of citrus by regulating rhizosphere microbial community, osmotic substances and enzyme activities. *Front. Plant Sci.* **2023**, *14*, 1161469. [\[CrossRef\]](#) [\[PubMed\]](#)
79. Mady, E.; Abd El-Wahed, A.H.M.; Awad, A.H.; Asar, T.O.; Al-Farga, A.; Abd El-Raouf, H.S.; Randhir, R.; Alnuzaile, E.S.; El-Taher, A.M.; Randhir, T.O.; et al. Evaluation of Salicylic Acid Effects on Growth, Biochemical, Yield, and Anatomical Characteristics of Eggplant (*Solanum melongena* L.) Plants under Salt Stress Conditions. *Agronomy* **2023**, *13*, 2213. [\[CrossRef\]](#)
80. Abdelkader, M.; Voronina, L.; Baratova, L.; Shelepova, O.; Zargar, M.; Puchkov, M.; Loktionova, E.; Amantayev, B.; Kipshakbaeva, A.; Arinov, B. Biostimulants-Based Amino Acids Augment Physio-Biochemical Responses and Promote Salinity Tolerance of Lettuce Plants (*Lactuca sativa* L.). *Horticulturae* **2023**, *9*, 807. [\[CrossRef\]](#)
81. Abdelaal, K.A.A.; Mazrou, Y.S.A.; Hafez, Y.M. Silicon Foliar Application Mitigates Salt Stress in Sweet Pepper Plants by Enhancing Water Status, Photosynthesis, Antioxidant Enzyme Activity and Fruit Yield. *Plants* **2020**, *9*, 733. [\[CrossRef\]](#) [\[PubMed\]](#)
82. Patani, A.; Prajapati, D.; Ali, D.; Kalasariya, H.; Yadav, V.K.; Tank, J.; Bagatharia, S.; Joshi, M.; Patel, A. Evaluation of the growth-inducing efficacy of various *Bacillus* species on the salt-stressed tomato (*Lycopersicon esculentum* Mill.). *Front. Plant Sci.* **2023**, *14*, 1168155. [\[CrossRef\]](#) [\[PubMed\]](#)
83. Nikolaou, K.E.; Chatzistathis, T.; Theocharis, S.; Argiriou, A.; Koundouras, S.; Zioziou, E. Effects of Salinity and Rootstock on Nutrient Element Concentrations and Physiology in Own-Rooted or Grafted to 1103 P and 101-14 Mgt Rootstocks of Merlot and Cabernet Franc Grapevine Cultivars under Climate Change. *Sustainability* **2021**, *13*, 2477. [\[CrossRef\]](#)
84. Zuluaga, M.Y.A.; Monterisi, S.; Roupheal, Y.; Colla, G.; Lucini, L.; Cesco, S.; Pii, Y. Different vegetal protein hydrolysates distinctively alleviate salinity stress in vegetable crops: A case study on tomato and lettuce. *Front. Plant Sci.* **2023**, *14*, 1077140. [\[CrossRef\]](#) [\[PubMed\]](#)
85. Abu Zeid, I.M.; Mohamed, F.H.; Metwali, E.M.R. Responses of two strawberry cultivars to NaCl-induced salt stress under the influence of ZnO nanoparticles. *Saudi J. Biol. Sci.* **2023**, *30*, 103623. [\[CrossRef\]](#) [\[PubMed\]](#)

86. Gajjar, P.; Ismail, A.; Islam, T.; Darwish, A.G.; Moniruzzaman, M.; Abuslima, E.; Dawood, A.S.; El-Saady, A.M.; Tsoleva, V.; El-Kereamy, A.; et al. Physiological Comparison of Two Salt-Excluder Hybrid Grapevine Rootstocks under Salinity Reveals Different Adaptation Qualities. *Plants* **2023**, *12*, 3247. [\[CrossRef\]](#) [\[PubMed\]](#)
87. Jiang, T.; Chen, J.; Huang, Y.; Chang, X.; Wu, Y.; Liu, G.; Wang, R.; Xu, K.; Lu, L.; Tian, S. Characteristics of bacterial communities in rhizosphere and bulk soil in Fe-deficient citrus growing in coastal saline-alkali land. *Front. Plant Sci.* **2024**, *14*, 1335843. [\[CrossRef\]](#) [\[PubMed\]](#)
88. Cristofano, F.; El-Nakhel, C.; Colla, G.; Cardarelli, M.; Pii, Y.; Lucini, L.; Roupahel, Y. Modulation of Morpho-Physiological and Metabolic Profiles of Lettuce Subjected to Salt Stress and Treated with Two Vegetal-Derived Biostimulants. *Plants* **2023**, *12*, 709. [\[CrossRef\]](#) [\[PubMed\]](#)
89. Ge, L.; Yang, X.; Liu, Y.; Tang, H.; Wang, Q.; Chu, S.; Hu, J.; Zhang, N.; Shi, Q. Improvement of Seed Germination under Salt Stress via Overexpressing Caffeic Acid O-methyltransferase 1 (*SICOMT1*) in *Solanum lycopersicum* L. *Int. J. Mol. Sci.* **2023**, *24*, 734. [\[CrossRef\]](#) [\[PubMed\]](#)
90. Auñón-Calles, D.; Pinciroli, M.; Nicolás, E.; Gil-Izquierdo, A.; Gabaldón, J.A.; Sánchez-Iglesias, M.P.; Carbonell-Barrachina, A.A.; Ferreres, F.; García, C.J.; Romero-Trigueros, C. Agronomic and Metabolic Responses of Citrus clementina to Long-Term Irrigation with Saline Reclaimed Water as Abiotic Factor. *Int. J. Mol. Sci.* **2025**, *26*, 3450. [\[CrossRef\]](#) [\[PubMed\]](#)
91. Jiang, J.; Ren, X.; Li, L.; Hou, R.; Sun, W.; Jiao, C.; Yang, N.; Dong, Y. H₂S Regulation of Metabolism in Cucumber in Response to Salt-Stress Through Transcriptome and Proteome Analysis. *Front. Plant Sci.* **2020**, *11*, 1283. [\[CrossRef\]](#) [\[PubMed\]](#)
92. Malekzadeh, M.R.; Roosta, H.R.; Kalaji, H.M. GO nanoparticles mitigate the negative effects of salt and alkalinity stress by enhancing gas exchange and photosynthetic efficiency of strawberry plants. *Sci. Rep.* **2023**, *13*, 8457. [\[CrossRef\]](#) [\[PubMed\]](#)
93. El-Ghamry, A.M.; El-Sherpiny, M.A.; Alkharpotly, A.E.A.; Ghazi, D.A.; Helmy, A.A.; Siddiqui, M.H.; Pessarakli, M.; Hossain, M.A.; Elghareeb, E.M. The synergistic effects of organic composts and microelements co-application in enhancing potato productivity in saline soils. *Heliyon* **2024**, *10*, e32694. [\[CrossRef\]](#) [\[PubMed\]](#)
94. Kacjan Maršić, N.; Štolfa, P.; Vodnik, D.; Košmelj, K.; Mikulič-Petkovšek, M.; Kump, B.; Vidrih, R.; Kokalj, D.; Piskernik, S.; Ferjančič, B.; et al. Physiological and Biochemical Responses of Ungrafted and Grafted Bell Pepper Plants (*Capsicum annuum* L. var. *grossum* (L.) Sendtn.) Grown under Moderate Salt Stress. *Plants* **2021**, *10*, 314. [\[CrossRef\]](#) [\[PubMed\]](#)
95. Carrasco, D.; Zhou-Tsang, A.; Rodriguez-Izquierdo, A.; Ocete, R.; Revilla, M.A.; Arroyo-García, R. Coastal Wild Grapevine Accession (*Vitis vinifera* L. ssp. *sylvestris*) Shows Distinct Late and Early Transcriptome Changes under Salt Stress in Comparison to Commercial Rootstock Richter 110. *Plants* **2022**, *11*, 2688. [\[CrossRef\]](#) [\[PubMed\]](#)
96. Abdeldym, E.A.; El-Mogy, M.M.; Abdellateaf, H.R.L.; Atia, M.A.M. Genetic Characterization, Agro-Morphological and Physiological Evaluation of Grafted Tomato under Salinity Stress Conditions. *Agronomy* **2020**, *10*, 1948. [\[CrossRef\]](#)
97. Fedeli, R.; Celletti, S.; Loppi, S. Wood Distillate Promotes the Tolerance of Lettuce in Extreme Salt Stress Conditions. *Plants* **2024**, *13*, 1335. [\[CrossRef\]](#) [\[PubMed\]](#)
98. Abdelkader, M.; Suliman, A.A.; Salem, S.S.; Assiya, A.; Voronina, L.; Puchkov, M.; Loktionova, E.; Bhuker, A.; Ataya, F.S.; Mahmoud, M.H.; et al. Studying the Combined Impact of Salinity and Drought Stress-Simulated Conditions on Physio-Biochemical Characteristics of Lettuce Plant. *Horticulturae* **2024**, *10*, 1186. [\[CrossRef\]](#)
99. Liu, J.; Yu, Y.; Huang, Z.; Hu, J.; Gao, N.; Wang, X.; Qi, J.; Cao, Y.; Li, X.; Wang, X.; et al. Organic amendment with mushroom residue alleviates salt stress in watermelon seedlings. *Sci. Hortic.* **2025**, *353*, 114508. [\[CrossRef\]](#)
100. Martinez-Alonso, A.; Nicolás-Espinosa, J.; Carvajal, M.; Bárzana, G. The differential expressions of aquaporins underline the diverse strategies of cucumber and tomato against salinity and zinc stress. *Physiol. Plant.* **2024**, *176*, e14222. [\[CrossRef\]](#) [\[PubMed\]](#)
101. Jalalian, S.; Ebrahimzadeh, A.; Zahedi, S.M.; Becker, S.J.; Hayati, F.; Hassanpouraghdam, M.B.; Rasouli, F. *Chlamydomonas* sp. extract meliorates the growth and physiological responses of ‘Camarosa’ strawberry (*Fragaria × ananassa* Duch) under salinity stress. *Sci. Rep.* **2024**, *14*, 22436. [\[CrossRef\]](#) [\[PubMed\]](#)
102. Zhang, J.; Liu, X.; Yin, Z.; Zhao, T.; Du, D.; Li, J.; Zhu, M.; Sun, Y.; Pan, Y. Genome- and Transcriptome-Wide Characterization and Expression Analyses of bHLH Transcription Factor Family Reveal Their Relevance to Salt Stress Response in Tomato. *Plants* **2025**, *14*, 200. [\[CrossRef\]](#) [\[PubMed\]](#)
103. Lu, X.; Chen, G.; Ma, L.; Zhang, C.; Yan, H.; Bao, J.; Nai, G.; Wang, W.; Chen, B.; Ma, S.; et al. Integrated transcriptome and metabolome analysis reveals antioxidant machinery in grapevine exposed to salt and alkali stress. *Physiol. Plant.* **2023**, *175*, e13950. [\[CrossRef\]](#) [\[PubMed\]](#)
104. Sarwar, M.; Anjum, S.; Ali, Q.; Alam, M.W.; Haider, M.S.; Mehboob, W. Triacntanol modulates salt stress tolerance in cucumber by altering the physiological and biochemical status of plant cells. *Sci. Rep.* **2021**, *11*, 24504. [\[CrossRef\]](#) [\[PubMed\]](#)
105. Li, F.; Yuan, Y.; Gong, P.; Imazumi, Y.; Na, R.; Shimizu, N. Comparative effects of mineral fertilizer and digestate on growth, antioxidant system, and physiology of lettuce under salt stress. *Hortic. Environ. Biotechnol.* **2023**, *64*, 379–391. [\[CrossRef\]](#)
106. Zi, Y.; Zhang, M.; Yang, X.; Zhao, K.; Yin, T.; Wen, K.; Li, X.; Liu, X.; Zhang, H. Identification of the sweet orange (*Citrus sinensis*) bHLH gene family and the role of *CsbHLH55* and *CsbHLH87* in regulating salt stress. *Plant Genome* **2024**, *17*, e20502. [\[CrossRef\]](#) [\[PubMed\]](#)

107. Farooq, M.A.; Zeeshan Ul Haq, M.; Zhang, L.; Wu, S.; Mushtaq, N.; Tahir, H.; Wang, Z. Transcriptomic Insights into Salt Stress Response in Two Pepper Species: The Role of MAPK and Plant Hormone Signaling Pathways. *Int. J. Mol. Sci.* **2024**, *25*, 9355. [\[CrossRef\]](#) [\[PubMed\]](#)
108. Danaeifar, A.; Khaleghi, E.; Zivdar, S.; Mehdikhanlou, K. Physiological, biochemical, and gene expression of Sour Orange (*Citrus aurantium* L.) to Iron(II)-Arginine Chelate under salinity, alkalinity, and salt-alkali combined stresses. *Sci. Hortic.* **2023**, *319*, 112146. [\[CrossRef\]](#)
109. Zhang, F.; Zhang, J.; Li, Q.; Yang, Y.; Sheng, Y. Exploring Candidate Genes and Regulatory Mechanisms for Salt-Alkali Tolerance in Cucumber. *Agronomy* **2024**, *14*, 543. [\[CrossRef\]](#)
110. Wang, L.; Li, R.; Li, K.; Qu, Z.; Zhou, R.; Lu, G.; Li, P.; Li, G. Genome-wide identification of the grapevine β -1,3-glucanase gene (*VviBG*) family and expression analysis under different stresses. *BMC Plant Biol.* **2024**, *24*, 911. [\[CrossRef\]](#) [\[PubMed\]](#)
111. Sadler, M.T.; Ali, A.A.M.; Alsadon, A.A.; Wahb-Allah, M.A. Long-Term Salinity-Responsive Transcriptome in Advanced Breeding Lines of Tomato. *Plants* **2025**, *14*, 100. [\[CrossRef\]](#) [\[PubMed\]](#)
112. Abdelkhalik, A.; Abd El-Mageed, T.A.; Mohamed, I.A.A.; Semida, W.M.; Al-Elwany, O.A.A.I.; Ibrahim, I.M.; Hemida, K.A.; El-Saadony, M.T.; AbuQamar, S.F.; El-Tarabily, K.A.; et al. Soil application of effective microorganisms and nitrogen alleviates salt stress in hot pepper (*Capsicum annum* L.) plants. *Front. Plant Sci.* **2023**, *13*, 1079260. [\[CrossRef\]](#) [\[PubMed\]](#)
113. Chourasia, K.N.; Lal, M.K.; Tiwari, R.K.; Dev, D.; Kardile, H.B.; Patil, V.U.; Kumar, A.; Vanishree, G.; Kumar, D.; Bhardwaj, V.; et al. Salinity Stress in Potato: Understanding Physiological, Biochemical and Molecular Responses. *Life* **2021**, *11*, 545. [\[CrossRef\]](#) [\[PubMed\]](#)
114. Lu, X.; Ma, L.; Zhang, C.; Yan, H.; Bao, J.; Gong, M.; Wang, W.; Li, S.; Ma, S.; Chen, B. Grapevine (*Vitis vinifera*) responses to salt stress and alkali stress: Transcriptional and metabolic profiling. *BMC Plant Biol.* **2022**, *22*, 528. [\[CrossRef\]](#) [\[PubMed\]](#)
115. Li, C.; Lu, X.; Liu, Y.; Xu, J.; Yu, W. Strigolactone Alleviates the Adverse Effects of Salt Stress on Seed Germination in Cucumber by Enhancing Antioxidant Capacity. *Antioxidants* **2023**, *12*, 1043. [\[CrossRef\]](#) [\[PubMed\]](#)
116. Zhang, G.; Wang, Y.; Wu, K.; Zhang, Q.; Feng, Y.; Miao, Y.; Yan, Z. Exogenous Application of Chitosan Alleviate Salinity Stress in Lettuce (*Lactuca sativa* L.). *Horticulturae* **2021**, *7*, 342. [\[CrossRef\]](#)
117. Duan, S.; Xu, Z.; Li, X.Y.; Liao, P.; Qin, H.K.; Mao, Y.P.; Dai, W.S.; Ma, H.-J.; Bao, M.L. Dodder-transmitted mobile systemic signals activate a salt-stress response characterized by a transcriptome change in *Citrus sinensis*. *Front. Plant Sci.* **2022**, *13*, 986365. [\[CrossRef\]](#) [\[PubMed\]](#)
118. Liu, T.; Amanullah, S.; Xu, H.; Gao, P.; Du, Z.; Hu, X.; Han, M.; Che, Y.; Zhang, L.; Qi, G.; et al. RNA-Seq Identified Putative Genes Conferring Photosynthesis and Root Development of Melon under Salt Stress. *Genes* **2023**, *14*, 1728. [\[CrossRef\]](#) [\[PubMed\]](#)
119. Chen, T.W.; Gomez Pineda, I.M.; Brand, A.M.; Stützel, H. Determining Ion Toxicity in Cucumber under Salinity Stress. *Agronomy* **2020**, *10*, 677. [\[CrossRef\]](#)
120. Batelli, G.; Ruggiero, A.; Esposito, S.; Venezia, A.; Lupini, A.; Nurcato, R.; Costa, A.; Palombieri, S.; Vitiello, A.; Mauceri, A.; et al. Combined salt and low nitrate stress conditions lead to morphophysiological changes and tissue-specific transcriptome reprogramming in tomato. *Plant Physiol. Biochem.* **2024**, *215*, 108976. [\[CrossRef\]](#) [\[PubMed\]](#)
121. Hosseinpour, A.; Haliloglu, K.; Tolga Cinisli, K.; Ozkan, G.; Ozturk, H.I.; Pour-Aboughadareh, A.; Pocza, P. Application of Zinc Oxide Nanoparticles and Plant Growth Promoting Bacteria Reduces Genetic Impairment under Salt Stress in Tomato (*Solanum lycopersicum* L. 'Linda'). *Agriculture* **2020**, *10*, 521. [\[CrossRef\]](#)
122. Silva, J.E.S.B.D.; Torres, S.B.; Leal, C.C.P.; Leite, M.D.S.; Guirra, K.S.; Nogueira Neto, F.A.; Dantas, B.F. Pre-germination treatments with plant growth regulators and bioactivators attenuate salt stress in melon: Effects on germination and seedling development. *Acta Sci. Agron.* **2023**, *45*, e60516. [\[CrossRef\]](#)
123. Kafi, M.; Nabati, J.; Ahmadi-Lahijani, M.J.; Oskoueian, A. Silicon Compounds and Potassium Sulfate Improve Salinity Tolerance of Potato Plants through Instigating the Defense Mechanisms, Cell Membrane Stability, and Accumulation of Osmolytes. *Commun. Soil Sci. Plant Anal.* **2021**, *52*, 843–858. [\[CrossRef\]](#)
124. Bello, A.S.; Ben-Hamadou, R.; Hamdi, H.; Saadaoui, I.; Ahmed, T. Application of Cyanobacteria (*Roholtiella* sp.) Liquid Extract for the Alleviation of Salt Stress in Bell Pepper (*Capsicum annum* L.) Plants Grown in a Soilless System. *Plants* **2021**, *11*, 104. [\[CrossRef\]](#) [\[PubMed\]](#)
125. Jin, Y.; Yang, P.; Li, J.; Yang, Y.; Yang, R.; Fu, H.; Li, J. Brassinosteroids Alleviate Salt Stress by Enhancing Sugar and Glycine Betaine in Pepper (*Capsicum annum* L.). *Plants* **2024**, *13*, 3029. [\[CrossRef\]](#) [\[PubMed\]](#)
126. Xu, J.; Sui, C.; Ge, J.; Ren, R.; Pang, Y.; Gan, H.; Du, Y.; Cao, H.; Sun, Q. Exogenous spermidine improved the salinity-alkalinity stress tolerance of grapevine (*Vitis vinifera*) by regulating antioxidant system, Na⁺/K⁺ homeostasis and endogenous polyamine contents. *Sci. Hortic.* **2024**, *326*, 112725. [\[CrossRef\]](#)
127. Brenes, M.; Pérez, J.; González-Orenga, S.; Solana, A.; Boscaiu, M.; Prohens, J.; Plazas, M.; Fita, A.; Vicente, O. Comparative Studies on the Physiological and Biochemical Responses to Salt Stress of Eggplant (*Solanum melongena*) and Its Rootstock *S. torvum*. *Agriculture* **2020**, *10*, 328. [\[CrossRef\]](#)

128. Babalik, Z.; Göktürk Baydar, N. Drought and salt stress alter the mineral composition of grapevines. *J. Plant Nutr.* **2024**, *47*, 1981–1995. [\[CrossRef\]](#)
129. Song, C.; Dong, S.; Schlisser, A.; Lupo, Y.; Rachmilevitch, S.; Lazarovitch, N.; Fait, A. Rootstock varietal ability in accumulation of chloride ions underpins improved physiology and metabolism of grapevine exposed to salinity. *Sci. Hortic.* **2024**, *328*, 112964. [\[CrossRef\]](#)
130. Palmitessa, O.D.; Renna, M.; De Angelis, D.; Signore, A.; Serio, F.; Summo, C.; Santamaria, P. Moderate saline waters are effective to enhance a landrace of unripe melon cultivated in a “water culture system” with high input efficiency. *Sci. Hortic.* **2024**, *337*, 113599. [\[CrossRef\]](#)
131. Zhao, F.; Zheng, T.; Liu, Z.; Fu, W.; Fang, J. Transcriptomic Analysis Elaborates the Resistance Mechanism of Grapevine Rootstocks against Salt Stress. *Plants* **2022**, *11*, 1167. [\[CrossRef\]](#) [\[PubMed\]](#)
132. Shen, L.; Xia, X.; Zhang, L.; Yang, S.; Yang, X. Genome-Wide Identification of Catalase Gene Family and the Function of *SmCAT4* in Eggplant Response to Salt Stress. *Int. J. Mol. Sci.* **2023**, *24*, 16979. [\[CrossRef\]](#) [\[PubMed\]](#)
133. Shumaila, S.; Ullah, S. Mitigation of Salinity-Induced Damages in *Capsicum annum* L. (Sweet Pepper) Seedlings Using Priming Techniques: A Future Perspective of Climate Change in the Region. *Commun. Soil Sci. Plant Anal.* **2020**, *51*, 1602–1625. [\[CrossRef\]](#)
134. Oliveira, V.K.N.; Silva, A.A.R.D.; Lima, G.S.D.; Soares, L.A.D.A.; Gheyi, H.R.; de Lacerda, C.F.; Azevedo, C.A.V.; Nobre, R.G.; Chaves, L.H.G.; Fernandes, P.D.; et al. Foliar Application of Salicylic Acid Mitigates Saline Stress on Physiology, Production, and Post-Harvest Quality of Hydroponic Japanese Cucumber. *Agriculture* **2023**, *13*, 395. [\[CrossRef\]](#)
135. Shen, H.; Zhou, Y.; Liao, C.; Xie, Q.; Chen, G.; Hu, Z.; Wu, T. The AlkB Homolog *SLALKBH10B* Negatively Affects Drought and Salt Tolerance in *Solanum lycopersicum*. *Int. J. Mol. Sci.* **2023**, *25*, 173. [\[CrossRef\]](#) [\[PubMed\]](#)
136. Ekinci, M.; Yildirim, E.; Turan, M. Ameliorating effects of hydrogen sulfide on growth, physiological and biochemical characteristics of eggplant seedlings under salt stress. *S. Afr. J. Bot.* **2021**, *143*, 79–89. [\[CrossRef\]](#)
137. Sanwal, S.K.; Kumar, P.; Kesh, H.; Gupta, V.K.; Kumar, A.; Kumar, A.; Meena, B.L.; Colla, G.; Cardarelli, M.; Kumar, P. Salinity Stress Tolerance in Potato Cultivars: Evidence from Physiological and Biochemical Traits. *Plants* **2022**, *11*, 1842. [\[CrossRef\]](#) [\[PubMed\]](#)
138. Shalaby, T.A.; Abd-Alkarim, E.; El-Aidy, F.; Hamed, E.S.; Sharaf-Eldin, M.; Taha, N.; El-Ramady, H.; Bayoumi, Y.; dos Reis, A.R. Nano-selenium, silicon and H₂O₂ boost growth and productivity of cucumber under combined salinity and heat stress. *Ecotoxicol. Environ. Saf.* **2021**, *212*, 111962. [\[CrossRef\]](#) [\[PubMed\]](#)
139. Aazami, M.A.; Rasouli, F.; Ebrahimzadeh, A. Oxidative damage, antioxidant mechanism and gene expression in tomato responding to salinity stress under in vitro conditions and application of iron and zinc oxide nanoparticles on callus induction and plant regeneration. *BMC Plant Biol.* **2021**, *21*, 597. [\[CrossRef\]](#) [\[PubMed\]](#)
140. Sheikhalipour, M.; Mohammadi, S.A.; Esmailpour, B.; Spanos, A.; Mahmoudi, R.; Mahdavinia, G.R.; Milani, M.H.; Kahnemoei, A.H.; Nouraein, M.; Antoniou, C.; et al. Seedling nanoprimer with selenium-chitosan nanoparticles mitigates the adverse effects of salt stress by inducing multiple defence pathways in bitter melon plants. *Int. J. Biol. Macromol.* **2023**, *242*, 124923. [\[CrossRef\]](#) [\[PubMed\]](#)
141. Gálvez, A.; Albacete, A.; Martínez-Andújar, C.; del Amor, F.M.; López-Marín, J. Contrasting Rootstock-Mediated Growth and Yield Responses in Salinized Pepper Plants (*Capsicum annum* L.) Are Associated with Changes in the Hormonal Balance. *Int. J. Mol. Sci.* **2021**, *22*, 3297. [\[CrossRef\]](#) [\[PubMed\]](#)
142. Liu, Y.; Wei, L.; Feng, L.; Zhang, M.; Hu, D.; Tie, J.; Liao, W. Hydrogen Sulfide Promotes Adventitious Root Development in Cucumber under Salt Stress by Enhancing Antioxidant Ability. *Plants* **2022**, *11*, 935. [\[CrossRef\]](#) [\[PubMed\]](#)
143. Snoussi, H.; Askri, H.; Nacouzi, D.; Ouerghui, I.; Ananga, A.; Najar, A.; El Kayal, W. Comparative Transcriptome Profiling of Salinity-Induced Genes in Citrus Rootstocks with Contrasted Salt Tolerance. *Agriculture* **2022**, *12*, 350. [\[CrossRef\]](#)
144. Kapadia, C.; Sayyed, R.Z.; El Enshasy, H.A.; Vaidya, H.; Sharma, D.; Patel, N.; Malek, R.A.; Syed, A.; Elgorban, A.M.; Ahmad, K.; et al. Halotolerant Microbial Consortia for Sustainable Mitigation of Salinity Stress, Growth Promotion, and Mineral Uptake in Tomato Plants and Soil Nutrient Enrichment. *Sustainability* **2021**, *13*, 8369. [\[CrossRef\]](#)
145. Li, W.; Wei, Y.; Zhang, L.; Wang, Y.; Song, P.; Li, X.; Han, D. *FvMYB44*, a Strawberry R2R3-MYB Transcription Factor, Improved Salt and Cold Stress Tolerance in Transgenic Arabidopsis. *Agronomy* **2023**, *13*, 1051. [\[CrossRef\]](#)
146. Wang, R.; Yang, X.; Chi, Y.; Zhang, X.; Ma, X.; Zhang, D.; Zhao, T.; Ren, Y.; Yang, H.; Ding, W.; et al. Regulation of hydrogen rich water on strawberry seedlings and root endophytic bacteria under salt stress. *Front. Plant Sci.* **2024**, *15*, 1497362. [\[CrossRef\]](#) [\[PubMed\]](#)
147. Guo, M.; Wang, X.S.; Guo, H.D.; Bai, S.Y.; Khan, A.; Wang, X.M.; Gao, Y.-M.; Li, J.-S. Tomato salt tolerance mechanisms and their potential applications for fighting salinity: A review. *Front. Plant Sci.* **2022**, *13*, 949541. [\[CrossRef\]](#) [\[PubMed\]](#)
148. Ramzan, M.; Haider, S.T.A.; Hussain, M.B.; Ehsan, A.; Datta, R.; Alahmadi, T.A.; Ansari, M.J.; Alharbi, S.A. Potential of kaempferol and caffeic acid to mitigate salinity stress and improving potato growth. *Sci. Rep.* **2024**, *14*, 21657. [\[CrossRef\]](#) [\[PubMed\]](#)
149. Gedeon, S.; Ioannou, A.; Balestrini, R.; Fotopoulos, V.; Antoniou, C. Application of Biostimulants in Tomato Plants (*Solanum lycopersicum*) to Enhance Plant Growth and Salt Stress Tolerance. *Plants* **2022**, *11*, 3082. [\[CrossRef\]](#) [\[PubMed\]](#)

150. Brenes, M.; Solana, A.; Boscaiu, M.; Fita, A.; Vicente, O.; Calatayud, Á.; Prohens, J.; Plazas, M. Physiological and Biochemical Responses to Salt Stress in Cultivated Eggplant (*Solanum melongena* L.) and in *S. insanum* L., a Close Wild Relative. *Agronomy* **2020**, *10*, 651. [\[CrossRef\]](#)
151. Campobenedetto, C.; Mannino, G.; Beekwilder, J.; Contartese, V.; Karlova, R.; Berteà, C.M. The application of a biostimulant based on tannins affects root architecture and improves tolerance to salinity in tomato plants. *Sci. Rep.* **2021**, *11*, 354. [\[CrossRef\]](#) [\[PubMed\]](#)
152. Zang, Y.; Huang, Y.; Chang, X.; Chen, J.; Jiang, T.; Wu, Z.; Lu, L.; Tian, S. High Soil pH and Plastic-Shed Lead to Iron Deficiency and Chlorosis of Citrus in Coastal Saline-Alkali Lands: A Field Study in Xiangshan County. *Horticulturae* **2023**, *9*, 437. [\[CrossRef\]](#)
153. Zhang, F.; Li, D.; Sa, R.; Wang, L.; Sheng, Y. Cloning and Function Analysis of the CsTAU1 in Response to Salt-Alkali Stress. *Genes* **2024**, *15*, 613. [\[CrossRef\]](#) [\[PubMed\]](#)
154. Roy Choudhury, A.; Trivedi, P.; Choi, J.; Madhaiyan, M.; Park, J.H.; Choi, W.; Walitang, D.I.; Sa, T. Inoculation of ACC deaminase-producing endophytic bacteria down-regulates ethylene-induced pathogenesis-related signaling in red pepper (*Capsicum annuum* L.) under salt stress. *Physiol. Plant.* **2023**, *175*, e13909. [\[CrossRef\]](#) [\[PubMed\]](#)
155. Darwish, H.; Al-Osaimi, G.S.; Al Kashgry, N.A.T.; Sonbol, H.; Alayafi, A.A.M.; Alabdallah, N.M.; Al-Humaid, A.; Al-Harbi, N.A.; Al-Qahtani, S.M.; Abbas, Z.K.; et al. Evaluating the genotoxicity of salinity stress and secondary products gene manipulation in lime, *Citrus aurantifolia*, plants. *Front. Plant Sci.* **2023**, *14*, 1211595. [\[CrossRef\]](#) [\[PubMed\]](#)
156. Bonnin, M.; Favreau, B.; Soriano, A.; Leonhardt, N.; Oustric, J.; Lourkisti, R.; Ollitrault, P.; Morillon, R.; Berti, L.; Santini, J. Insight into Physiological and Biochemical Determinants of Salt Stress Tolerance in Tetraploid Citrus. *Antioxidants* **2023**, *12*, 1640. [\[CrossRef\]](#) [\[PubMed\]](#)
157. Gohari, G.; Zareei, E.; Rostami, H.; Panahirad, S.; Kulak, M.; Farhadi, H.; Amini, M.; Martinez-Ballesta, M.d.C.; Fotopoulos, V. Protective effects of cerium oxide nanoparticles in grapevine (*Vitis vinifera* L.) cv. Flame Seedless under salt stress conditions. *Ecotoxicol. Environ. Saf.* **2021**, *220*, 112402. [\[CrossRef\]](#) [\[PubMed\]](#)
158. Petoumenou, D.G.; Liava, V. Sustainable Foliar Applications to Improve Grapevine Responses to Drought, High Temperatures, and Salinity: Impacts on Physiology, Yields, and Berry Quality. *Plants* **2025**, *14*, 2157. [\[CrossRef\]](#) [\[PubMed\]](#)
159. García-López, J.V.; Redondo-Gómez, S.; Flores-Duarte, N.J.; Rodríguez-Llorente, I.D.; Pajuelo, E.; Mateos-Naranjo, E. PGPR-based biofertilizer modulates strawberry photosynthetic apparatus tolerance responses by severe drought, soil salinization and short extreme heat event. *Plant Stress* **2024**, *12*, 100448. [\[CrossRef\]](#)
160. Suarez, D.L.; Celis, N.; Ferreira, J.F.S.; Reynolds, T.; Sandhu, D. Linking genetic determinants with salinity tolerance and ion relationships in eggplant, tomato and pepper. *Sci. Rep.* **2021**, *11*, 16298. [\[CrossRef\]](#) [\[PubMed\]](#)
161. Ntanos, E.; Kekelis, P.; Assimakopoulou, A.; Gasparatos, D.; Denaxa, N.K.; Tsafouros, A.; Roussos, P.A. Amelioration Effects against Salinity Stress in Strawberry by Bentonite-Zeolite Mixture, Glycine Betaine, and *Bacillus amyloliquefaciens* in Terms of Plant Growth, Nutrient Content, Soil Properties, Yield, and Fruit Quality Characteristics. *Appl. Sci.* **2021**, *11*, 8796. [\[CrossRef\]](#)
162. Parthasarathi, T.; Ephrath, J.E.; Lazarovitch, N. Grafting of tomato (*Solanum lycopersicum* L.) onto potato (*Solanum tuberosum* L.) to improve salinity tolerance. *Sci. Hortic.* **2021**, *282*, 110050. [\[CrossRef\]](#)
163. Li, Y.; Jiang, F.; Niu, L.; Wang, G.; Yin, J.; Song, X.; Ottosen, C.-O.; Rosenqvist, E.; Mittler, R.; Wu, Z.; et al. Synergistic regulation at physiological, transcriptional and metabolic levels in tomato plants subjected to a combination of salt and heat stress. *Plant J.* **2024**, *117*, 1656–1675. [\[CrossRef\]](#) [\[PubMed\]](#)
164. Amorim, T.L.; Santos, H.R.B.; Neto, J.B.; Hermínio, P.J.; Silva, J.R.I.; Silva, M.M.A.; Simões, A.N.; Souza, E.; Ferreira-Silva, S.L. Resistant rootstocks mitigate ionic toxicity with beneficial effects for growth and photosynthesis in grapevine grafted plants under salinity. *Sci. Hortic.* **2023**, *317*, 112053. [\[CrossRef\]](#)
165. Chevilly, S.; Dolz-Edo, L.; Martínez-Sánchez, G.; Morcillo, L.; Vilagrosa, A.; López-Nicolás, J.M.; Blanca, J.; Yenush, L.; Mulet, J.M. Distinctive Traits for Drought and Salt Stress Tolerance in Melon (*Cucumis melo* L.). *Front. Plant Sci.* **2021**, *12*, 777060. [\[CrossRef\]](#) [\[PubMed\]](#)
166. Montanaro, G.; Briglia, N.; Lopez, L.; Amato, D.; Panara, F.; Petrozza, A.; Cellini, F.; Nuzzo, V. A synthetic cytokinin primes photosynthetic and growth response in grapevine under ion-independent salinity stress. *J. Plant Interact.* **2022**, *17*, 789–800. [\[CrossRef\]](#)
167. Modarelli, G.C.; Roupheal, Y.; De Pascale, S.; Öztekin, G.B.; Tüzel, Y.; Orsini, F.; Gianquinto, G. Appraisal of Salt Tolerance under Greenhouse Conditions of a Cucurbitaceae Genetic Repository of Potential Rootstocks and Scions. *Agronomy* **2020**, *10*, 967. [\[CrossRef\]](#)
168. Abou-Sreya, A.I.B.; Azzam, C.R.; Al-Taweel, S.K.; Abdel-Aziz, R.M.; Belal, H.E.E.; Rady, M.M.; Abdel-Kader, A.A.S.; Majrashi, A.; Khaled, K.A.M. Natural Biostimulant Attenuates Salinity Stress Effects in Chili Pepper by Remodeling Antioxidant, Ion, and Phytohormone Balances, and Augments Gene Expression. *Plants* **2021**, *10*, 2316. [\[CrossRef\]](#) [\[PubMed\]](#)
169. Younis, U.; Danish, S.; Datta, R.; Al Obaid, S.; Ansari, M.J. Synergistic effects of boron and saponin in mitigating salinity stress to enhance sweet potato growth. *Sci. Rep.* **2024**, *14*, 12988. [\[CrossRef\]](#) [\[PubMed\]](#)

170. Ors, S.; Ekinci, M.; Yildirim, E.; Sahin, U.; Turan, M.; Dursun, A. Interactive effects of salinity and drought stress on photosynthetic characteristics and physiology of tomato (*Lycopersicon esculentum* L.) seedlings. *S. Afr. J. Bot.* **2021**, *137*, 335–339. [\[CrossRef\]](#)
171. Baggett, J.P.; Habibsadeh, S.; Toups, H.S.; Cochetel, N.; Ghan, R.; Robinson, M.L.; Barrios-Masias, F.H.; Cramer, G.R. Is foliar Cl⁻ concentration the cause of photosynthetic decline in grapevine during mild salinity? *OENO One* **2021**, *55*, 33–48. [\[CrossRef\]](#)
172. Hannachi, S.; Steppe, K.; Eloudi, M.; Mechi, L.; Bahrini, I.; Van Labeke, M.C. Salt Stress Induced Changes in Photosynthesis and Metabolic Profiles of One Tolerant ('Bonica') and One Sensitive ('Black Beauty') Eggplant Cultivars (*Solanum melongena* L.). *Plants* **2022**, *11*, 590. [\[CrossRef\]](#) [\[PubMed\]](#)

Disclaimer/Publisher's Note: The statements, opinions and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of MDPI and/or the editor(s). MDPI and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions or products referred to in the content.