Salt stress and hydroxyectoine enhance phosphate solubilisation and plant colonisation capacity of *Kosakonia radicincitans*

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**Highlights**

- *K. radicincitans* enhance P solubilization responding to osmotic stress and hydroxyectoine.
- Hydroxyectoine uptake by *K. radicincitans* cells can increase the production of acid phosphatases.
- Pre-conditioning of *K. radicincitans* by osmoadaptation improved radish plant yields.
- Osmoadaptation and the enrichment with hydroxyectoine increase plant colonisation.
- Successful microfermentation of *K. radicincitans* cells via the BioLector approach.

**Abstract**

Gram-negative bacterial endophytes have attracted research interest caused by their advantageous over epiphytic bacteria in plant nutrition and protection. However, research on these typically Gram-negative endophytes has deficiencies concerning the role of cultivation and pre-formulation strategies on further plant colonisation capabilities. Besides, the influence of cultivation conditions and osmotic stress within bacterial endophytes on their phosphate solubilising ability has not yet been addressed. By preconditioning cells with an osmoadaptation and a hydroxyectoine accumulation approach, this research aimed at enhancing the capability of the plant growth promoting bacterium *Kosakonia radicincitans* strain DSM 16656 to both solubilise phosphate and colonise plant seedlings. The results showed that halotolerant bacterial phenotypes increased the root-colonising capability by approximately 3-fold and presented growth-promoting effects in radish plants. Interestingly, findings also demonstrated that salt stress in the culture media along with the accumulation of hydroxyectoine led to an increase in the *in vitro* phosphate-solubilising ability by affecting the production of acid phosphatases, from 1.24 to 3.34 U mg⁻¹ for non-salt stressed cells and hydroxyectoine-added cells respectively. Thus, this approach
Introduction

Bacterial endophytes have attracted considerable attention because of their capability to promote plant growth through direct or indirect mechanisms [1,2]. Many bacterial endophytes can support host plants by countering negative impacts in the environment and are classified as plant growth-promoting endophytic bacteria (PGPEB) [3,4]. Colonisation by PGPEB is essential for providing benefits to host plants. Endophytic colonisation involves the entry, growth and proliferation of bacterial populations within the plant [2,3]. The colonisation patterns of PGPEB in plant tissues are strongly dependent on several biotic and abiotic factors [6,7]. Moreover, rhizosphere microbial communities and metabolic pathway profiles may be entirely different from those of endophytes [8]. Hence, variations in abiotic factors such as temperature [9], soil type [10], pH [11], and nutrients availability have been described to influence bacterial plant colonisation behaviour. Especially, soil salinity alters bacterial community composition and their functional activities [5,12,13]. Thus, bacterial endophytes may help the plant to withstand such sudden osmolarity peak conditions in their ecological niche.

Organic osmolytes enable organisms to adapt to environmental conditions by protecting cells or molecules against salt stress. These organic osmolytes are generally "compatible" with the metabolism of the cell without adversely affecting macromolecules or physiological processes and are referred to as compatible solutes [14]. Solutes accumulate either by synthesis or by transport from the extracellular medium through osmotically regulated transporters and mechanosensitive channels [15]. Moreover, compatible solutes provide beneficial enzyme functions, protecting against high temperature, desiccation, salinity, freeze-thaw procedures and even drying [16–18]. Among these, intracellular solutes such as hydroxyectoine is well known in halophytic bacteria [19,20] and is considered a protein protectant [21]. Despite these benefits, the effect of hydroxyectoine enrichment in bacterial cells on bacterial plant colonisation behaviour is so far missing.

To date, four genomes of endophytes belonging to Kosakonia radicincitans have been completed and published [22–24]. Isolation procedures of these strains revealed their endophytic capacity in different crops such as wheat, maize, banana and yerba mate. Within these isolates, the endophytic capability of the Gram-negative bacterium Kosakonia radicincitans DSM 166567 (syn. Enterobacter radicincitans) [25], formerly Pantoea agglomerans has been previously demonstrated [26]. K. radicincitans was isolated from the winter wheat leaves of the temperate regions [27]. Additionally, this facultative endophyte is known to stimulate the growth and yield of a range of plant hosts such as wheat (Triticum aestivum), maize (Zea mays) and radish (Raphanus sativus), among others [26,28–30]. The strain is able to fix atmospheric nitrogen biologically, solubilise calcium phosphate and to produce phytohormones such as auxins and cytokinins [31–33]. Finally, there already exists an efficient commercial product "AbiVital", which increases crop yield and even fruit quality parameters of radish and tomato plants [28,29,34].

The course of nutrients use including the phosphorus uptake emphasises the necessity to stimulate its availability during early plant growth stages and the endophytic establishment [33,35]. Thus, beyond the well-documented plant growth promoting benefits of this bacterial strain, and its inorganic phosphate solubilisation capability, the response of this endophyte to both the osmotic stress and the accumulation of compatible solutes, and how these conditions may influence its phosphate solubilisation ability and subsequent plant colonisation activity are still unknown. Here, we hypothesised that providing exogenous hydroxyectoine during the adaptation at high salinities in culture media may synergistically influence the phosphatase enzymes of K. radicincitans, enhancing its physiological machinery for phosphate solubilisation and subsequent plant colonisation activity.

Members of the Brassicaceae family are economically important crops, which the benefits of associated bacterial endophytes have tested successfully [36]. Previously studies demonstrated the ability of K. radicincitans to colonise internal tissue of Brassicaceae species such as radish (Raphanus sativus L. var. sativus) [29], independently of the site of application. Radish has features that allow the assessing of effects caused by environmental variables such as the short growth cycle (24–30 days), small size enabling a large number of treatments, easy determination of growth parameters and nutrient uptake from soil [37]. In particular, radish easily responds to the application of PGP and especially P-solubilizing bacteria with leaf and tuber growth improvement and P-uptake responses [38]. Therefore, this plant was selected as a model for testing plant growth-promoting and endophytic activity of pre-conditioned cells. Thus, this research investigates the effects of osmotic stress and hydroxyectoine accumulation in K. radicincitans DSM 166567 cells as endophytic bacteria model on the radish colonisation capability and P-solubilising activity.

Material and methods

Compatible solute standard hydroxyectoine (H-ectoine) was acquired from Sigma Aldrich (Cat: 70709, Sigma Aldrich Corporation, Darmstadt, Germany). All other materials used were of analytical reagent grade and used as received.

Bacteria and growth conditions

The bacterial strain K. radicincitans DSM 166567 was provided by the Leibnitz Institute of Vegetable and Ornamental Crops in Grossbeeren, Germany. Chemically defined growth medium (DM) was routinely used (g L−1): glycerol (15), yeast extract (8), K2HPO4 (2.74), KH2PO4 (1.31), MgSO4 7H2O (0.5), FeSO4·H2O (60 ppm), MnSO4 (10 ppm) at pH 7.4. Pre-conditioning of bacteria before plant colonisation assays was carried out by amending the DM with NaCl [1, 4%] and providing hydroxyectoine [1 mM] to DM 4% NaCl.

Bacterial suspensions for plant colonisation experiments were prepared as follows: DM (100 mL) was poured into 250 mL baffled Erlenmeyer flasks that were autoclaved at 121 °C under 1.5 atm, for 30 min. The initial inoculum concentration in the media was adjusted at 106 cells mL−1. The cultures were maintained at 190 rpm in a rotary incubator at 30 ± 1 °C (IKA KS 4000 IC Control, Staufen, Germany). Hydroxyectoine were sterilised separately by filtration through a 0.2 μm membrane filter (Durapore® 0.2 μm polyvinylidene fluoride (PVDF), Millipore, Darmstadt, Germany). Actively growing cells were harvested at the exponential phase after 20 h (OD600 of 0.7–0.9) by centrifugation at 6875g for 15 min (Centrifuge 5810R, Eppendorf, Wesseling, Germany), and the obtained pellet of bacteria was washed and centrifuged twice.
with corresponding NaCl solution [1, 4%] to maintain the osmotic pressure. The three bacteria type treatments [NaCl 1%, 4% and 4% + H-ectoine] were stored in the same NaCl solution adjusted to OD<sub>560 </sub>≈ 1.0 until use in plant colonisation assays. The intracellular hydroxyectoine concentration before endophytic establishments experiments was determined via HPLC [39].

Mineral phosphate solubilisation in liquid media

The efficiency of K. radicincitans of both pre-conditioned osmoadapted cells and osmoadapted hydroxyectoine-added cells for phosphate solubilisation were measured via Pikovkaya's (PVK) liquid media (pH 7.2 ± 0.2) [40]. PVK media were amended with 4% NaCl and 4% NaCl plus hydroxyectoine [1 mM] to reveal the effects of osmotic stress and the osmolyte on bacterial phosphate solubilisation ability. Each flask containing 100 mL of PVK was inoculated with 100 µL of bacterial suspension at 10<sup>8</sup> cells mL<sup>-1</sup>. The flasks were incubated for approximately nine days at 30 ± 1 °C under shaking at 190 rpm on a rotary shaker (IKA KS 4000 IC Control, Staufen, Germany). Four independent replicates per treatment were tested. Quantitative spectrophotometric analysis of the soluble phosphate was performed at 24 h, 48 h, 120 h and 200 h according to a standard protocol [41]. In parallel, the influence of osmotic stress and the addition of hydroxyectoine on pH evolution in PVK medium was monitored online, using microtiter plate cultivations (MPCs) (RoboLector-BioLector system, m2p-labs, Baesweiler, Germany). The following adjustments were used: Scattered light (620 nm filter, gain 3), pH-optode (Filter pH [HP8] Ex (nm) = 470 nm; Em (nm) = 525, gain 7), and 1000 µL of PVK media incubated in 48-well MTP-48-BH flower-plates, Lot No: 1808 at 30 °C and 1200 rpm.

Phosphatase enzyme activity

Phosphatase activity was determined using p-nitrophenyl phosphate disodium (PNPP, 0.025 M) as a colourimetric substrate. For the assay, 2 mL of 0.5 M modified universal buffer (MUB) buffer adjusted to pH 6.5 (acid phosphatases) and 11 (alkaline phosphatases), and 0.5 mL of the substrate were added to 0.5 mL of PVK supernatant medium. Cell-free supernatant samples were obtained by centrifuging 2 mL of culture at 21382 g for 10 min (Mikro HT 200R, Hettich GmbH & Co. KG, Tuttingen, Germany). Reactions were carried out at 37 ± 1 °C for 60 min and stopped by the addition of 0.5 mL of 0.5 M CaCl<sub>2</sub> and 2 mL of 0.5 M NaOH. Samples were filtered by using filter paper (grade 401) retention time 12–15 µm pore size. The p-nitrophenol (PNP) formed was measured spectrophotometrically at 400 nm (Genesys 10S UV-Vis, Thermo Fisher Scientific, Waltham, MA, USA) [42]. Four independent replicates per treatment were tested. Controls including the chemical hydroxyectoine were analysed under the same experimental conditions. The unit of enzyme activity (U) were expressed as micrograms of PNPP released per millilitre per hour (µg mL<sup>-1</sup> h<sup>-1</sup>) and normalised to dry biomass (mg) produced after 24 h.

Promotion of radish plant growth by osmoadapted K. radicincitans cells

The effects of both osmoadaptation and the incorporation of hydroxyectoine into K. radicincitans on plant growth promotion were tested under glasshouse conditions. Radish (Raphanus sativus L. var. sativus) seeds of cultivar Rondar (an F1 hybrid; S & G GmbH, Kleve, Germany) were used in all experiments. Radish inoculation with osmoadapted bacterial cells and hydroxyectoine-added cells was conducted by immersing radish seeds into a bacterial suspension (10<sup>8</sup> cells mL<sup>-1</sup>) for 5 min. Afterwards, 10 inoculated seeds were placed in pots (10 pots per treatment), filled with 1.5 L of a 1:1 (v/v) quartz-sand soil mixture (Fruhstorfer Erde type T25: P<sub>2</sub>O<sub>5</sub>: 200–300 mg L<sup>-1</sup>, Hawita Gruppe GmbH Vechta, Germany). The pots were then placed randomly on trivets to avoid transfer of bacteria between individual pots [29]. Seeds treated with NaCl solution [1, 4%] were used as controls. Seedlings were irrigated manually with 50 mL of tap water per day (conductivity 0.005–0.05 S/m). Plants were maintained under natural light conditions in the glasshouse at an average temperature of 18 ± 4 °C and an air humidity >45%. At one and four weeks post-planting, the plants were harvested.

One-week-old seedlings samples from three different locations per pot were taken, rinsed thoroughly with sterile water for removing soil with loosely adhering bacteria and flash frozen for nucleic acid extraction. Afterwards, the plants were equally thinned to five plants per pot. Total fresh mass of tuber and leaves material, as well as the tuber diameter of each plant were measured at four weeks old plants. The leaves were separated from the roots, and oven dried at 60 °C until constant weight, after which the dry weight of tubers and leaves were measured. The complete experiment was repeated twice.

Nucleic acid extraction and quantification of K. radicincitans in planta using qPCR

DNA was extracted from approximately 50 mg of lyophilised plant root material (one-week-old seedlings) using DNeasy Plant Mini Kit (Qiagen, Hilden GmbH, Germany) according to the manufacturer’s instructions. The lysis of bacterial cells was ensured by the addition of 5 mm sterile metal beads and by mechanical cell disruptor (Retsch MM200, Haan, Germany) at 30 rpm for 5 min. DNA quality and purity were assessed photometrically (NanoDrop, Thermo Fisher Scientific, Darmstadt, Germany). Quantitative real-time PCR (qPCR) was conducted using Advanced TM Universal SYBR<sup>®</sup> Green I Dye Supermix (Bio-Rad Laboratories, Hercules, CA, USA). K. radicincitans DSM 16656<sup>®</sup> species-specific primers and the plant TEF reference gene were used for in planta bacterial quantification [43]. The fold colonisation of K. radicincitans treated plants with respect to the reference gene and the control plants was calculated and represented with the 2<sup>−AΔAeq</sup> method [44].

Statistical analysis

The data were analysed using SPSS Statistics v.2 software (SPSS, Chicago, IL). Data are presented as mean values ± standard deviations (SD) or standard error (SE). The means were tested for significant differences by one-way analysis of variance (ANOVA) followed by the Tukey post hoc test. Repeated measures ANOVA was carried out for the phosphate solubilisation data. The level of significance was set at P < 0.05.

Results

Effects of osmotic stress on phosphate solubilisation capability

To look into the influence of osmotic unbalance on phosphate solubilisation capability PVK-amended liquid media with NaCl was used. Fig. 1A shows the positive effects on K. radicincitans ability to solubilise calcium phosphate in the PVK media under osmotic stress caused by 4% NaCl, compared with that in the PVK control media (NaCl 0.02%), solubilisation in the salt-amended media increased by up to 23.3%. The bacterial uptake of hydroxyectoine was detected by HPLC analysis after 15 h in response to high salinity, reaching more than 500 µmol g<sup>-1</sup> of dry biomass at 24 h. No hydroxyectoine was detected in cells grown in PVK in the absence of salt. Interestingly, repeated measures ANOVA indicated signifi-
cant differences \((F_{2, 9} = 28.229; P < 0.001)\) in response to the accumulation of hydroxyectoine on phosphate solubilisation during cultivation in PVK medium at 4% NaCl, and additional phosphate release increased by up to 9.39% after five days (Fig. 1A). Thus, \textit{K. radicincitans} can solubilise phosphate in a range of 150–400 mg L\(^{-1}\) after four days of cultivation. Online pH monitoring by the BioLector during the cultivation in PVK media, demonstrated a decrease in pH, with minimal pH values of 5.43, 5.66 and 5.82 recorded after 12.3 h, 21.6 h and 22.3 h for PVK control, PVK 4% NaCl with hydroxyectoine and PVK 4% NaCl media respectively. (Fig. 1B).

Phosphatase enzyme activity

Compared to the PVK control and the PVK 4% NaCl media, the addition of hydroxyectoine induced a positive effect on phosphatase enzyme activity. Acid phosphatase activity was highly induced by the presence of hydroxyectoine in the PVK media (4% NaCl) after 24 h \((F_{2, 11} = 12.80; P = 0.0029)\), 48 h \((F_{2, 11} = 70.67; P < 0.0001)\) and 120 h \((F_{2, 11} = 35.22; P = 0.0001)\) (Fig. 2A). Conversely, alkaline phosphatase activity was affected only after 24 h \((F_{2, 11} = 35.52; P = 0.0001)\), since at 48 h \((F_{2, 11} = 1.17; P = 0.3542)\) and at 120 h \((F_{2, 11} = 1.32; P = 0.3140)\), all treatments presented similar activities (Fig. 2B). Chemical control hydroxyectoine had not a detectable reaction for inducing P-solubilising.

Plant growth promotion in radish

Generally, plant growth promotion by non-osmoadapted and osmoadapted \textit{K. radicincitans} cells was observed in all inoculated radish plants. Interestingly, when hydroxyectoine at 1 mM was added during the cultivation of bacterial cells before plant seed inoculation, the fresh matter of tuber and leaves increased significantly by 41.1\% \((F_{3, 39} = 9.80, P = 0.0001)\) and 5.4\% \((F_{3, 39} = 3.86, P = 0.0172)\) in comparison to the non-inoculated control (Fig. 3A). Notably, compared with the osmoadapted cells in DM 4% NaCl, the hydroxyectoine amended cells increased the dry matter of either tubers or leaves by 16.20\% \((F_{3, 39} = 3.01, P = 0.0426)\) and 3.96\% \((F_{3, 39} = 3.60, P = 0.0672)\) respectively (Fig. 3B). In line with the plant weight increase, the tuber diameter also significantly increased in all cases of \textit{K. radicincitans} cells compared to the non-inoculated control \((F_{3, 34} = 6.70, P = 0.0013)\). The tuber diameter varied from 15.74 ± 1.82 mm in the native control up to 19.18 ± 1.41 mm for the hydroxyectoine pre-conditioned treatment (Fig. 3C).

\textit{K. radicincitans} plant colonisation

Regarding the relative gene expression response, compared with non-pre-conditioned bacteria cells at 1 NaCl, bacteria cells pre-conditioned with 4% NaCl colonised significantly tissue of eight-day-old seedlings \((F_{2, 14} = 10.803; P = 0.033)\), in comparison...
to non-pre-conditioned cells at 1% NaCl (Fig. 4). Consistent with the biomass production of radish plants, the endophytic colonisation was relatively strong in the presence of intracellular hydroxy-ylactone in *K. radicincitans* cells that were osmoadapted at 4% NaCl; the colonisation was approximately 3-fold greater under the evaluated treatment conditions ($F_{2, 14} = 10.803; P = 0.002$) (Fig. 4).

**Discussion**

Beyond the undeniable plant growth-promoting capability of *K. radicincitans*, results of this study suggest that it is feasible to significantly improve plant colonisation ability of bacterial endophytes, by physiologically growing them under conditions that promote the uptake or synthesis of advantageous compatible solutes such as hydroxy-ylactone. The bacterial ability to solubilise rock phosphate has been extensively reported as a vital mechanism that promotes plant growth [45,46]. In this study, it was demonstrated that *K. radicincitans* cells under osmotic stress could increase its phosphate solubilisation capability up to $357.7 \pm 9.38 \text{ mg L}^{-1}$ in comparison to non-stressed cells with 290.3 $\pm 42.6 \text{ mg L}^{-1}$ after four days of cultivation. Comparable yields are congruent with other pure culture experiments with Gram-negative bacterial endophytes [47]. Therefore, bacteria may respond to high salinity no only by accumulating exogenous osmo-lytes such as hydroxy-ylactone but also by up-regulating central metabolic pathways involved in the organic acid synthesis, which is well known for its role as crucial factors in phosphate solubilisation [48]. Since the pH values were lower in the treatment with hydroxy-ylactone compared to the treatment in which the bacterial cells were grown solely in PVK 4% NaCl media, a higher concentration of organic acids or the consumption of alkaline components within the media is feasible in conjunction with the osmo-adaptation process [33].

Widely distributed in microorganisms, plants and animals, phosphatases are hydrolases that catalyse the hydrolysis of orthophosphate monooesters thereby releasing phosphate [49]. The general classification of acid and alkaline phosphatases relies only on the optimum pH for enzymatic activity. A major production of enzymes may be correlated with the intracellular content of amino acids, since phosphatases are built on the basis of amino acid sequences and the phosphoryl group acceptors. It is expected that the physiological changes caused by either osmoadaptation or the addition of hydroxy-ylactone trigger metabolic pathways for induction of amino acids, either aromatic or branched-chain ones, which can favour enzyme biosynthesis and secretion. At high salinity, as a homeostasis response, trehalose biosynthesis in *K. radicincitans* cells may also lead to high concentrations of trehalose-6-phosphate phosphatase (T6PP), which catalyses the hydrolysis of trehalose 6-phosphate (T6P) to not only trehalose but also inorganic phosphate [50,51]. A greater production of enzymes, including phosphatases, may also be attributed to the construction of osmo-remedial mutations-epimutations in *K. radicincitans* cells, since some proteins are nonfunctional when the cells are grown in media of low osmotic strength but regain activity at elevated osmolarities, suggesting that cells may be undergoing phenotypic modulation [52,53]. Here the increase in acid phosphatase activity of the endophyte *K. radicincitans* exposed to osmotic shock in the presence of hydroxy-ylactone was evident. However, further research is needed to elucidate the types of phosphatases (nonspecific vs specific for certain substrates) that have been up-regulated in the production process.

Since osmo-adaptation in this bacterium leads to increased acid phosphatase enzyme production, it can be expected that this sub-lethal pre-conditioning procedure also supports the metabolite-
rich arsenal to face the extremely competitive rhizosphere conditions before entering the plant. It has been demonstrated with bacteria such as \textit{Pantoea agglomerans}, that osmotic stress can alter quorum sensing or the quenching of produced metabolites such as 1,3-propanediol, which are suggested to act as signals in the plant microbiome, resulting in tolerance to abiotic stress \cite{54,55}. Therefore, \textit{K. radicincitans} has a fraction of genes involved in propanediol degradation (PDD-pduABDEFLMPQ) and dha genes (dhaBDKLMt) involved in glycerol transformation (GT) \cite{56}.

\textit{K. radicincitans} cells were able to promote growth in radish plants. These findings are in line with previous studies \cite{29}, in which the weight of radish tubers and leaves increased from 20 to roughly 50\%, in response to either seed-inoculated or two-leaf sprayed plants with fresh cultivated cells. However, since in these pot experiments the majority of plant growth yields were not significantly different between \textit{K. radicincitans} cells grown in DM 1\% and DM 4\% NaCl, the physiological advantages conferred to the osmoadapted cells may affect mainly during early colonisation stages, as shown here for one-week old plants. Thus, the relative high endophytic lifestyle preference for salt-stressed bacteria cells and hydroxyectoine-added cells may be due to the alteration of signalling types of metabolites that \textit{K. radicincitans} can secrete into the microenvironments surrounding plant roots, modifying plant-defence and plant-competition mechanisms along with plant metabolite synthesis \cite{57}. Besides, external stimuli such as the presence of root exudates may up-regulate the expression of the chemotaxis and motility-related genes of these bacteria \cite{36}.

Consistent with the biomass production of radish plants, plant colonisation was stronger in the presence of intracellular hydroxyectoine within \textit{K. radicincitans} cells, in which the approximately 3-fold increase in colonisation was significant under the evaluated conditions. Generally, these results may indicate that symbiotic performance with plants increased by synergistic effects of pre-conditioned cells by osmoadaptation and the physiological changes caused by hydroxyectoine. However, despite the endophytic colonisation ability of the strain, it also growth and colonises the root surfaces \cite{58,59}. These cells may contribute to phosphorus solubilization and improved plant P-uptake which has been previously demonstrated \cite{33}. Nevertheless, additional studies are required to determine the performance of halotolerant bacterial endophytic cells on plant phosphorus uptake under saline conditions.

The reinforcement of \textit{in vitro} phosphate solubilisation capability, including the production of acid phosphatase enzymes and the accumulation of hydroxyectoine, also suggests an intracellular metabolic re-ordering within bacterial cells in response to high salinity. It is proposed that these substantial and significant alterations in metabolites levels represent the activation of a phenotypic shift as an osmoadaptation mechanism for conferring both survivals under such stress conditions and advantages during rhizosphere competition establishment, orchestrating nutrient exchange and mediating associations within the plant. To the best of our knowledge, the current study is the first to address the pre-conditioning of bacterial endophytic cells as an alternative to increasing plant colonisation abilities.

\section*{Conclusions}

The results of this study showed that physiological modifications of \textit{K. radicincitans} by osmotic stress treatments and by the accumulation of compatible solutes during cultivation constitute a feasible strategy to improve the ability of the bacterium to solubilise phosphate and to promote the root colonisation. Nevertheless, additional studies are needed for understanding the related mechanisms that are triggered by osmoadaptation and compatible solutes inclinations that favour endophyte-plant signalling. Moreover, pre-conditioning could be a promising alternative prior to formulating bacterial endophytes.

\section*{Conflict of interest}

The authors have declared no conflict of interest.

\section*{Compliance with Ethics requirements}

This article does not contain any studies with human or animal subjects.

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