Citramalic acid and salicylic acid in sugar beet root exudates solubilize soil phosphorus

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Abstract

Background: In soils with a low phosphorus (P) supply, sugar beet is known to intake more P than other species such as maize, wheat, or groundnut. We hypothesized that organic compounds exuded by sugar beet roots solubilize soil P and that this exudation is stimulated by P starvation.

Results: Root exudates were collected from plants grown in hydroponics under low- and high-P availability. Exudate components were separated by HPLC, ionized by electrospray, and detected by mass spectrometry in the range of mass-to-charge ratio (m/z) from 100 to 1000. Eight mass spectrometric signals were enhanced at least 5-fold by low P availability at all harvest times. Among these signals, negative ions with an m/z of 137 and 147 were shown to originate from salicylic acid and citramalic acid. The ability of both compounds to mobilize soil P was demonstrated by incubation of pure substances with Oxisol soil fertilized with calcium phosphate.

Conclusions: Root exudates of sugar beet contain salicylic acid and citramalic acid, the latter of which has rarely been detected in plants so far. Both metabolites solubilize soil P and their exudation by roots is stimulated by P deficiency. These results provide the first assignment of a biological function to citramalic acid of plant origin.

Background

Sugar beet and wheat are similar in their phosphorus (P) efficiency with regard to shoot production [1] but they appear to use different mechanisms to overcome low availability of soil P. Wheat has a large root system that compensates for low P influx when P availability is low, whereas sugar beet is able to achieve high P influx despite low P availability in soil [1]. The higher P influx of sugar beet compared to other plant species cannot solely be due to a more efficient uptake physiology. At low P availability, soil P transport is the limiting factor in P uptake [2]. Hence, the high P influx of sugar beet is attributed to the ability of the plant to mobilize, i.e. solubilize, P in the soil. This mobilization is most likely due to chemical modification of the rhizosphere by root exudates.

Plants exude up to 30% of assimilated carbon into the rhizosphere [3-5]. The composition of root exudates is complex and includes high molecular weight (HMW) and low molecular weight (LMW) molecules. HMW exudates include secreted enzymes and mucilage, which consists mainly of polysaccharides. Liebersbach et al. [6] showed that HMW exudates can increase P availability for plants, probably because carboxyl groups of polysaccharides interact with P-binding sites in the soil, which releases P into the soil solution. The long chains of polygalacturonate in HMW exudates may also cover soil particles and reduce the re-adsorption of phosphate [7]. Furthermore, the ability of HMW exudates to swell and absorb water may facilitate P diffusion toward the root [8]. LMW exudates include organic acids, sugars, phenolics, amino acids, phytosiderophores, flavonoids, vitamins and other compounds [4,5,9]. Phenolics might affect the speciation of iron (Fe) by complexation and thus might increase the availability of P occluded by Fe-oxides [10]. Organic acids, especially citrate, malate, and oxalate, are the root exudates most frequently investigated with regard to P mobilization.

P deficiency usually increases the root exudation rate and alters the composition of exudates [11,6], which results in an enhanced release of organic acids into the soil [12,13]. Hernandez [14] showed that the amount of organic acids was less in P-stressed roots than in P-sufficient roots of common bean; the reduced amount of organic acids in the P-deficient roots likely reflected exudation of organic acids...
from the root into the rhizosphere. These observations suggest that organic acids are involved in P acquisition.

Different mechanisms have been proposed for the facilitation of P acquisition by organic acids. For instance, citrate might increase the availability of P in the soil by binding calcium (Ca) and thus reducing the formation of insoluble complexes of Ca with P [15]. Citrate may also replace phosphate in complexes with Fe- and Al-oxides/hydroxides (ligand exchange). These processes release P into the soil solution as shown by Gerke et al. [16], who described a positive relation between citrate sorption and P-solution concentration. Furthermore, citrate can complex Fe, forming a citrate-Fe-P polymer that is soluble and can diffuse to the roots. The root reduces the Fe, thus breaking the polymer and releasing P directly at the root surface [17]. Hence, in soil incubation experiments with different organic acids, citrate often had the largest effect on P release.

It is still not clear, however, whether the beneficial effect of citrate and other organic acids on P availability actually occur in the rhizosphere. Most soil incubation experiments investigating the P-releasing effect of citrate were performed at relatively high concentrations of experiments investigating the P-releasing effect of citrate actually occur in the rhizosphere. Most soil incubation experiments investigating the P-releasing effect of citrate were achieved with the high P concentration (Figure 1). In the low P treatment, P concentration in shoot dry matter was 0.14-0.21%, which is much less than the range considered sufficient (0.35-1.10%, [22]), indicating that the plants grown under low P were P deficient. The high-P plants achieved P concentrations of about 1.5% in dry matter (Figure 1).

Exudation rate
At each harvest, root exudates were collected in a trap solution over a 2 h period. The trap solution was lyophilized, and the dry matter was weighed and was considered to indicate the quantity of root exudates. The exudation rate was 4-5 times greater in the P-deficient plants than in the well-supplied plants at all three harvests (Figure 2). The exudation rate remained constant over the whole growing period. A high exudation rate under P deficiency—especially a high exudation of carboxylates—has often been shown [13,16,23]; see also review of Jones [18].

The objective of this study was to use HPLC-ESI-MS to identify components of exudates of sugar beet roots that might increase P availability in the soil. Assuming that P-deficient plants would exude larger amounts of the compounds that solubilize soil P, root exudates were collected from sugar beet plants grown in nutrient solution with sufficient and with low P supply. Components of root exudates produced in larger amounts under low P supply were identified by differential metabolic profiling and were tested for their ability to solubilize P in soil.
The focus of this study was the difference in the composition of the exudates between low-P and high-P plants.

**HPLC-MS analysis of root exudates**

Full-scan HPLC-MS data were collected on root exudates as described in Methods section. The signals were normalized to compare low-P with high-P samples separately for each harvest. The normalized data were manually inspected to remove signals originating from solvent adducts and isotope peaks. Background peaks were removed that occurred in the controls. The profiles of high-P and low-P exudates were compared, and a total of 55 and 12 signals were detected in a negative and positive ionization mode, respectively, that were at least 5-times higher in low-P exudates than in high-P exudates (Table 1). Most of these signals, however, occurred only at a single harvest. Only 14 signals in negative mode were higher in exudates collected under low-P conditions than under high-P conditions in at least two harvests. Among these, ions [M-H]- with m/z of 137 and 147 were selected for further investigation because of their distinct MS signals and the high reproducibility of the signals among the harvests.

**Phosphorus mobilization in soil**

Candidate compounds were tested for the ability to solubilize P from soil. Six compounds were selected from the KEGG database [24] based on their molecular mass, the presence of carboxylic acids and commercial availability: 4-hydroxbenzoic acid, urocanic acid, salicylic acid, pantoic acid, D-arabino-1,4-lactone, and citramalic acid. Compounds with carboxylic acid groups were selected because the effect of organic anions on the mobilization of phosphate in soil is mediated by their functional groups [25], among which carboxyls are known to play a central role [26]. The results of P solubilization experiment are shown in Figure 3. Only salicylic and citramalic acid (molecular weight 138 and 148, respectively) increased P concentration significantly compared to the water control. The increase in concentration was by a factor of 2 and 6, respectively. The use of 1 mmol L⁻¹ solutions of candidate compounds in a ratio of 5 mL g⁻¹ of soil resulted in the addition of 5 μmol of compounds per g of soil. This quantity was based on the exudation rate of 22 μg m⁻¹ h⁻¹, and by assuming that this rate was maintained for 7 h, that the radius of the rhizosphere was 0.5 mm, and that the exudate had a molecular weight of 147. The calculated value was increased by a factor of 5, which corresponds to a higher exudation rate than actually measured. However, local concentration of root exudates is higher than the concentration in bulk soil solution. High concentrations of the candidates for P-solubilizing metabolites were also used to generate prominent, unequivocal effects. Fox et al. [27] used the same concentration (1 mmol L⁻¹) and the same soil-solution ratio (1:5) to compare the ability of organic acids to mobilize P in a Spodic Horizon.

**Identification of salicylic and citramalic acids in root exudates of sugar beet**

The mass spectrometric signals originating from root exudate components with putative molecular weights 138 and 148 were compared with the signals generated by pure salicylic and citramalic acid standards, respectively. For this comparison, samples originating from the same level of P supply for all three harvests were pooled to give a high-P pool and a low-P pool. A comparison of retention times and fragmentation patterns, based on reverse phase chromatography for salicylic acid and HILIC for citramalic acid (see Methods section), proved that the root exudate signal with an m/z value of 137 [M-H]- originated from

![Figure 2 Root exudation rates](image-url)

**Figure 2 Root exudation rates**. Rate of root exudation of sugar beet grown at low (2 μM) and high (500 μM) P is shown. Values originate from three harvest times and are given as means ± standard error of 3 replicates.

| Harvest | Negative ionization (m/z 100-400) | Negative ionization (m/z 400-1000) | Positive ionization (m/z 100-1000) |
|---------|----------------------------------|-----------------------------------|----------------------------------|
| 1st     | 35                               | 1                                 | 1                                |
| 2nd     | 12                               | 8                                 | 7                                |
| 3rd     | 15                               | 4                                 | 6                                |

The criterion for the inclusion of a signal in the list was a threshold of at least 5-times enhancement of signal intensity after normalization for low-P as compared to high-P conditions.

Table 1 Number of HPLC-MS signals in sugar beet root exudates enhanced under low-P conditions

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Khorassani et al. *BMC Plant Biology* 2011, 11:121
http://www.biomedcentral.com/1471-2229/11/121

Page 3 of 8
salicylic acid and that the signal with an m/z value of 147 [M-H]⁻ originated from citramalic acid. Salicylic and citramalic acids were present in root exudates originating from both low- and high-P treatments but concentrations of the two compounds were higher with the low-P than high-P treatment (Figures 4 and 5).

**Discussion**

Root exudates were collected from roots grown in nutrient solution which may differ from soil grown roots by, for example, the lack of root hairs. However, dense root hairs are formed in nutrient solution under conditions of P deficiency [28]; actually the effect of P deficiency on root hairs has often been studied on hydroponically grown roots [29]. It therefore appears safe to assume that the root exudation patterns caused by different P supply are comparable between solution and soil grown plants.

The ability of salicylic and citramalic acid to mobilize P from soil is related to their carboxyl and hydroxyl functionalities (Figure 6). The effect of organic anions on the mobilization of phosphate from soils and metal oxides increases with the number of carboxylic groups [26]. The relative position of hydroxyl and carboxyl groups is important [21] because it controls the ability of the substance to chelate Fe ions in soils where P is associated with Fe.

The effect of salicylic acid on P availability in soil has been studied by Staunton and Leprice [30], who showed
that salicylate increased the P concentration in the soil solution of a neutral calcic Luvisol. The competition between salicylic acid and P for sorption sites in two variable-charge soils was documented as well by Xu et al. [31]. On the other hand, Fox et al. [27] compared 16 organic acids and found no significant differences between salicylic acid and water in P solubilization. The discrepancy between their results and Staunton and Leprice [30], Xu et al. [31], and this study was likely caused by differences in the chemical form of P in the different soils.

Citramalic acid is known as a bacterial metabolite and an important chiral building block for synthetic pharmaceuticals, but it has rarely been detected in plants. Only three reports of citramalic acid in plants exist. It was found in apple peels 57 years ago [32], it was detected in tomato juice [33], and it was identified as one of uncommon metabolites during metabolic profiling of Arabidopsis thaliana [34]. Neither the biosynthetic pathway nor the physiological function of citramalic acids is known in plants. Our results provide the first putative assignment of a biological role to this uncommon plant metabolite.

We identified two among the eight mass spectrometric signals that were consistently enhanced in sugar beet root exudates under low-P conditions. Further work is needed to identify the remaining compounds and assess their ability to solubilize soil P.

**Conclusions**

The enhanced exudation of citramalic acid and salicylic acid by sugar beet roots under low P availability and solubilization of soil P by these metabolites strongly indicate that the function of citramalic acid and salicylic acid in root exudates is to increase the availability of P. While the presence of salicylic acid in root exudates and its function is established, citramalic acid has rarely been observed in plants before. Solubilization of soil P is the first biological role assigned to citramalic acid in plants.

**Methods**

**Hydroponic experiment**

Sugar beet plants were grown in hydroponic culture in a growth chamber. Sugar beet seeds were sown in sand and grown only with distilled water for 14 days. Seedlings of similar size were selected, washed free of sand with distilled water, and carefully transferred into pots containing 12 L of aerated nutrient solution. The experiment included two levels of P, three harvest dates, and three replicate pots for each combination of P level and harvest date. Plants were grown in a growth chamber at 16/8 h light/dark cycle, 25°C/16°C day/night temperature, 60%/75% relative humidity, and 41 W m⁻² PAR (photosynthetically active radiation during the day time). The composition of the nutrient solution was 1 mM Ca(NO₃)₂·4H₂O, 0.1 mM NH₄NO₃, 0.2 mM KCl, 0.1 mM MgSO₄·7H₂O, 17.9 μM
Fe-EDTA, 16 μM H$_3$BO$_3$, 9.1 μM MnCl$_2$, 4H$_2$O, 0.8 μM ZnSO$_4$, 7H$_2$O, 0.5 μM (NH$_4$)$_6$Mo$_7$O$_24$.4H$_2$O, 0.3 μM CuSO$_4$, 5H$_2$O. Phosphorus was added as NaH$_2$PO$_4$ in the concentration of 2 or 500 μM, designated as low-P and high-P, respectively. The number of seedlings per pot depended on the planned harvest date for that pot: 15, 10, and 6 plants were placed in pots intended for the first, second, and third harvest, respectively. Plants were harvested every 2 weeks after transplanting. The nutrient solution was replaced and pH was measured every second day. At each harvest, root exudates were collected (see next section), shoot dry weight was recorded, and the shoot P concentration was determined after digestion of 0.3 g of finely ground shoot material in 4 mL of HNO$_3$ and 2 mL of H$_2$O$_2$ under pressure at 175°C for 1 h in a microwave oven. P was determined by the molybdate-vanadate method of Scheffer and Pajenkamp [35]. Root length was determined with the line-intercept method [36].

Root exudate sampling

Whole root systems of intact plants were carefully washed with running de-ionized water to remove the nutrient solution. For collection of exudates, the whole root system was dipped into aerated de-ionized water in a glass container, the volume of which depended on the size (age) of the root system. The container was covered with aluminium foil to create dark conditions for roots. The roots were kept in the water for 2 h under the same controlled climate conditions described for plant growth. Röhmheld and Neumann [23] showed that for a short exudation period of 2 h, the roots were not harmed by the de-ionized water and no significant microbial degradation of the exudates occurred [23]. The collected root exudates were immediately frozen at -30°C, freeze dried, and weighed.

Preparation of samples and standards for HPLC-MS analysis

A 0.5-mg sample of each freeze-dried root exudate was weighed into an HPLC vial, and 100 μL of acetonitrile was added. After 30 min, samples were vigorously shaken. Then 900 μL of HPLC-quality water was added, and the samples were shaken again and filtered through Teflon membrane filters (0.2-μm pore size, 13 mm diameter, Optiflow-TF, Wicom Germany GmbH, Heppenheim, Germany). Standards of citramalic acid and salicylic acid were prepared as 10 mg/mL stock solutions in methanol:water (50:50 v/v) and diluted to 100 ng/mL in mobile phase used for column equilibration.

Metabolic profiling of root exudates

For metabolic profiling of root exudates, an HPLC system coupled to an ion trap mass spectrometer was used. HPLC separation was carried out as described [37]. Mass spectrometry was performed with the 500-MS LC ion trap equipped with an electrospray ion source (Varian, Darmstadt, Germany). Nebulizer and drying gas pressures were set to 50 and 20 psi (345 and 138 kPa), respectively. Drying gas temperature was set to 350°C at the beginning of the gradient and was reduced gradually to 250°C with the increasing proportion of methanol in the eluate. For the detection of positive and negative ions, needle voltages were set to 5000 and -3500 V and shield voltages to 600 and -600 V, respectively. The capillary voltage was +/-50 V. In positive ionization mode, ions with an m/z of 100 to 1000 were collected in a single run, while in negative mode ranges of m/z 50-400 and m/z 400-1000 were scanned separately. The scan rate was 5000 Da/s, and three consecutive scans were averaged. MS data were transformed into chromatograms using MS Data Review 6.9 (Varian) and converted into netCDF format. Data from positive and negative ionization modes were separately processed as follows: minor differences in retention times were corrected by peak alignment performed with XCMS version 1.5.2 [38] run under R package 2.4.0. The resulting data were normalized to compensate for differences caused by uneven preparation of the samples as described previously [39].

Identification and analysis of selected metabolites

HPLC-MS-MS detection of specific compounds was performed on an identical HPLC system with a triple quadrupole mass spectrometer (1200 L, Varian). Chromatography was performed on a polar-modified RP-18 phase and on a HILIC (hydrophilic interaction chromatography) phase. Root exudates and test substances were separated under identical conditions. RP-18 chromatography was conducted as described before [40]. For HILIC, the buffer system consisted of (A) 25 mM ammonium acetate in water and (B) acetonitrile. A 10-μL volume of sample was loaded onto a ZIC-HILIC (Sequant, Haltern am See, Germany) equilibrated with 95% B and separated in a linear gradient from 95-10% B in 10 min. The column was kept at 40°C and the flow rate was 0.2 mL/min. Salicylic acid was identified by the mass transition 137 > 93 [M-H-CO$_2$] and a retention time of 7.15 min on the RP-18 phase. Citramalic acid was identified by the mass transition 147 > 87 [M-H$_2$CO$_2$] and a retention time of 9.2 min on the HILIC phase.

P solubility experiment

Solutions of 4-hydroxy benzoic acid (Fluka, Germany), urocanic acid (Fluka, Germany), salicylic acid (Merck, Germany), citramalic acid (Aldrich, Germany), D-arabino-1,4-lactone (Dextera, UK), and pantoic acid (generated from pantolactone purchased from Aldrich, Germany, by hydrolysis in 0.1 M NaOH) were prepared in water at the concentration of 1 mmol L$^{-1}$. pH of the solutions was...
adjusted to 5.6 with NaOH or HCl. Highly P fixing fossil Oxisol, containing mainly P bound to Al and Fe with 788 mg P kg⁻¹ as Fe/Al-P and 330 mg P kg⁻¹ as Ca-P, clay content 50%, pH(CaCl₂) 5.6, P concentration in soil solution 0.17 μmol L⁻¹ and Ca acetate-lactate (CAL) extractable P of 4.3 mg kg⁻¹ was fertilized with 100 mg P kg⁻¹ as Ca(H₂PO₄)₂·H₂O and moisturized to 22% w/w water content. After 10 days of equilibration, 9.76 g of moist soil (equivalent to 8.0 g of dry soil) was mixed with 40 mL of the solutions of the compounds listed above or with distilled water. Microbial degradation was prevented by adding two drops of toluene. Each treatment was replicated three times. The samples were shaken on a reciprocal shaker at 150 cycles per minute for 5 h, centrifuged for 15 min at the relative centrifugation force of 2660 g, and the supernatant was filtered through a 0.45-μm nylon membrane filter. Inorganic P concentration was determined by a molybdenum blue colorimetric method [41].

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Authors’ contributions
RK performed hydroponic experiments and P solubilization tests. UH and AR performed metabolic profiling and analysis of root exudate components. BS designed and supervised the P solubilization experiment. PK guided performed metabolic profiling and wrote scripts for data processing. NC conceived the study. All authors evaluated the data, wrote parts of the manuscript and approved the manuscript for submission.

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