The effect of pH on morphological and physiological root traits of *Lupinus angustifolius* treated with struvite as a recycled phosphorus source

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Abstract

**Aims** Phosphorus (P) recovery from specific waste streams is necessary to develop environmentally sustainable and efficient fertilizers, achieving maximum productivity with minimum losses. A promising example of a P-recovery product is struvite (MgNH₄PO₄·6H₂O). Phosphorus availability from struvite is profoundly influenced by soil pH and/or processes in the rhizosphere. Root exudates (e.g., organic anions) and root morphology affect fertilizer bioavailability. The overall objective of our study was to identify root morphological and physiological traits of the narrow-leaf lupine (*Lupinus angustifolius* L. subsp. *angustifolius*, cultivar: blau “Boregine”) involved in the acquisition of P from struvite, compared with KH₂PO₄ as a soluble P source. The study included different pH conditions, as soil pH is one of the main factors affecting P availability.

**Methods** Narrow-leaf lupine plants were grown in river sand under three pH conditions (4.5, 6.5 and 7.5). Three different P treatment conditions were used: 1) KH₂PO₄ (KP); 2) MgNH₄PO₄·6H₂O (Struvite), both supplied at 15 μg P g⁻¹ dry sand; and 3) no P addition (Nil-P), as control. Organic acids in the rhizosheath were collected. Root morphological parameters such as specific root length and root diameter were analyzed.

**Results** There was no significant difference in total plant biomass detected under any pH condition between struvite and KP treatments. In both acidic and alkaline conditions, the P-uptake efficiency (PUE: mg P plant⁻¹/cm² root surface area) with struvite was significantly greater than with KP. At neutral pH, there was no difference in PUE between plants supplied with KP or struvite. Plants growing at neutral pH showed greater root exudation of carboxylates (mainly citrate) when struvite was added compared with KP. At alkaline pH, the exudation per unit root surface area was greater than that at acidic or neutral pH. Plants growing in acidic pH had a higher specific root length (SRL) compared with those grown at alkaline or neutral pH.

**Conclusions** Similar P-uptake efficiency from struvite and KH₂PO₄ at neutral pH in conjunction with the higher total biomass compared to the Nil-P treatment (70% higher) suggests very effective mobilization of P from struvite by carboxylate exudation. Application of struvite, while taking into account the different
strategies for nutrient mobilization, can increase the use efficiency of this recovered P source.

**Keywords** Carboxylate exudation · Fertilizer-use efficiency · Struvite · Recycled phosphorus · Rhizosphere pH · Root diameter · Specific root length

**Abbreviations**

- DCL: Diameter class length
- DM: Dry mass
- Nil-P: No phosphorus treatment
- P: Phosphorus
- PUE: Phosphorus uptake efficiency
- SRL: Specific root length

**Introduction**

Phosphorus (P) is an essential macro-element and plays a crucial role in many key processes (Jouhet et al. 2007; Misson et al. 2005). The total amount of P in soil is generally high after fertilization, but a large portion is not in solution and thereby not directly available to plants. Recovery of the P supplied by fertilizers is thereby a slow process which takes years of cropping. In the first year after application, crops often take up less than 30% of the applied P (Vance et al. 2003). This is due to precipitation reactions with Ca, Fe, Al and Mn as well as sorption to clays, Fe or Al oxides and hydroxides present in soil, thus making P unavailable for plants (Shen et al. 2011). Furthermore, the conversion of P into organic forms (immobilization by the microbial community) (Balemi and Negisho 2012) may also further constrain the P availability to plants (Delgado and Scalenghe 2008). This is because the bioavailability of P from organic pools is mainly determined by the susceptibility of these compounds to plant and microbial phosphatases (Sanyal and De Datta 1991).

Application of large amounts of mineral P fertilizers derived from rock phosphate is the most common way to enhance plant-available P in regions with P-deficient soils. However, as a non-renewable natural resource, global rock phosphate reserves are being depleted (Cordell and White 2011). Therefore a move towards a more efficient and sustainable fertilization requires a change in the P stewardship (Fixen and Johnston 2012) by identifying alternative P sources and evaluating their potential as P fertilizers.

A promising example of an alternative source of P investigated in this study is struvite (MgNH₄PO₄·6H₂O), a precipitate recovered from municipal wastewater streams that is free of contaminants and has a stable nutrient composition. Struvite has previously been proposed as a slow-release fertilizer (Rahman et al. 2014). The slow release can ensure a steady nutrient supply for plants which might promote root development in the early stages and improve nutrient-use efficiency (defined by Blair (1993) as the ability of a genotype to acquire nutrients from growth medium and/or utilize them for the production of shoot and root biomass), consequently stimulating plant growth.

A key factor for the management of struvite and similar recycled P sources is to increase the bioavailability and plant P-uptake efficiency (uptake per unit of root surface area) under different soil conditions. Struvite fertilizers can increase biomass in maize (Zea mays) and narrow-leaf lupine (Lupinus angustifolius) in acidic conditions (Robles-Aguilar et al., unpublished results). However, contrary to the highly soluble commercial P fertilizers (i.e., potassium phosphate – KP), which are bioavailable over a broad pH range, uptake of P from struvite is slow at neutral to alkaline pH (Bhuiyan et al. 2007; Massye et al. 2009). Therefore, knowing how to increase struvite P-use efficiency at different soil pH values will make its use more reliable as a substitute for rock phosphate. This will improve the management of recycled P sources which is desirable from both an economic and an environmental perspective (Manske et al. 2001; Shenoy and Kalagudi 2005).

Plants have evolved a number of strategies to improve their P-acquisition efficiency (Lambers et al. 2006; Raghothama and Karthikeyan 2005). Three main strategies are i) changes in the relative investment in root to shoot biomass, ii) root-morphological changes that improve P acquisition by increasing the soil volume that is explored (Lynch 2007), and iii) P-mining strategies to enhance the solubility of P from sparingly-available sources by root exudates (organic anions) (Richardson et al. 2011). Significant morphological traits that affect P acquisition include the increase of root surface area through higher root branching density and longer root hairs that enable foraging for plant-available P and its uptake from the soil solution (Hammond et al. 2004). But also changes in root architecture including growth of roots in regions with higher P availability are important strategies (Postma et al. 2014). One noteworthy physiological adaptation to increase the P recovery is
the exudation of carboxylates (low molecular weight organic anions) into the rhizosphere when plants experience P deficiency (Richardson et al. 2011; Veneklaas et al. 2003). Most likely, plants can adjust at both levels by carboxylate exudation, as well as modifying root morphology (Pearse et al. 2006). Narrow-leaf lupine is well known for its ability to grow well in sandy acidic soils that contain low P concentrations. This is attributed to its high capability to exude carboxylates, allowing the mobilization of sparingly soluble P sources in the soil (Pearse et al. 2006). However, no proteoid roots are produced by narrow-leaf lupine, implying that the exudation of carboxylates is not associated with the formation of proteoid roots. Hence, we hypothesized that narrow-leaf lupine can compensate for the low availability of struvite-P at neutral to alkaline conditions not only through carboxylate exudation, but also through specific root morphological adaptations. Therefore, we evaluated if narrow-leaf lupine produces more biomass when struvite is supplied as a P source in sandy soil under different pH conditions ranging from acidic to alkaline. The strategy of growing P-efficient crops with appropriate concentrations of P via a nutrient solution to a final concentration of 15 \( \mu \)g P g \(^{-1}\) dry sand; and iii) no P addition (Nil-P), as the control. The supply of appropriate concentrations of \( \text{NH}_4\text{NO}_3 \) and \( \text{K}_2\text{SO}_4 \) to the P treatments evened out the nitrogen and potassium provided by the struvite and the \( \text{KH}_2\text{PO}_4 \) treatment respectively. The pH of the nutrient solution was adjusted from the original pH 4.5 to a neutral solution (pH 6.5) by adding 10 ml KOH (200 mM) L \(^{-1}\) solution, and to an alkaline solution (pH 8) by adding 25 ml KOH (200 mM) L \(^{-1}\) solution, in order to maintain the desired pH in the sand after its application. The sand was initially moistened to 80% of field capacity with nutrient solution, which was adjusted to the respective pH. Thereafter, soil moisture was maintained for the duration of the experiment by watering pots to a specific weight with deionized water every other day. Watering was controlled to make sure that no water leached from the bottom of the pots.

Three seedlings of narrow-leaf lupine (\( \text{Lupinus angustifolius} \) \( \text{L. subsp. angustifolius} \), cultivar: blau "Boregine", Kiepenkerl, Germany) were planted in each pot and thinned to one plant per pot after 1 week. Each pot was inoculated with 1 g of peat-based Group G Rhizobium solution (New Edge Microbials, Albury, NSW, Australia). The inoculation was done by applying a few drops to the seedlings using a plastic syringe containing a mixture of rhizobia in peat and water. There were nine treatment combinations in the study, with five replicates per treatment. In addition, two pots without plants were used as control for each treatment to monitor the sand pH development during

**Materials and methods**

**Plant cultivation**

The experiment was carried out during 6 weeks from February to March 2016 in a temperature-controlled glasshouse under natural light conditions at the University of Western Australia, Perth, Australia (31.57°S, 115.47°E) with an average daytime temperature of 24 °C and average nighttime temperature of 21 °C, and a mean relative humidity of 65%. Pure coarse river sand was washed, sterilized and air-dried. It was found to it contained trace amounts of nutrients as indicated by Pang et al. (2010) with 3 \( \mu \)g g \(^{-1}\) bicarbonate extractable P (Colwell 1963) (16% of the total P after fertilization), and had an initial pH of 6.3 analyzed in a solution of 0.01 \( M \) CaCl\(_2\) (Ahern et al. 1995). To obtain a pH of 4.5 and 7.5, the sand was mixed with 3 g iron sulfate kg \(^{-1}\) sand (FeSO\(_4\)) or 50 g calcium carbonate kg \(^{-1}\) sand (CaCO\(_3\)) in an end-over-end shaker for 15 min at 99 rpm, respectively; these amounts were determined beforehand to reach the desired pH. Pots (8.5 \( \times \) 8.5 \( \times \) 18 cm) were filled with 1.3 kg of sand with pH of either 4.5, 6.5 or 7.5. To ensure that the nutrient supply was adequate for plant growth for the duration of the experiment, all essential nutrients other than P were provided via a nutrient solution to a final concentration of 30 N, 40 S, 24 Ca, 11.5 Mg, 0.5 Cu, 5 Fe, 55 Cl, and 80 K, all values given in \( \mu \)g g \(^{-1}\) dry sand. In order to evaluate the P recovery from struvite, we used \( \text{KH}_2\text{PO}_4 \) (KP), a soluble form of P as the control. Three P treatments were applied: i) \( \text{KH}_2\text{PO}_4 \) was added as a nutrient solution; ii) struvite (\( \text{MgNH}_4\text{PO}_4\cdot6\text{H}_2\text{O} \)) in powder form was mixed with sand, both to a final concentration of 15 \( \mu \)g P g \(^{-1}\) of dry sand; and iii) no P addition (Nil-P), as the control. The supply of appropriate concentrations of \( \text{NH}_4\text{NO}_3 \) and \( \text{K}_2\text{SO}_4 \) to the P treatments evened out the nitrogen and potassium provided by the struvite and the \( \text{KH}_2\text{PO}_4 \) treatment respectively. The pH of the nutrient solution was adjusted from the original pH 4.5 to a neutral solution (pH 6.5) by adding 10 ml KOH (200 mM) L \(^{-1}\) solution, and to an alkaline solution (pH 8) by adding 25 ml KOH (200 mM) L \(^{-1}\) solution, in order to maintain the desired pH in the sand after its application. The sand was initially moistened to 80% of field capacity with nutrient solution, which was adjusted to the respective pH. Thereafter, soil moisture was maintained for the duration of the experiment by watering pots to a specific weight with deionized water every other day. Watering was controlled to make sure that no water leached from the bottom of the pots.
the experiment. Pots on the greenhouse bench were randomized every second day.

Rhizosheath carboxylates extraction

After 6 weeks, the plants were removed from the pots, and the root systems were shaken gently. The sand that was attached to the roots was defined as the rhizosheath (Pang et al. 2017). Rhizosheath extracts for the analyses of carboxylates were collected according to Pang et al. (2010). The entire root system was submerged in 0.2 mM CaCl2 and shaken to remove the rhizosheath sand. The volume of CaCl2 needed to cover the whole root system varied according to the root size from 20 to 100 ml. Subsamples of the extracts were taken and filtered through a 0.22 μm syringe filter. One ml of the filtered extract was transferred into a 1 ml high-performance liquid chromatography (HPLC) vial. HPLC analysis of the extracts was performed as described by Cawthray (2003) using working standards of malic, malonic, lactic, acetic, maleic, citric, succinic, cis-aconitic and trans-aconitic acid (ICN Biomedicals Inc., Aurora OH, USA) to identify and quantify carboxylates.

Plant measurements

Immediately after collection of carboxylates, plants were separated into shoots and roots, and leaf area was measured using a leaf area meter (LI-3100, LI-COR, Lincoln, Nebraska, USA). Subsequently, cleaned roots were scanned with a transparent adapted tray (acrylic box) positioned in the scanner (Epson Expression Scan 1680, Long Beach, Canada). Roots were spread out in the acrylic box (size A4) containing tap water to minimize the number of overlaps. Based on the scanned images of the root systems, total length, mean diameter, total surface area, and root volume were measured using the WinRHIZO™ regular V.2009 software (Regent Instruments Inc., Quebec, Canada). The root length measurements were partitioned into five diameter classes: <0.5, 0.5–1, 1–1.5, 1.5–2, and > 2 mm. The following parameters were based on observed and/or computed data: root-to-shoot ratio (root dry mass/shoot dry mass), specific root length (SRL) = root length/root dry mass (m g−1) and relative diameter class length (rDCL) = DCL/total root length (yielding a proportion of root length to normalize disparity between plant sizes). Visual scoring of nodulation was done at harvest using the Field guide for nodulation and nitrogen fixation (BC Ministry of Forests 1991). Shoot and root dry mass was recorded after drying for 48 h at 65 °C in a forced-draft oven.

Phosphorus analyses

To determine P concentrations, dried shoot and root samples were milled using a stainless steel ball mill. Weighed subsamples of the fine powder (200 mg) were digested using a hot concentrated nitric-perchloric (3:1) acid mixture. Total P concentrations in the shoot and root samples were determined by the malachite green method (Motomizu et al. 1983) using a UV-VIS spectrophotometer (Shimadzu Corporation, Japan). Phosphorus-uptake efficiency (PUE) was calculated based on Hammond et al. (2009), who described PUE as the increase of plant total P content per unit of added P fertilizer (g P plant−1 g−1 P fertilizer). We modified the formula, normalizing the P uptake per unit of root surface area instead of per amount of P applied (PUE: mg P plant−1 cm−2 root surface area).

Statistical analyses

Analyses were performed using the statistical program R.2.16.3 (R: A Language and Environment for Statistical Computing (2012) http://www.R-project.org/). Measurements were compared with two-way analysis of variance (ANOVA). Tukey’s HSD post hoc test after ANOVA at a significance level of p < 0.05 was used to check which level of a factor differed from one another. Data were calculated as arithmetic means ± standard error of the mean of the indicated replicates. Statistical analyses for slopes were conducted by analyses of covariance (ANCOVA). Details of all statistical analyses and data are provided as an electronic supplementary material (ESM) file.

Results

The effect of struvite-P fertilizer on shoot growth

When P was supplied, plant growth and shoot biomass varied significantly among the pH conditions (Fig. 1a, exemplary for struvite fertilization). The ANOVA showed significant effects (p < 0.05) of both factors, P treatment and pH, as well as their interactions (p < 0.05) in shoot biomass (electronic supplementary material,
ESM Tables 1 and 2). The similar effectiveness of struvite compared with KP was confirmed at neutral (pH 6.5) and acidic (pH 4.5) conditions (Fig. 1a as exemplary for neutral pH, and Fig. 2). Under both pH 6.5 and 4.5, shoot biomass did not differ between struvite- and KP-treated plants. Shoot biomass under both P treatments was greater than that in the Nil-P treatment \( (p < 0.05) \) (Fig. 2). At alkaline conditions, there was no difference between P-treated plants and the Nil-P treatment. Leaf area showed a similar trend to that of shoot biomass.

Both high and a low soil pH significantly reduced plant growth compared to plants growing at neutral pH \( (p < 0.05) \). At pH 7.5, the total biomass was 80% less compared to growth at neutral pH. At pH 4.5, the biomass was 65% less compared to growth at neutral pH 6.5, which was the optimum pH for narrow-leaf lupine.

The allometric analyses offers a visual description of root and shoot biomass distribution, which is not confounded by plant size. Figure 3 shows the root:shoot relationship under different pH values and P sources. ANCOVA analyses showed significant differences in slope \( (p < 0.05) \) between the Nil-P treatment and the other P sources, as well as between the alkaline pH compared with acidic and neutral pH (ESM Tables 7 and 8). This allocation pattern shows that the plants invested relatively more in roots when the pH of the sand was suboptimal for growth (alkaline) or when no P was applied (Nil-P) (Fig. 3). The results indicate that the effect of struvite was similar to that of KP–P in terms of allocation, as well as total biomass.

![Fig. 1 a Effects of pH on narrow-leaf lupine growth under struvite treatment in acidic (pH 4.5, left), neutral (pH 6.5, middle), and alkaline (pH 7.5, right) sand. b Effects of different phosphorus sources applied on narrow-leaf lupine growth at neutral pH treated with no phosphorus (Nil-P, left), struvite (middle), and potassium phosphate (KP, right)](image)

![Fig. 2 Plant biomass of narrow-leaf lupine grown with different phosphorus (P) sources: struvite (S), potassium phosphate (KP) and no P (Nil-P) applied at different sand pH (acidic 4.5, neutral 6.5 and alkaline 7.5). The line that divides the box represents the median of the data. The end of the box shows the upper and lower quartiles. The extreme lines out of the box show the highest and lowest value excluding outliers (shown as dots); n = 5](image)
Plant P-uptake efficiency as affected by pH and phosphorus source applied

Total P content (mg P plant$^{-1}$) increased corresponding with the solubility of the P source (Table 1). As a result, the highest total P content was found in plants treated with KP, which is the most soluble P source, followed by struvite and Nil-P, which is further reflected by biomass production (Fig. 2). The highest plant total P content and plant growth was observed at neutral pH. Plant P concentration (mg P g$^{-1}$ dry mass) showed a different pattern (Table 1). Plants treated with struvite had a higher P concentration than those treated with KP (significant only at alkaline pH). In contrast to plant biomass, plants growing at alkaline pH had the highest P concentration, followed by those at acidic and neutral pH (Table 1). Further statistical data and analyses are provided as electronic supplementary material, ESM Tables 9 and 10.

The P-uptake efficiency was calculated to compare the efficiency of the plants to take up P per unit of root surface area, as an indicator of other physiological processes occurring in the root that might increase the efficiency of P uptake.

ANOVA analyses indicated significant effects of P treatment and pH on PUE ($p < 0.05$). At neutral pH, there was no difference in PUE between KP and struvite. In acidic and in alkaline conditions, PUE from struvite was higher than that from KP (Table 2). At alkaline pH, the PUE was higher than that in the other pH conditions. Further statistical data and analyses are provided as electronic supplementary material, ESM Tables 13 and 14.

Root morphology

The root system, in general, was dominated by the taproot and primary lateral roots (first-order branches); however, significant differences in the rooting pattern were observed among pH conditions (Fig. 4).
Narrow-leaf lupine plants showed significant variation in total root length and root surface area in response to pH and P applied \((p < 0.05)\). When plants grew in neutral conditions, total root length and root surface area were greatest with struvite, followed by KP, and the lowest root length and surface area were found when no P was applied (Table 3). In acidic conditions, plants supplied with KP had almost twice more total root length compared with those supplied with struvite \((p < 0.05)\). In alkaline pH, secondary roots did not elongate (not growing more than 1.5 mm from the major axes in most cases), and surface area was reduced by 90% compared with roots growing at pH 6.5 in all the P treatments (Table 3). Nodulation was also inhibited at alkaline pH (data not shown).

The average root diameter in acidic conditions with struvite was significantly lower compared to KP. For every diameter range, the relative diameter class length was calculated for each pH (rDCL). Values are shown in Table 4 and represent the percent of the total length that belongs to each root diameter class in each particular pH. At acidic pH, thin roots (<1 mm) accounted for almost 80% of total root length, compared with the percentage of thin roots at alkaline (57%) and neutral (71%) pH. In alkaline conditions, the percentage of thick roots (>2 mm) increased up to 9%, significantly more than the percentage calculated for neutral (3.5%) or acidic (2.7%) conditions.

The specific root length (SRL) was significantly affected by pH \((p < 0.05)\), but not by the P treatment \((p = 0.06)\). The highest SRL was found in acidic conditions, with KP, followed by struvite and lowest in Nil-P treatment. The SRL reached its minimum when soil was alkaline, being reduced from 2.4 m g\(^{-1}\) root with KP to 2.0 m g\(^{-1}\) root with no P addition (Table 3). Further statistical data and analyses are provided as electronic supplementary material, ESM Tables 15 to 22.

**Carboxylate release**

At neutral pH, the total amount of carboxylates (μmol plant\(^{-1}\)), mainly citrate and malate, measured in the rhizosheath of narrow-leaf lupine was significantly increased with struvite treatment in comparison with KP treatment. Under acidic conditions, exudation of carboxylates was less than at neutral pH, with no significant differences between P sources. At acidic pH, the amount of carboxylates measured was even slightly greater with KP compared with that in the struvite treatment. The carboxylate amount per unit root surface area (mmol m\(^{-2}\) root) was greater as the pH was increased from 4.5 (acidic) to 7.5 (alkaline) (Fig. 5). At neutral pH (6.5) the exudation was greater in the struvite treatment compared with that in the KP treatment. At alkaline pH, root growth was strongly inhibited, and the total amount of carboxylates was strongly reduced, but the carboxylate amount per unit root surface area was greatest (Fig. 5).

**Phosphorus source, pH, and their interaction had a significant effect on the total amount of carboxylates in the rhizosheath \((p < 0.05)\) and in the amount per unit of root surface area. Roots modified the exudation of carboxylates, mainly citrate, in response to the different P sources applied, which in turn was affected by the sand pH.**

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**Table 1** Shoot and root phosphorus concentration (mg P g\(^{-1}\) dry mass, DM) of narrow-leaf lupine plants treated with potassium phosphate (KP), struvite or no phosphorus application (Nil-P) growing at acidic (pH 4.5), neutral (pH 6.5) and alkaline (pH 7.5) conditions in sand; mean values \((n = 5)\) with the same letter across shoot or root biomass are not significantly different based on Tukey Honest Sign. Diff. test \((p < 0.05)\)

| P source | Acidic | Neutral | Alkaline |
|----------|--------|---------|----------|
| Shoot    |        |         |          |
| Struvite | 11.8 b | 5.6 b   | 22.4 a   |
| KP       | 6.6 b  | 4.6 b   | 10.2 b   |
| Nil-P    | 8.3 b  | 7.9 b   | 12.7 b   |
| Root     |        |         |          |
| Struvite | 17.9 a | 6.6 c   | 15.4 ab  |
| KP       | 6.1 c  | 6.7 bc  | 9.1 bc   |
| Nil-P    | 9.6 abc| 7.4 bc  | 8.1 bc   |

Shoot and root values were tested separately. For ANOVA see ESM Table 9 to Table 12

**Table 2** Phosphorus uptake efficiency (PUE) from struvite, potassium phosphate (KP) and no phosphorus (Nil-P) calculated for narrow-leaf lupine plants growing at acidic (pH 4.5), neutral (pH 6.5) and alkaline (pH 7.5) conditions in sand depended on an interaction between P treatment and soil pH. PUE: mg P plant\(^{-1}\) cm\(^{-2}\) root surface area

| P source | Acidic | Neutral | Alkaline |
|----------|--------|---------|----------|
| Struvite | 308ab  | 162c    | 1546a    |
| KP       | 121b   | 175c    | 531a     |
| Nil-P    | 219ab  | 311ab   | 620a     |

Values represent the mean of \(n = 5\). For ANOVA see ESM Table 13
Rhizosheath pH measurements

The rhizosheath pH was significantly affected by the initial pH condition \((p < 0.05)\). In alkaline conditions, the pH of the rhizosheath of Nil-P plants was higher \((8.1)\) than that of struvite- and KP-treated plants \((8\) and \(7.8,\) respectively). Those pH values were higher than the initial pH values of the sand at alkaline condition, monitored in the non-planted control pots \((7.5)\). In neutral conditions, pH of the Nil-P was slightly higher \((6.0)\) compared to struvite \((5.6)\) and KP \((5.5)\). In this case, the pH measured in the rhizosheath was lower than the initial pH measured in neutral condition \((6.5)\). In contrast to what observed in alkaline and neutral pH, struvite treatment in acidic conditions slightly increased the pH \((4.9)\) in comparison with KP \((4.2)\) and Nil-P \((4.5)\). However, no significant difference in rhizosheath pH \((p = 0.2)\) was found between different P sources.

Discussion

Root morphology in response to pH and P source

The root morphology of narrow-leaf lupine was markedly altered by the soil pH. High pH inhibited root elongation, with a reduction of up to 90% in surface area between plants growing at pH 7.5 and pH 6.5,
comparable to results previously observed by Tang et al. (1992). Plants growing in acidic pH had a greater SRL than plants growing in alkaline or neutral pH. A strategy to increase the overall SRL and therefore presumably increase nutrient uptake is reducing the length while increasing the thickness of the primary root and increasing the length of thin lateral roots (Hammond et al. 2004). For plants growing in acidic pH, thin roots (0.1–1 mm) accounted for almost 80% of total root length, compared with 57% under alkaline and 71% under neutral conditions. This reduction in mean root diameter at acidic pH is in line with previous studies of relative diameter class length of lupine (Chen et al. 2012), where approximately 70% of total root length was in the diameter classes between 0.1 and 1.0 mm. Lupine plants in the present study grew well on low-pH sandy soils, but not on neutral to alkaline soils, consistent with results from a previous study (Tang et al. 1992). One of the strategies to grow at low pH might be the increase of SRL as a consequence of decreasing root diameter, as observed in our experiment in acidic conditions. This might be a beneficial trait to increase P uptake in acidic pH, as shown previously in a range of species by Hill et al. (2006).

In contrast to what was observed in acidic pH, alkaline conditions modified root morphology by increasing the proportion of thick roots. Usually, thicker roots to increase the overall SRL and therefore presumably increase nutrient uptake is reducing the length while increasing the thickness of the primary root and increasing the length of thin lateral roots (Hammond et al. 2004). For plants growing in acidic pH, thin roots (0.1–1 mm) accounted for almost 80% of total root length, compared with 57% under alkaline and 71% under neutral conditions. This reduction in mean root diameter at acidic pH is in line with previous studies of relative diameter class length of lupine (Chen et al. 2012), where approximately 70% of total root length was in the diameter classes between 0.1 and 1.0 mm. Lupine plants in the present study grew well on low-pH sandy soils, but not on neutral to alkaline soils, consistent with results from a previous study (Tang et al. 1992). One of the strategies to grow at low pH might be the increase of SRL as a consequence of decreasing root diameter, as observed in our experiment in acidic conditions. This might be a beneficial trait to increase P uptake in acidic pH, as shown previously in a range of species by Hill et al. (2006).

In contrast to what was observed in acidic pH, alkaline conditions modified root morphology by increasing the proportion of thick roots. Usually, thicker roots

| pH    | P source | Total root length (cm) | Average root diam. (mm) | Root surface area (cm²) | Specific root length (m g⁻¹ root) |
|-------|----------|------------------------|-------------------------|-------------------------|----------------------------------|
| Acidic| Struvite  | 538b                   | 0.8c                    | 143bc                   | 4.1a                             |
|       | KP       | 1025a                  | 1.7b                    | 273ab                   | 4.8a                             |
|       | Nil-P    | 404bc                  | 0.9c                    | 111c                    | 4.0a                             |
| Neutral| Struvite | 1327a                 | 2.6a                    | 404a                    | 3.2b                             |
|        | KP       | 1218a                  | 2.5a                    | 344a                    | 3.1b                             |
|        | Nil-P    | 280bc                  | 0.9c                    | 82c                     | 3.1b                             |
| Alkaline| Struvite | 165bc                  | 1.1bc                   | 55c                     | 2.4c                             |
|          | KP       | 104bc                  | 1.2bc                   | 39c                     | 2.0c                             |

For ANOVA see ESM Tables 15–22
account for approximately 4% of total root length (Chen et al. 2012), a similar proportion to what was found at acidic and neutral pH in the present study. However, at alkaline pH, this increased to 9%. The explanation might not be related to nutrient availability, but to the high pH itself. Under nutrient stress, an increase of SRL would be expected. However, plants growing in alkaline pH without P showed that the development of lateral and secondary roots was strongly inhibited; roots were thicker and mostly belonged to the top root part near the shoot and showed no further lateral root growth.

The SRL was affected by pH but did not vary among P applications. Plants with Nil-P showed no significant difference in SRL in comparison with the P-treated plants at the same pH. Differences in total root length between P applications were only observed in acidic pH. At pH 4.5, the total root length of KP-treated plants almost doubled in comparison with those grown with struvite, despite showing no differences in SRL. To what extent the plant benefits from the increased SRL in terms of additional P uptake is still a matter of debate (Zobel et al. 2007). As discussed below, narrow-leaf lupine’s physiological responses to different P sources were apparently more pronounced than morphological root modifications.

Physiological root adaptations to P source applied

Root-induced chemical changes in the rhizosphere or rhizosheath have a distinct effect on the availability of P (Marschner et al. 1986). Our study found that narrow-leaf lupine released a greater amount of carboxylates at neutral conditions when struvite was applied than with the readily available KP. We hypothesized that the increase in carboxylates exudation would enhance the bioavailability of P from struvite because of an increase in P solubility from struvite at lower pH. We observed lower rhizosheath pH values in neutral condition in comparison with the bulk-sand pH. However, pH measurements did not differ between the struvite treatment and the easily available KP. Therefore, the effect of an increased P solubility may be explained also by the displacement of P from the matrix, in this case of struvite, by anion exchange, as described before for other P sources (Lambers et al. 2006). This agrees with a study by Hinsinger et al. (2003) stating that the contribution of organic acids to acidification is likely less important than the contribution by the cation-anion balance to increased P solubility.

The exudation of carboxylates depended not only on the P source applied, but also on the initial soil pH at which the plants were grown. The total amount of carboxylates was very low in alkaline conditions, because root growth was strongly reduced. Interestingly, a
greater exudation per root surface area was observed at alkaline pH without differences between P treatments. This suggests that at high pH, carboxylate exudation is a key strategy of narrow-leaf lupine to reduce pH as well as to mobilize P.

Our results agree with those of Talboys et al. (2016) who affirmed that struvite’s effectiveness as a P fertilizer is greater for crop species that exude large quantities of organic anions. They compared *Triticum aestivum* (spring wheat), a species exhibiting little exudation, and *Fagopyrum esculentum* (buckwheat), a species exhibiting abundant exudation, treated with either struvite, triple-superphosphate or di-ammonium phosphate. They found that buckwheat grew better on struvite than spring wheat did, suggesting that the larger amount of exudates of buckwheat can explain its higher yield and apparent fertilizer recovery (Talboys et al. 2016). Thus, the use of species that release large amounts of exudates would be a good strategy for alkaline soils where struvite is not initially soluble. However, calcicole species that are less sensitive to a high pH than *L. angustifolius* should be further investigated, e.g., *L. pilosus* and *L. cosentinii* (Brand et al. 2002; Ding et al. 2018).

Growth of narrow-leaf lupine under different pH and P sources

The response of narrow-leaf lupine to P application showed substantial differences in biomass under different pH, but no differences among struvite and KP at any pH. Allocation patterns showed that both P sources are functioning similarly in terms of biomass production. As expected, plant growth was reduced without the addition of P in all pH conditions. Likewise, plant growth was inhibited at alkaline conditions, with no significant differences in biomass production among the P treatments. The chlorosis of old leaves at alkaline pH or when no P was supplied was observed in most of the plants, possibly reflecting the process of translocation of P and other nutrients to young leaves. Other nutrients other than P were applied at sufficient concentrations for plant growth. Generally, high pH was not optimal for plant growth; however, the high content of mineral nutrients in large-seeded species like narrow-leaf lupine, and their translocation from cotyledons during early growth possibly enabled seedling growth under stress conditions for 2 weeks (Milberg et al. 1998). A pH above 6.0 specifically reduces nodulation in some lupine species (Tang and Robson 1993), as also observed in this experiment at alkaline conditions.

The biomass of narrow-leaf lupine when P was supplied (indistinctly of the P form) was 80% lower at alkaline pH and 65% lower at acidic conditions compared with plants grown at pH 6.5. The optimal pH for growth of narrow-leaf lupine is between 5.0 and 5.5 (Tang et al. 1992), similar to the rhizosheath pH found in the present study at neutral condition. At acidic pH, the solubility of struvite is higher due to its physical-chemical properties. The struvite crystallization process is affected by several factors such as temperature or pH (Huchzermeier and Tao 2012). High pH favors the formation of struvite crystals (pH range of 7 to 11), whereas low pH favors its solubilization (Bhuiyan et al. 2007). Therefore, as expected, the effectiveness of struvite and KP as P-fertilizers at acidic pH was similar. At neutral pH, the effectiveness of both P sources was also similar despite the lower solubility of struvite at this pH. Both P treatments resulted in higher total biomass compared to plants where no P was added, with no significant difference between them. This observation is likely due to the carboxylate exudation by the narrow-leaf lupine roots when struvite was applied, mobilizing the struvite-P at similar rates as the readily soluble KP-P.

Phosphorus-uptake efficiency and P concentration in shoots and roots

As for the biomass, it was expected that P concentration of narrow-leaf lupine would decrease at high pH, as shown previously in other lupine species grown at a pH above 6 (Jessop and Mahoney 1982). However, narrow-leaf lupine grown in alkaline pH exhibited significantly higher P concentration compared with acidic and neutral conditions. At this pH, where plants only reached early developmental stage, P was likely not the limiting nutrient. In plants grown at lower pH conditions the P concentration in old leaves decreased with age (Imo and Timmer 1992), as measured in plants growing in acidic pH.

The P-uptake efficiency (PUE) from struvite at acidic pH was 60% greater than that from KP. The higher P concentration and PUE of struvite-treated plants in acidic pH was related to a higher SRL of narrow-leaf lupine at acidic pH, but also due to the high solubility of struvite at this pH. The measured shoot P concentration in the plants grown at alkaline pH was 22 mg P g−1 dry
mass and 11 mg P g\(^{-1}\) dry mass at acidic pH, both under struvite treatment, which can be considered as toxic (Römheld 2012; Shane et al. 2004a,b). This might explain why struvite-treated plants in acidic conditions were slightly smaller than those treated with KP, as for other lupine species like \(L.\) micranthus high sensitivity to P toxicity has been reported, showing growth inhibition after exposure to a high P supply (Abdolzadeh et al. 2010). This indicates a low capacity to down-regulate the P-uptake system which might also be the case in narrow-leaf lupine, with the increase of the SRL. However, no visual signs of P or any other nutrient toxicity were observed.

At neutral pH, no difference in PUE was found. This is probably related to the greater exudation of carboxylates in the struvite treatment as discussed above.

We suggest that for plants adapted to neutral or acidic conditions such as narrow-leaf lupine that are able to exude large amounts of carboxylates, no difference in biomass between struvite and KP treatments is expected at those pH values. Lower amount of struvite than of KP in acidic conditions might be recommended, as the P concentration in plant tissues measured in the struvite treatment was at the level considered toxic, even though no P-toxicity symptoms were observed.

Conclusions

The effectiveness of struvite as a P source can be enhanced in crop species that exude large amounts of carboxylates by P mobilization via cation exchange. Narrow-leaf lupine released greater amounts of organic anions at neutral pH when struvite was applied than when the readily available KP was used. Large quantities of malate and citrate were found in the rhizosheath, reaching concentrations in the low mM range. As a consequence, biomass at neutral pH was similar for plants treated with struvite and with the highly plant-available KP. This confirms the hypothesis that narrow-leaf lupine can increase struvite P solubility, resulting in an enhanced plant P availability. However, this was not observed at alkaline pH, when root growth of the calcifuge narrow-leaf lupine was strongly inhibited. Interestingly, even though the total amount of exudates was low in alkaline conditions, the amount of carboxylates per unit root surface area was even higher than under neutral pH. This suggests that carboxylate exudation is a general strategy of narrow-leaf lupine to acidify the rhizosphere and to mobilize soil P. At acidic pH, plants supplied with struvite had a higher P concentration and PUE compared to plants supplied with KP, explained by a higher SRL, and also due to the high solubility of struvite at this pH. The P concentration in plant tissue under acidic conditions can be considered as toxic and might explain why struvite-treated plants grown at this pH were slightly smaller than those treated with KP.

Our study showed that struvite used as a slow-release fertilizer, could be utilized by crop species with root morphological and physiological strategies to mobilize P from a crop root system. Struvite might be more efficient for the production of biomass compared to conventional highly soluble P fertilizers, given the consideration of the plant’s adaptation to specific pH conditions.

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